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(54) **OXALOBACTER FORMIGENES (OF)-DERIVED FACTORS FOR THE TREATMENT/PREVENTION OF EXCESS OXALATE LEVELS**

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CPC *A61K 35/741* (2013.01); *A61K 38/164* (2013.01)

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(57) **ABSTRACT**

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Provided herein are compositions comprising *Oxalobacter formigenes* (OF)-derived factors and variants and fragments thereof, and method of use thereof for the treatment/prevention excess oxalate levels and conditions and diseases related thereto.

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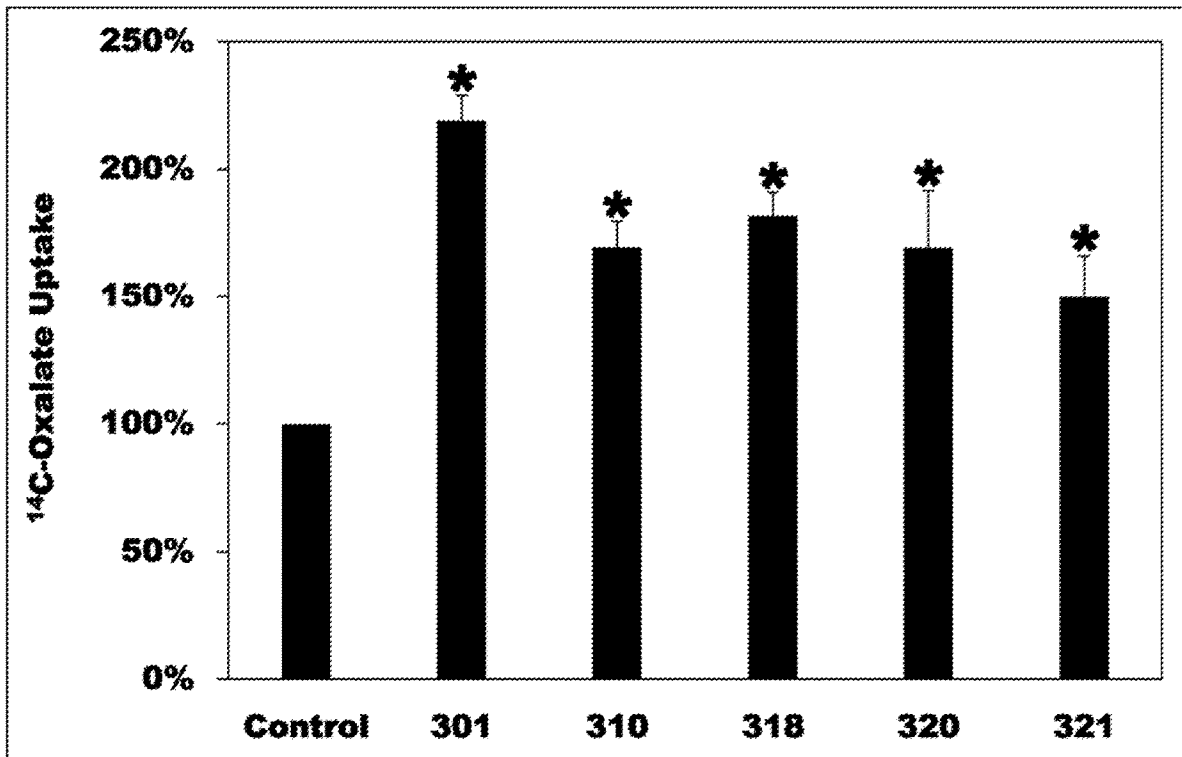


FIG. 1

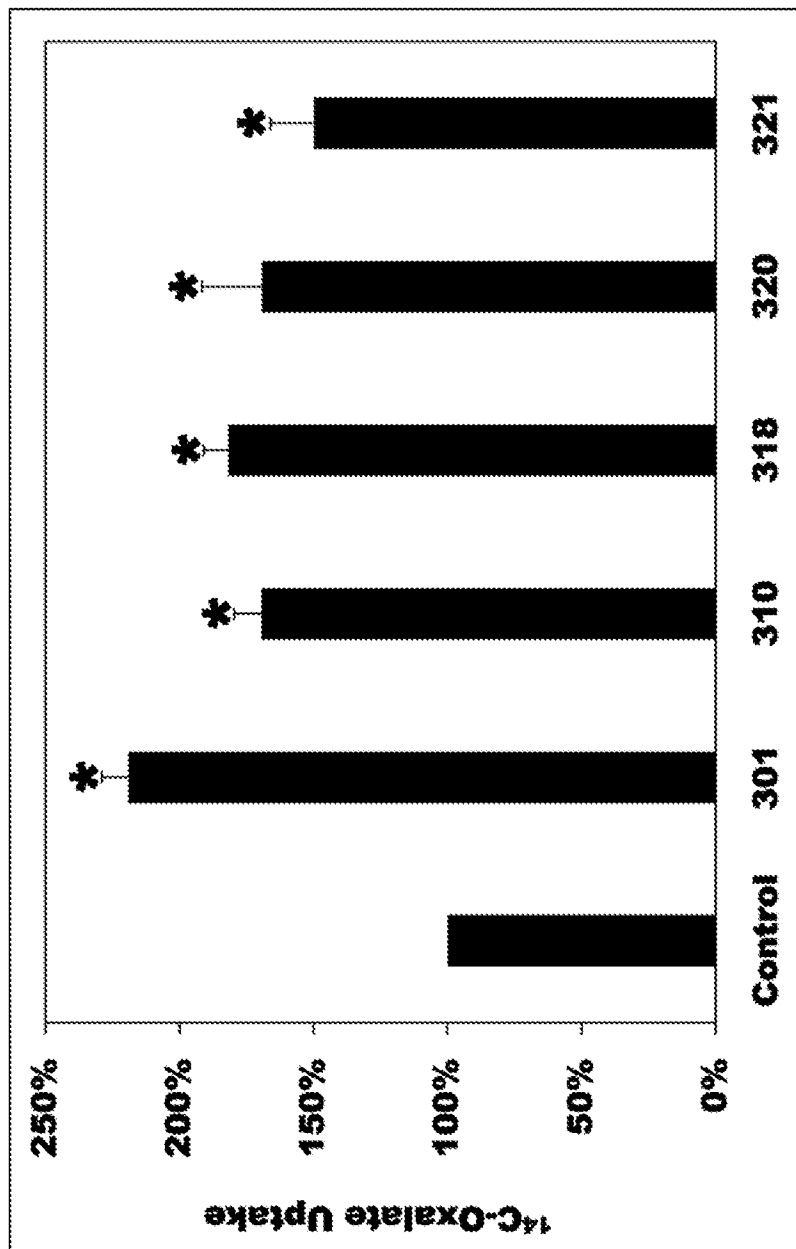


FIG. 2

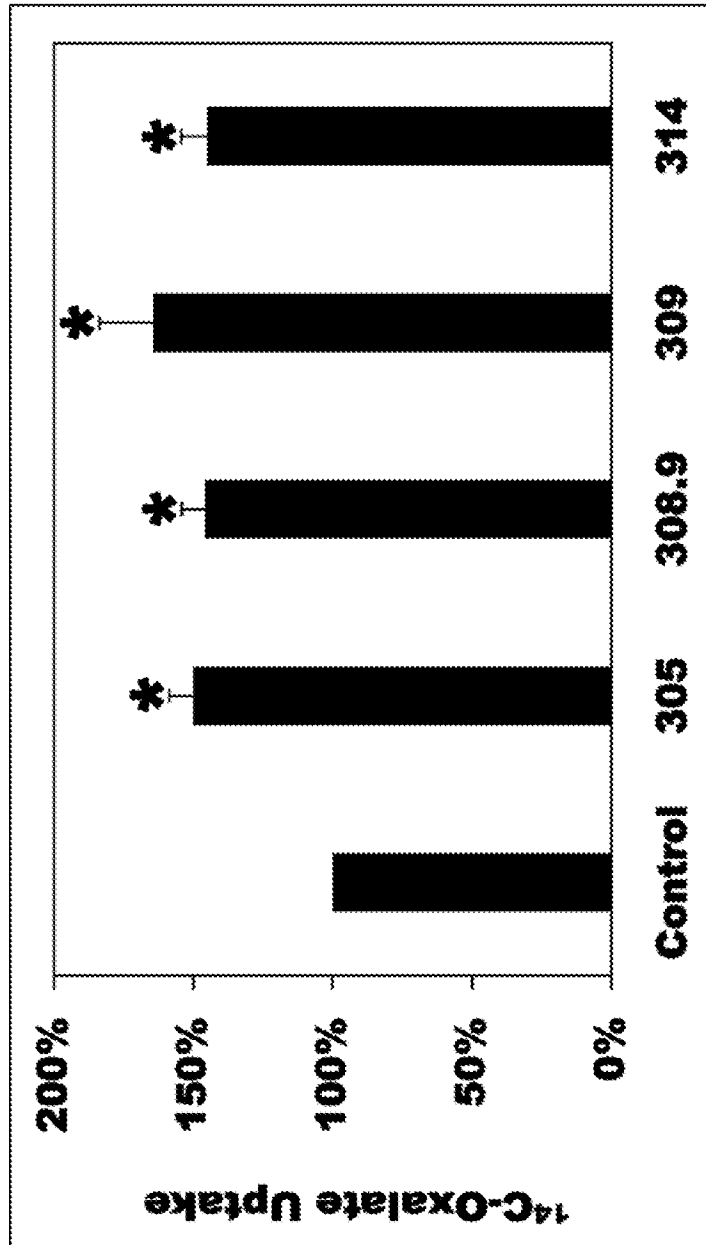


FIG. 3

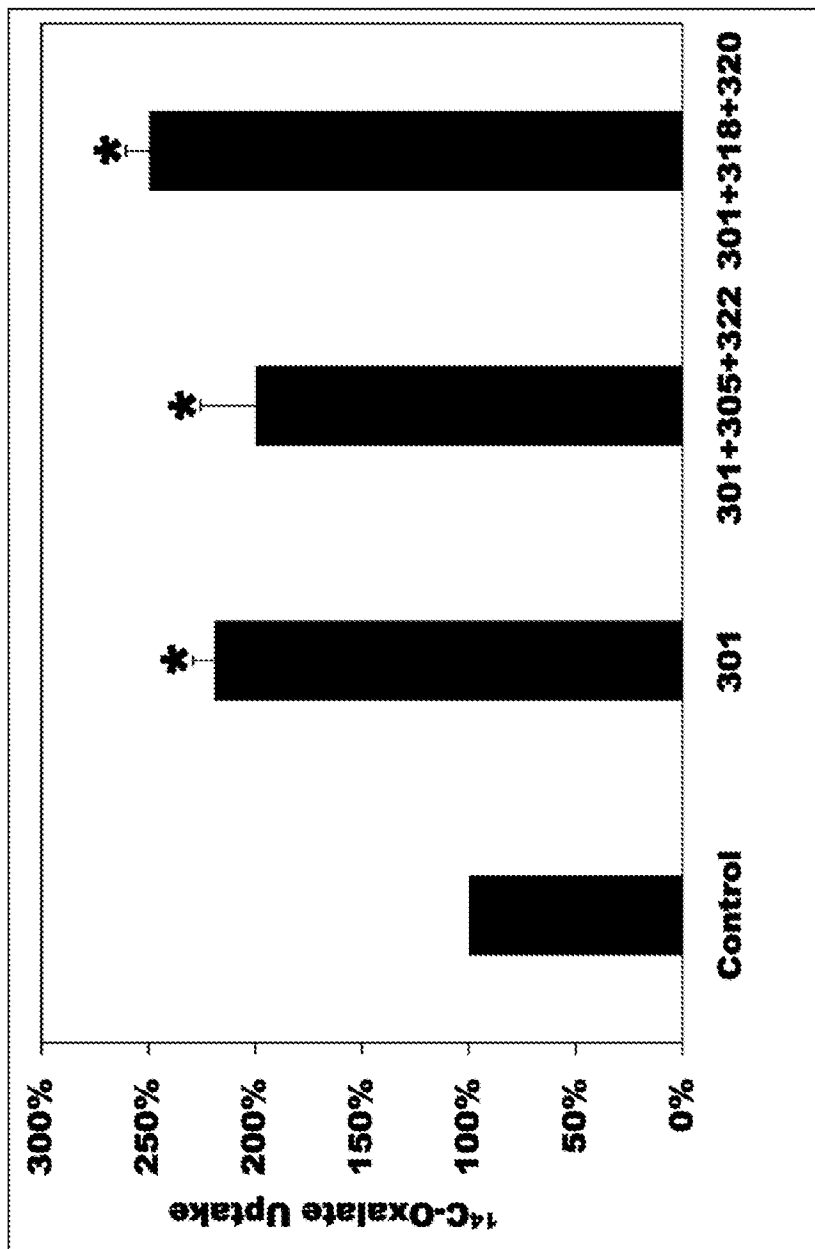


FIG. 4

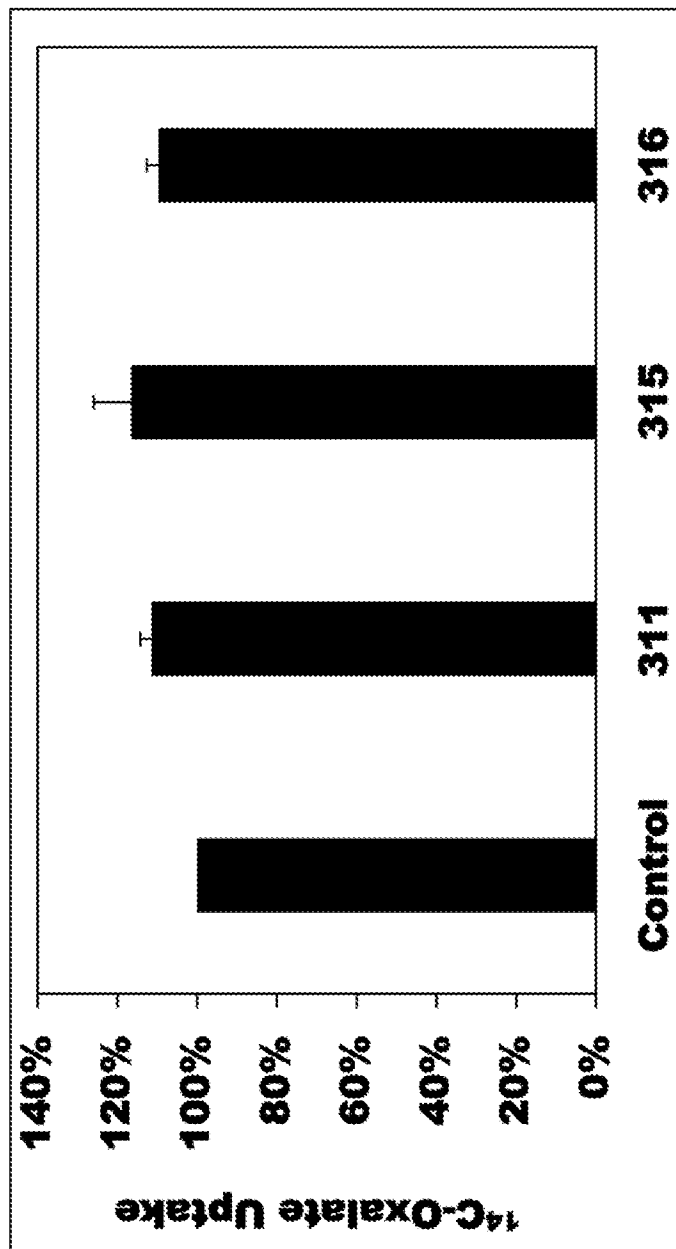


FIG. 5

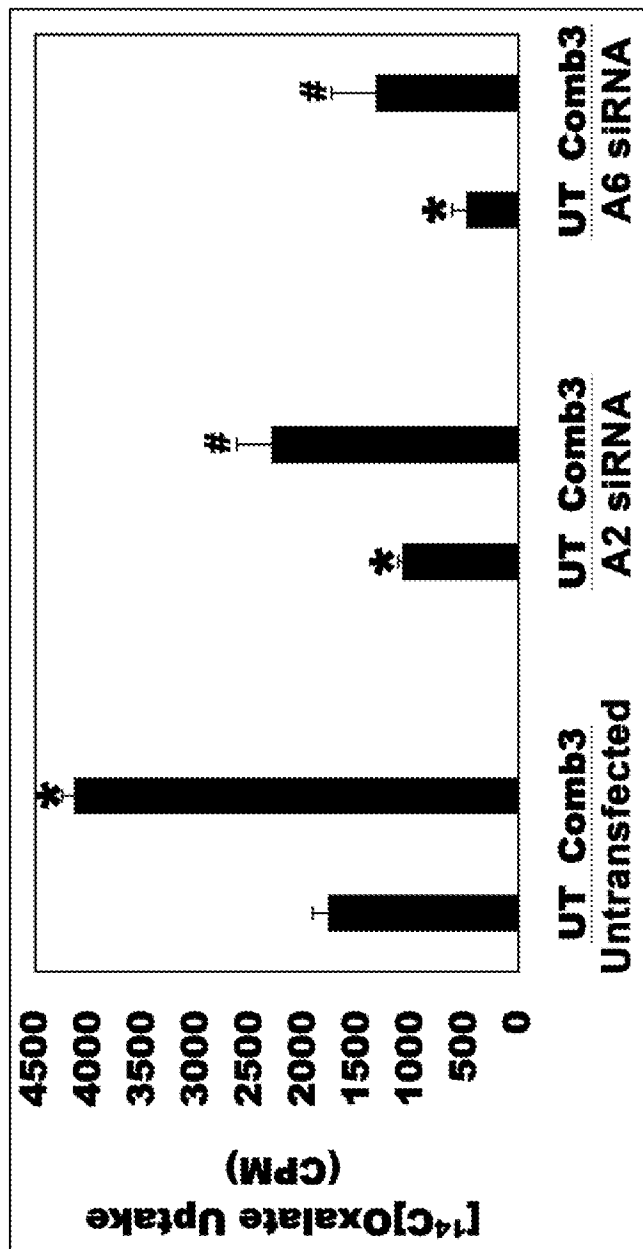
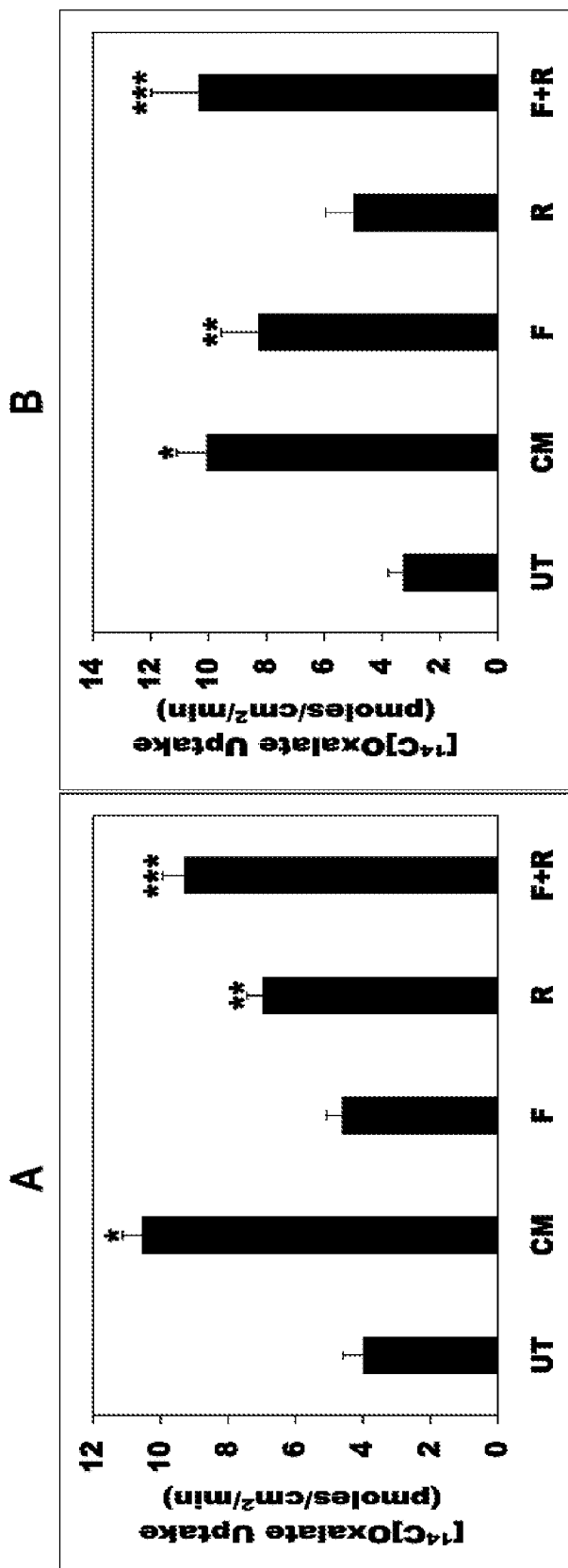


FIG. 6



**OXALOBACTER FORMIGENES
(OF)-DERIVED FACTORS FOR THE
TREATMENT OF
TREATMENT/PREVENTION OF EXCESS
OXALATE LEVELS**

CROSS-REFERENCE TO RELATED
APPLICATIONS

[0001] The present invention claims priority to U.S. Provisional Patent Application Ser. No. 62/448,178, filed Jan. 19, 2017, which is incorporated by reference in its entirety.

STATEMENT REGARDING FEDERAL
FUNDING

[0002] This invention was made with government support under Grant No. DK067245 and Grant No. DK042086 awarded by the National Institute of Health. The government has certain rights in the invention.

FIELD

[0003] Provided herein are compositions comprising *Oxalobacter formigenes* (O)-derived factors and variants and fragments thereof, and method of use thereof for the treatment/prevention excess oxalate levels and conditions and diseases related thereto.

BACKGROUND

[0004] Nephrolithiasis, or the formation of mineral deposit blockages in the kidney (kidney stones (KS)), is the second most prevalent kidney disease in USA after hypertension, with a rising prevalence and complications including advanced chronic kidney disease (CKD) and end stage renal disease (ESRD). It remains a major source of patient discomfort and disability, lost working days, and health-care expenditure, with an annual economic cost approaching \$10 billion. Hyperoxaluria (HO) is a major risk factor for KS, and 70-80% of KS are composed of calcium oxalate. Urinary oxalate is an important determinant of supersaturation, and the risk for stone formation is affected by small increases in urine oxalate. Oxalate is a metabolic end product that cannot be further metabolized and is highly toxic. The mammalian intestine plays a crucial role in oxalate homeostasis, by regulating the amount of absorbed dietary oxalate and providing an avenue for enteric oxalate excretion. Anion exchanger SLC26A6 (A6)-mediated intestinal oxalate secretion plays a critical role in preventing hyperoxaluria and calcium oxalate kidney stones (COKS). Inflammatory bowel disease patients have a significantly increased risk of KS due to the associated enteric hyperoxaluria. Obesity is a risk factor for KS and obese stone formers often have mild to moderate hyperoxaluria. Hyperoxaluria is also emerging as a major complication (developing in >50% of patients) of bariatric surgery for obesity. With the rising prevalence of obesity and increased utilization of bariatric surgery, it is expected that the incidence of hyperoxaluria and related COKS (including the associated cost burden) will continue to increase at a significant rate. Primary hyperoxaluria (PH) is an inherited disease in which there is endogenous oxalate overproduction, which leads to recurrent KS and/or progressive nephrocalcinosis, ESRD, as well as significant hyperoxalemia, systemic oxalosis and premature death. Systemic deposition of calcium oxalate (oxalosis) leads to bone disease, cardiac arrhythmias, car-

diomyopathy, skin ulcers, erythropoietin refractory anemia, and digital gangrene. The only treatment known to fully correct the underlying metabolic defect is liver transplantation or combined kidney-liver transplantation once ESRD develops. In addition, significant hyperoxalemia is also seen in ESRD. Cardiovascular diseases are the leading cause of morbidity and mortality in ESRD patients, and a recent report suggest that the ESRD-associated hyperoxalemia may contribute to this increased risk.

[0005] Unfortunately, there is currently no specific therapy that effectively lowers urine and/or plasma oxalate level(s), and the risk of recurrent COKS, nephrocalcinosis, oxalate nephropathy, ESRD, and systemic oxalosis remains substantial in the absence of treatment. *Oxalobacter formigenes* (Of) is an anaerobic bacterium that utilizes oxalate as its exclusive energy source. Of colonization correlates with reduced risk of COKS formation in a number of studies, presumably by reducing intestinal oxalate absorption and urinary oxalate excretion. In addition to degrading intraluminal dietary oxalate, Of also interacts with colonic epithelium by inducing distal colonic oxalate secretion, leading to reduced urinary excretion via a potential unknown secretagogue. Of colonization of PH1 mice (a mouse model of primary hyperoxaluria type 1) significantly reduced serum and urinary oxalate levels due to induction of colonic oxalate secretion. However, all PH1 mice lost colonization within 18 days when switched from a high oxalate/low calcium diet (1.5% oxalate/0.5% calcium; needed to induce and maintain colonization) to regular mouse chow (0.25% oxalate/1% calcium). In addition, colonization cannot be maintained without reducing dietary calcium, which contradicts the current recommendations to increase dietary calcium for preventing recurrent KS. Moreover, it has been suggested from studies in PH patients & PH1 mice that the intraluminal environment in PH is not supportive of sustained Of colonization. Collectively, maintaining Of colonization in the absence of high exogenous oxalate remains problematic (and therefore making use of live Of as a potential therapeutic agent impractical). Treatments and/or therapies for reducing and/or maintaining healthy serum and urinary oxalate levels are needed.

SUMMARY

[0006] Provided herein are compositions comprising *Oxalobacter formigenes* (Of)-derived factors and variants and fragments thereof, and method of use thereof for the treatment/prevention of excess oxalate levels and conditions and diseases related thereto. In some embodiments, the Of-derived factors are derived from the OxB strain and/or OXCC13 strain.

[0007] In some embodiments, provided herein are methods comprising administering to the subject one or more *Oxalobacter formigenes* (Of)-derived factors, and/or bioactive variants and/or fragments thereof, that result in stimulation of oxalate transport. In some embodiments, conditioned media (CM) (or factors derived therefrom) stimulates oxalate transport by human intestinal Caco2-BBE (C2) cells by activating protein kinase A (PKA) activation and increasing the transport activity of SLC26A6. In some embodiments, Of-derived factors are administered by administering Of CM to the subject. In some embodiments, the Of CM is fractionated, purified, and/or otherwise processed prior to administration. In some embodiments, one or more Of-derived factors are administered by administering

NO: 24, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, and/or, SEQ ID NO: 102. In some embodiments, the pharmaceutical composition is formulated for rectal administration, oral administration, and/or injection.

[0010] In some embodiments, provided herein are methods of treating or preventing hyperoxaluria and/or hyperoxalemia comprising administering a pharmaceutical described herein to a subject. In some embodiments, treating or preventing hyperoxaluria and/or hyperoxalemia lowers the subject's risk of the risk of calcium oxalate kidney stones, nephrocalcinosis, oxalate nephropathy, end stage renal disease, and/or systemic oxalosis.

BRIEF DESCRIPTION OF THE DRAWINGS

[0011] FIG. 1. Sel1 Proteins 301, 310, 318, 320, and 321 significantly stimulate ^{14}C -oxalate influx into Caco2-BBE (C2) cells.

[0012] FIG. 2. Non-Sel1 proteins 305, 308.9, 309, and 314 significantly stimulate ^{14}C -oxalate influx into Caco2-BBE (C2) cells.

[0013] FIG. 3. Combination of 301+318+320 stimulate ^{14}C -oxalate influx into Caco2-BBE (C2) cells more than 301 alone.

[0014] FIG. 4. Non-Sel1 proteins 311, 315, and 316 have no significant effect on ^{14}C -oxalate influx into Caco2-BBE cells.

[0015] FIG. 5. siRNA knockdown of the oxalate transporters SLC26A6 (A6) and SLC26A2 (A2) greatly reduced the Sel1 (Comb3=301+318+320)-induced stimulation of ^{14}C -oxalate influx into C2 cells, indicating that Sel1 proteins stimulate oxalate transport by C2 cells through mechanisms including enhanced A6 and A2 transport activities.

[0016] FIG. 6. Effect of selective ultrafiltration using 10 kDa (Panel A) and 30 kDa (Panel B) cutoff spin columns on the CM-induced stimulation of ^{14}C -oxalate uptake by Caco2-BBE (C2) cells. C2 cells were untreated (UT) or were treated with the conditioned medium (CM), the filtrate (F), the retentate (R), or the combined fractions (F+R). The results indicate that the molecular weights of the factors are largely between 10-30 kDa. Since F+R have a better stimulatory effect than F, while R has no effect, such data suggest the possibility that these factors might exist as a multifunctional complex requiring a bacterial product of >30 kDa for optimal functioning. These results support testing different combinations of Sel1 and other identified proteins to obtain a level of oxalate transport stimulation similar to that observed with the CM.

DEFINITIONS

[0017] Although any methods and materials similar or equivalent to those described herein can be used in the practice or testing of embodiments described herein, some preferred methods, compositions, devices, and materials are described herein. However, before the present materials and methods are described, it is to be understood that this invention is not limited to the particular molecules, compositions, methodologies or protocols herein described, as these may vary in accordance with routine experimentation and optimization. It is also to be understood that the terminology used in the description is for the purpose of describ-

ing the particular versions or embodiments only, and is not intended to limit the scope of the embodiments described herein.

[0018] Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. However, in case of conflict, the present specification, including definitions, will control. Accordingly, in the context of the embodiments described herein, the following definitions apply.

[0019] As used herein and in the appended claims, the singular forms “a”, “an” and “the” include plural reference unless the context clearly dictates otherwise. Thus, for example, reference to “an Of-derived factor” is a reference to one or more Of-derived factors and equivalents thereof known to those skilled in the art, and so forth.

[0020] As used herein, the term “comprise” and linguistic variations thereof denote the presence of recited feature(s), element(s), method step(s), etc. without the exclusion of the presence of additional feature(s), element(s), method step(s), etc. Conversely, the term “consisting of” and linguistic variations thereof, denotes the presence of recited feature(s), element(s), method step(s), etc. and excludes any unrecited feature(s), element(s), method step(s), etc., except for ordinarily-associated impurities. The phrase “consisting essentially of” denotes the recited feature(s), element(s), method step(s), etc. and any additional feature(s), element(s), method step(s), etc. that do not materially affect the basic nature of the composition, system, or method. Many embodiments herein are described using open “comprising” language. Such embodiments encompass multiple closed “consisting of” and/or “consisting essentially of” embodiments, which may alternatively be claimed or described using such language.

[0021] As used herein, the term “hyperoxaluria” refers to the excessive urinary excretion of oxalate by a subject (e.g., >25 mg/day).

[0022] As used herein, the term “hyperoxalemia” refers to excessive plasma levels of oxalate in a subject. Various studies report a normal range of oxalate in the plasma of 1 to 3 μmol per liter. Subjects with levels exceeding that range are considered to suffer from hyperoxalemia.

[0023] The term “amino acid” refers to natural amino acids, unnatural amino acids, and amino acid analogs, all in their D and L stereoisomers, unless otherwise indicated, if their structures allow such stereoisomeric forms.

[0024] Natural amino acids include alanine (Ala or A), arginine (Arg or R), asparagine (Asn or N), aspartic acid (Asp or D), cysteine (Cys or C), glutamine (Gln or Q), glutamic acid (Glu or E), glycine (Gly or G), histidine (His or H), isoleucine (Ile or I), leucine (Leu or L), Lysine (Lys or K), methionine (Met or M), phenylalanine (Phe or F), proline (Pro or P), serine (Ser or S), threonine (Thr or T), tryptophan (Trp or W), tyrosine (Tyr or Y) and valine (Val or V).

[0025] Unnatural amino acids include, but are not limited to, azetidinecarboxylic acid, 2-amino adipic acid, 3-amino adipic acid, beta-alanine, naphthylalanine (“naph”), aminopropionic acid, 2-aminobutyric acid, 4-aminobutyric acid, 6-aminocaproic acid, 2-aminoheptanoic acid, 2-aminoisobutyric acid, 3-aminoisobutyric acid, 2-aminopimelic acid, tertiary-butylglycine (“tBuG”), 2,4-diaminoisobutyric acid, desmosine, 2,2'-diaminopimelic acid, 2,3-diaminopropionic acid, N-ethylglycine, N-ethylasparagine, homoproline

("hPro" or "homoP"), hydroxylysine, allo-hydroxylysine, 3-hydroxyproline ("3Hyp"), 4-hydroxyproline ("4Hyp"), isodesmosine, allo-isoleucine, N-methylalanine ("MeAla" or "Nime"), N-alkylglycine ("NAG") including N-methylglycine, N-methylisoleucine, N-alkylpentylglycine ("NAPG") including N-methylpentylglycine, N-methylvaline, naphthylalanine, norvaline ("Norval"), norleucine ("Norleu"), octylglycine ("OctG"), ornithine ("Orn"), pentylglycine ("pG" or "PGly"), pipercolic acid, thioproline ("ThioP" or "tPro"), homoLysine ("hLys"), and homoArginine ("hArg").

[0026] The term "amino acid analog" refers to a natural or unnatural amino acid where one or more of the C-terminal carboxy group, the N-terminal amino group and side-chain functional group has been chemically blocked, reversibly or irreversibly, or otherwise modified to another functional group. For example, aspartic acid-(beta-methyl ester) is an amino acid analog of aspartic acid; N-ethylglycine is an amino acid analog of glycine; or alanine carboxamide is an amino acid analog of alanine. Other amino acid analogs include methionine sulfoxide, methionine sulfone, S-(carboxymethyl)-cysteine, S-(carboxymethyl)-cysteine sulfoxide and S-(carboxymethyl)-cysteine sulfone.

[0027] As used herein, the term "peptide" refers a short polymer of amino acids linked together by peptide bonds. In contrast to other amino acid polymers (e.g., proteins, polypeptides, etc.), peptides are of about 50 amino acids or less in length. A peptide may comprise natural amino acids, non-natural amino acids, amino acid analogs, and/or modified amino acids. A peptide may be a subsequence of naturally occurring protein or a non-natural (synthetic) sequence.

[0028] As used herein, the term "mutant peptide" or "variant peptide" refers to a peptide having a distinct amino acid sequence from the most common variant occurring in nature, referred to as the "wild-type" sequence. A mutant peptide may be a subsequence of a mutant protein or polypeptide (e.g., a subsequence of a naturally-occurring protein that is not the most common sequence in nature) or may be a peptide that is not a subsequence of a naturally occurring protein or polypeptide. For example, a "mutant SLR peptide" (e.g., a "mutant Sell protein") may be a subsequence of a mutant version of SLR protein (e.g., Sell protein) or may be distinct sequence not found in naturally-occurring SLR proteins (e.g., Sell proteins).

[0029] As used herein, the term "mutant polypeptide" or "variant polypeptide" refers to a polypeptide having a distinct amino acid sequence from the "wild-type" sequence. A mutant polypeptide may be a naturally-occurring protein that is not the most common sequence in nature (or a polypeptide fragment thereof) or may be a polypeptide that is not a subsequence of a naturally occurring protein or polypeptide. For example, a "mutant SLR polypeptide" may be a naturally occurring SLR protein (e.g., Sell protein), a polypeptide fragment of a SLR protein (e.g., Sell protein), or may be distinct sequence not found in naturally-occurring SLR proteins (e.g., Sell proteins).

[0030] As used herein, the term "artificial peptide" or "artificial polypeptide" refers to a peptide or polypeptide having a distinct amino acid sequence from those found in natural peptides and/or proteins. An artificial protein is not a subsequence of a naturally occurring protein, either the wild-type (i.e., most abundant) or mutant versions thereof. For example, an artificial SLR peptide or polypeptide is not

a subsequence of naturally occurring SLR protein (e.g., Sell protein). An artificial peptide or polypeptide may be produced or synthesized by any suitable method (e.g., recombinant expression, chemical synthesis, enzymatic synthesis, etc.).

[0031] The terms "peptide mimetic" or "peptidomimetic" refer to a peptide-like molecule that emulates a sequence derived from a protein or peptide. A peptide mimetic or peptidomimetic may contain amino acids and/or non-amino acid components. Examples of peptidomimetics include chemically modified peptides, peptoids (side chains are appended to the nitrogen atom of the peptide backbone, rather than to the α -carbons), β -peptides (amino group bonded to the R carbon rather than the α carbon), etc.

[0032] As used herein, a "conservative" amino acid substitution refers to the substitution of an amino acid in a peptide or polypeptide with another amino acid having similar chemical properties, such as size or charge. For purposes of the present disclosure, each of the following eight groups contains amino acids that are conservative substitutions for one another:

1) Alanine (A) and Glycine (G);

[0033] 2) Aspartic acid (D) and Glutamic acid (E);

3) Asparagine (N) and Glutamine (Q);

4) Arginine (R) and Lysine (K);

5) Isoleucine (I), Leucine (L), Methionine (M), and Valine (V);

6) Phenylalanine (F), Tyrosine (Y), and Tryptophan (W);

7) Serine (S) and Threonine (T); and

8) Cysteine (C) and Methionine (M).

[0034] Naturally occurring residues may be divided into classes based on common side chain properties, for example: polar positive (histidine (H), lysine (K), and arginine (R)); polar negative (aspartic acid (D), glutamic acid (E)); polar neutral (serine (S), threonine (T), asparagine (N), glutamine (Q)); non-polar aliphatic (alanine (A), valine (V), leucine (L), isoleucine (I), methionine (M)); non-polar aromatic (phenylalanine (F), tyrosine (Y), tryptophan (W)); proline and glycine; and cysteine. As used herein, a "semi-conservative" amino acid substitution refers to the substitution of an amino acid in a peptide or polypeptide with another amino acid within the same class.

[0035] In some embodiments, unless otherwise specified, a conservative or semi-conservative amino acid substitution may also encompass non-naturally occurring amino acid residues that have similar chemical properties to the natural residue. These non-natural residues are typically incorporated by chemical peptide synthesis rather than by synthesis in biological systems. These include, but are not limited to, peptidomimetics and other reversed or inverted forms of amino acid moieties. Embodiments herein may, in some embodiments, be limited to natural amino acids, non-natural amino acids, and/or amino acid analogs. Non-conservative substitutions may involve the exchange of a member of one class for a member from another class.

[0036] As used herein, the term "sequence identity" refers to the degree to which two polymer sequences (e.g., peptide,

polypeptide, nucleic acid, etc.) have the same sequential composition of monomer subunits. The term “sequence similarity” refers to the degree with which two polymer sequences (e.g., peptide, polypeptide, nucleic acid, etc.) differ only by conservative and/or semi-conservative amino acid substitutions. The “percent sequence identity” (or “percent sequence similarity”) is calculated by: (1) comparing two optimally aligned sequences over a window of comparison (e.g., the length of the longer sequence, the length of the shorter sequence, a specified window, etc.), (2) determining the number of positions containing identical (or similar) monomers (e.g., same amino acids occurs in both sequences, similar amino acid occurs in both sequences) to yield the number of matched positions, (3) dividing the number of matched positions by the total number of positions in the comparison window (e.g., the length of the longer sequence, the length of the shorter sequence, a specified window), and (4) multiplying the result by 100 to yield the percent sequence identity or percent sequence similarity. For example, if peptides A and B are both 20 amino acids in length and have identical amino acids at all but 1 position, then peptide A and peptide B have 95% sequence identity. If the amino acids at the non-identical position shared the same biophysical characteristics (e.g., both were acidic), then peptide A and peptide B would have 100% sequence similarity. As another example, if peptide C is 20 amino acids in length and peptide D is 15 amino acids in length, and 14 out of 15 amino acids in peptide D are identical to those of a portion of peptide C, then peptides C and D have 70% sequence identity, but peptide D has 93.3% sequence identity to an optimal comparison window of peptide C. For the purpose of calculating “percent sequence identity” (or “percent sequence similarity”) herein, any gaps in aligned sequences are treated as mismatches at that position.

[0037] As used herein, the term “subject” broadly refers to any animal, including but not limited to, human and non-human animals (e.g., dogs, cats, cows, horses, sheep, poultry, fish, crustaceans, etc.). As used herein, the term “patient” typically refers to a human subject that is being treated for a disease or condition.

[0038] As used herein, the term “effective amount” refers to the amount of a sufficient to effect beneficial or desired results. An effective amount can be administered in one or more administrations, applications or dosages and is not intended to be limited to a particular formulation or administration route.

[0039] As used herein, the terms “administration” and “administering” refer to the act of giving a drug, prodrug, or other agent, or therapeutic treatment to a subject or in vivo, in vitro, or ex vivo cells, tissues, and organs. Exemplary routes of administration to the human body can be through space under the arachnoid membrane of the brain or spinal cord (intrathecal), the eyes (ophthalmic), mouth (oral), skin (topical or transdermal), nose (nasal), lungs (inhalant), oral mucosa (buccal), ear, rectal, vaginal, by injection (e.g., intravenously, subcutaneously, intratumorally, intraperitoneally, etc.) and the like.

[0040] As used herein, the terms “co-administration” and “co-administering” refer to the administration of at least two agent(s) or therapies to a subject. In some embodiments, the co-administration of two or more agents or therapies is concurrent. In other embodiments, a first agent/therapy is administered prior to a second agent/therapy. Those of skill

in the art understand that the formulations and/or routes of administration of the various agents or therapies used may vary. The appropriate dosage for co-administration can be readily determined by one skilled in the art. In some embodiments, when agents or therapies are co-administered, the respective agents or therapies are administered at lower dosages than appropriate for their administration alone. Thus, co-administration is especially desirable in embodiments where the co-administration of the agents or therapies lowers the requisite dosage of a potentially harmful (e.g., toxic) agent(s), and/or when co-administration of two or more agents results in sensitization of a subject to beneficial effects of one of the agents via co-administration of the other agent.

[0041] As used herein, the term “treatment” means an approach to obtaining a beneficial or intended clinical result. The beneficial or intended clinical result may include alleviation of symptoms, a reduction in the severity of the disease, inhibiting a underlying cause of a disease or condition, steadying diseases in a non-advanced state, delaying the progress of a disease, and/or improvement or alleviation of disease conditions.

[0042] As used herein, the term “pharmaceutical composition” refers to the combination of an active agent (e.g., Of-derived factor) with a carrier, inert or active, making the composition especially suitable for diagnostic or therapeutic use in vitro, in vivo or ex vivo. The terms “pharmaceutically acceptable” or “pharmacologically acceptable,” as used herein, refer to compositions that do not substantially produce adverse reactions, e.g., toxic, allergic, or immunological reactions, when administered to a subject.

[0043] As used herein, the term “pharmaceutically acceptable carrier” refers to any of the standard pharmaceutical carriers including, but not limited to, phosphate buffered saline solution, water, emulsions (e.g., such as an oil/water or water/oil emulsions), and various types of wetting agents, any and all solvents, dispersion media, coatings, sodium lauryl sulfate, isotonic and absorption delaying agents, disintegrants (e.g., potato starch or sodium starch glycolate), and the like. The compositions also can include stabilizers and preservatives. For examples of carriers, stabilizers and adjuvants, see, e.g., Martin, Remington’s Pharmaceutical Sciences, 15th Ed., Mack Publ. Co., Easton, Pa. (1975), incorporated herein by reference in its entirety.

DETAILED DESCRIPTION

[0044] Provided herein are compositions comprising *Oxalobacter formigenes* (Of)-derived factors and variants and fragments thereof, and methods of use thereof for the treatment/prevention excess oxalate levels and conditions and diseases related thereto.

[0045] Most kidney stones (KS) are composed of calcium oxalate, and small increases in urine oxalate affect the stone risk. The mammalian intestine plays a crucial role in oxalate homeostasis. Intestinal oxalate secretion mediated by anion exchanger SLC26A6 (A6) plays a major role in limiting net intestinal absorption of ingested oxalate; thereby preventing hyperoxaluria and calcium oxalate kidney stones (COKS). Hyperoxaluria and a high incidence of KS are commonly seen in IBD patients. Hyperoxaluria is also emerging as a major complication of bariatric surgery for obesity. Primary hyperoxaluria (PH) is an inherited disease in which there is endogenous oxalate overproduction. Enhancing intestinal oxalate secretion is expected to lead to reduced urine and

plasma oxalate levels. In addition to degrading intraluminal dietary oxalate, the probiotic bacterium *oxalobacter formigenes* (Of) also interacts with colonic epithelium by inducing colonic oxalate secretion, leading to reduced urinary excretion. Significant difficulties exist in sustaining Of colonization in animals and humans in the absence of high exogenous oxalate.

[0046] Experiments were conducted during development of embodiments herein to determine whether Of CM affects intestinal oxalate transport using the human intestinal Caco2-BBE (C2) cells. DIDS (anion exchange inhibitor)-sensitive apical oxalate influx was measured in the presence of an outward Cl gradient as an assay of Cl-oxalate exchange, $\geq 50\%$ of which is mediated by A6. Compared with control medium, Of CM significantly stimulated oxalate uptake (>2.4 -fold), whereas CM from *Lactobacillus acidophilus* (La) did not. Treating the CM with heat or pepsin completely abolished this bioactivity, and selective ultrafiltration of the CM revealed that the Of-derived factors have molecular masses of 10-30 kDa. Treatment with the PKA inhibitor H89 or DIDS completely blocked the CM-induced oxalate transport. A6 Knockdown also significantly restricted the induction of oxalate transport by CM. In a mouse model of primary hyperoxaluria type 1, rectal administration of Of CM significantly reduced ($>32.5\%$) urinary oxalate excretion and stimulated ($>42\%$) distal colonic oxalate secretion, reflecting the in vivo retention of biologic activity and the therapeutic potential of these factors.

[0047] Experiments conducted during development of embodiments herein to identify the Of-derived bioactive factor(s) inducing colonic oxalate secretion determined that small molecular weight protein(s) and/or peptide(s) secreted by Of in its culture conditioned medium (CM) significantly stimulate(s) oxalate transport (>2.4 -fold) by human intestinal Caco2-BBE cells through mechanisms including PKA activation and increased A6 transport activity. Rectal administration of Of CM significantly reduced ($>32.5\%$) urinary oxalate excretion and stimulated ($>42\%$) distal colonic oxalate secretion in PH1 mice. Probiotic bacteria have several health benefits; however, the difficulties in determining intestinal bacterial bioavailability and biosafety concerns when administering live probiotics are problems facing current probiotic clinical applications. These issues are compounded by the difficulties described above in maintaining Of colonization in the absence of high exogenous dietary oxalate.

[0048] Experiments conducted during development of embodiments herein demonstrate that Of-derived factors retain their biological activity when administered in vivo and effectively reduce urinary oxalate excretion in hyperoxaluric mice, thereby demonstrating the utility of the Of-derived factors as therapeutic agents for prevention and/or treatment of hyperoxaluria, hyperoxalemia, COKS, and related diseases/conditions.

[0049] Experiments conducted during development of embodiments herein have identified Sel1 repeat proteins as among the Of-derived factors responsible for stimulation of oxalate transport by C2 cells. Sel1 repeat proteins are involved in signal transduction pathways, and this is very important since we found that Of CM signals through PKA to stimulate oxalate transport by C2 cells. 12 Sel1 repeat proteins were purified and the effects of different concentrations and different incubation periods of the purified proteins on oxalate transport by C2 cells were assessed. Sel1

proteins 301, 310, 318, 320, and 321 significantly stimulate (~ 1.5 - 2.2 -fold) oxalate transport by C2 cells (FIG. 1). The combination of 301+318+320 have a better stimulatory effect (2.5-fold) compared to 301 (FIG. 3). 301+318 also stimulated oxalate transport by ~ 2.5 -fold. Sel1 proteins 304, 317, 319, 322, 323, 324, and 325 also stimulated oxalate transport by C2 cells by ~ 1.3 - 1.6 -fold. Since all of the 12 tested Sel1 proteins significantly stimulated oxalate transport by C2 cells, it is very likely that the remaining 32 Sel1 proteins (individually or in combination) will also stimulate oxalate and this will be tested. siRNA knockdown of the oxalate transporters SLC26A2 and SLC26A6 greatly reduced Sel1 (Comb3=301+318+320)-induced stimulation of oxalate transport (FIG. 5) as observed with the CM. In addition, 301-induced stimulation of oxalate transport is completely blocked by the PKA inhibitor H89, indicating that Sel1 proteins act through the PKA signaling pathway to stimulate oxalate transport, which is similar to the CM. Collectively, Sel1 proteins almost fully mimic the effects of the CM. Importantly, 3 non Sel1 proteins (311, 315, and 316) have no significant effects on oxalate transport by C2 cells (FIG. 4), strongly indicating that the Sel1-induced stimulation of oxalate transport by C2 cells is specific. In addition to Sel1 proteins, 4 other non Sel1 proteins (305, 308.9, 309, and 314) also stimulated oxalate transport (FIG. 2). In some embodiments, compositions are provided herein that comprise one or more Sel1 proteins or Sel1-derived variants or fragments (e.g., Sel1-derived peptides or polypeptides that stimulate oxalate transport). In some embodiments, methods are provided of treating/preventing excess oxalate levels in the urine and/or plasma by the administration of Sel1 proteins or Sel1-derived variants or fragments (e.g., Sel1-derived peptides or polypeptides that stimulate oxalate transport) to a subject. In some embodiments, compositions are provided herein that comprise one or more Sel1 proteins or Sel1-derived variants or fragments (e.g., Sel1-derived peptides or polypeptides that stimulate oxalate transport). In some embodiments, methods are provided of treating/preventing excess oxalate levels in the urine and/or plasma by the administration of Sel1 proteins or Sel1-derived variants or fragments (e.g., Sel1-derived peptides or polypeptides that stimulate oxalate transport) to a subject.

[0050] Sel1-like repeat (SLR) proteins (e.g. Sel1, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, and MerG) are involved in signal transduction pathways. In some embodiments, Of conditioned medium (CM) signals (e.g., through PKA) to stimulate oxalate transport (e.g., by C2 cells) as a result of the signal transduction functions of the SLR proteins (e.g., Sel1 proteins); although embodiments herein are not limited to any particular mechanism of action and an understanding of the mechanism of action is not necessary to practice such embodiments. SLR proteins (e.g., Sel1 proteins) have repeat units (e.g., repeat peptides). Most repeats are 5 to 40 amino acids (e.g., 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, or ranges therebetween), but longer repeat peptides (e.g., 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, 100, or longer or ranges therebetween) are within the scope of the SLR proteins (e.g., Sel1 proteins) herein. In some embodiments, repeat units fold into two to four secondary structural elements. In some embodiments, SLR proteins (e.g., Sel1 proteins) serve as adaptor proteins for the assembly of membrane-bound macromolecular complexes. Several bac-

terial and eukaryotic SLR proteins (e.g. Sell & Hrd3) are activated upon cellular stress. In some embodiments, Of Sell proteins are activated when oxalate is low in the culture medium (e.g., as evidenced by the observation of a CM of higher (>2-fold) bioactivity under this condition). Bacterial LpnE, EnhC, HcpA, ExoR, and AlgK proteins mediate the interactions between bacterial and eukaryotic host cells. In some embodiments, the SLR motif establishes a link between signal transduction pathways from eukaryotes and bacteria. In some embodiments, an SLR protein (e.g., Sell proteins) comprises leader sequences. In some embodiments, SLR proteins (e.g., Sell proteins) without leader sequences, such as PodJ or leaderless analogs of natural SLR proteins (e.g., Sell proteins), are active in the periplasmic space. In some embodiments, bacterial SLR proteins (e.g., Sell proteins), such as HcpA, ExoR, EnhC and LpnE are responsible for the adaptation of bacteria to different eukaryotic hosts.

[0051] The Of genome has 44 Sell proteins and one SelR domain, with many of these proteins having molecular masses between 10-30 kDa (e.g., where most of the CM

stimulatory activity lies) (Table 1). In addition, several other Sell proteins with molecular masses between 32-139 kDa were also identified (Table 1). Many of the Sell proteins are predicted to have signal peptides and therefore are secreted proteins. Moreover, Sell proteins with molecular masses of 25 (#1361), 33 (#1414), and 68 (#1362) kDa blast in common with another Sell protein having a molecular mass of 59 (#1344) kDa. Five Sell proteins (# s 1360-1364) resides in an operon, with carbon starvation protein CstA located immediately upstream of this operon. Sell proteins #1343, 1344, and 1356 are also located near this operon. In some embodiments, although most of the CM stimulatory activity is mediated by factors with molecular masses between 10-30 kDa, the results of selective ultrafiltration using 30 kDa cutoff column (FIG. 6B) indicates that the factors exist as a multifunctional complex requiring a bacterial product of >30 kDa for optimal functioning. The latter result indicates that different combinations of Sell and other identified proteins will provide a level of oxalate transport stimulation similar to (or greater than) that observed with the CM.

TABLE 1

Of-derived proteins				
Name	Protein	Description	DNA seq	Protein seq
301	194	Sell repeat :: PF08238	SEQ ID NO: 1	SEQ ID NO: 2
317	1343	Sell repeat :: PF08238	SEQ ID NO: 15	SEQ ID NO: 16
310	1344	Sell repeat :: PF08238	SEQ ID NO: 21	SEQ ID NO: 22
318	1356	Sell repeat :: PF08238	SEQ ID NO: 9	SEQ ID NO: 10
319	1360	Sell repeat :: PF08238	SEQ ID NO: 11	SEQ ID NO: 12
320	1361	Sell repeat :: PF08238	SEQ ID NO: 3	SEQ ID NO: 4
321	1362	Sell repeat :: PF08238	SEQ ID NO: 5	SEQ ID NO: 6
304	1363	Sell repeat :: PF08238	SEQ ID NO: 7	SEQ ID NO: 8
322	1364	Sell repeat :: PF08238	SEQ ID NO: 23	SEQ ID NO: 24
323	1414	Sell repeat :: PF08238	SEQ ID NO: 13	SEQ ID NO: 14
324	1548	Sell repeat :: PF08238	SEQ ID NO: 17	SEQ ID NO: 18
325	1549	Sell repeat :: PF08238	SEQ ID NO: 19	SEQ ID NO: 20
	193	Sell repeat :: PF08238	SEQ ID NO: 25	SEQ ID NO: 26
	235	Sell repeat :: PF08238	SEQ ID NO: 100	
	238	Sell repeat :: PF08238	SEQ ID NO: 27	SEQ ID NO: 28
	240	Sell repeat :: PF08238	SEQ ID NO: 29	SEQ ID NO: 30
	275	Sell repeat :: PF08238	SEQ ID NO: 31	SEQ ID NO: 32
	841	Sell repeat :: PF08238	SEQ ID NO: 33	SEQ ID NO: 34
	1112	Sell repeat :: PF08238	SEQ ID NO: 35	SEQ ID NO: 36
	1124	Sell repeat :: PF08238	SEQ ID NO: 37	SEQ ID NO: 38
	1143	Sell repeat :: PF08238	SEQ ID NO: 39	SEQ ID NO: 40
	1250	Sell repeat :: PF08238	SEQ ID NO: 41	SEQ ID NO: 42
	1257	SelR domain :: PF01641	SEQ ID NO: 43	SEQ ID NO: 44
	1412	Sell repeat :: PF08238	SEQ ID NO: 45	SEQ ID NO: 46
	1419	Sell repeat :: PF08238	SEQ ID NO: 47	SEQ ID NO: 48
	1423	Sell repeat :: PF08238	SEQ ID NO: 49	SEQ ID NO: 50
	1551	Sell repeat :: PF08238	SEQ ID NO: 51	SEQ ID NO: 52
	1715	Sell repeat :: PF08238	SEQ ID NO: 53	SEQ ID NO: 54
	1790	Sell repeat :: PF08238	SEQ ID NO: 55	SEQ ID NO: 56
	1942	Sell repeat :: PF08238	SEQ ID NO: 57	SEQ ID NO: 58
	1945	Sell repeat :: PF08238	SEQ ID NO: 59	SEQ ID NO: 60
	1954	Sell repeat :: PF08238	SEQ ID NO: 61	SEQ ID NO: 62
	1955	Sell repeat :: PF08238	SEQ ID NO: 63	SEQ ID NO: 64
	1960	Sell repeat :: PF08238	SEQ ID NO: 65	SEQ ID NO: 66
	1969	Sell repeat :: PF08238	SEQ ID NO: 67	SEQ ID NO: 68
	1970	Sell repeat :: PF08238	SEQ ID NO: 69	SEQ ID NO: 70
	2000	Sell repeat :: PF08238	SEQ ID NO: 71	SEQ ID NO: 72
	2001	Sell repeat :: PF08238	SEQ ID NO: 73	SEQ ID NO: 74
	2022	Sell repeat :: PF08238	SEQ ID NO: 75	SEQ ID NO: 76
	2025	Sell repeat :: PF08238	SEQ ID NO: 77	SEQ ID NO: 78
	2123	Sell repeat :: PF08238	SEQ ID NO: 79	SEQ ID NO: 80
	2176	Sell repeat :: PF08238	SEQ ID NO: 81	SEQ ID NO: 82
	2239	Sell repeat :: PF08238	SEQ ID NO: 83	SEQ ID NO: 84
	2249	Sell repeat :: PF08238	SEQ ID NO: 85	SEQ ID NO: 86
	2270	Sell repeat :: PF08238	SEQ ID NO: 87	SEQ ID NO: 88

TABLE 1-continued

Of-derived proteins				
Name	Protein	Description	DNA seq	Protein seq
314	8	LD-carboxypeptidase :: PF02016	SEQ ID NO: 89	SEQ ID NO: 90
309	1525	Peptidase family S49 :: PF01343	SEQ ID NO: 91	SEQ ID NO: 92
305	1524	haloacid dehalogenase-like hydrolase :: PF00702	SEQ ID NO: 98	SEQ ID NO: 101
315	272	Outer membrane efflux protein :: PF02321	SEQ ID NO: 93	SEQ ID NO: 94
316	1285	Outer membrane efflux protein :: PF02321	SEQ ID NO: 95	SEQ ID NO: 96
308	769	EAL domain :: PF00563	SEQ ID NO: 99	SEQ ID NO: 102
311	1847	MotA/TolQ/ExbB proton channel family :: PF01618		

[0052] To evaluate whether one or more of these SLR proteins (e.g., Sel1 proteins) are responsible for mediating the CM-induced stimulation of oxalate transport, a subset of Sel1 proteins were cloned and overexpressed in *E. coli* and

preliminary studies (FIG. 2). On the other hand, outer membrane efflux proteins (# s 272 and 1285) and MotA/TolQ/ExbB proton channel family (1847=311) failed to stimulate oxalate transport by C2 cells.

TABLE 2

Purified Of-derived proteins				
Protein name	Protein		DNA seq	Protein seq
301	194 Sel1 repeat :: PF08238		SEQ ID NO: 1	SEQ ID NO: 2
320	1361 Sel1 repeat :: PF08238		SEQ ID NO: 3	SEQ ID NO: 4
321	1362 Sel1 repeat :: PF08238		SEQ ID NO: 5	SEQ ID NO: 6
304	1363 Sel1 repeat :: PF08238		SEQ ID NO: 7	SEQ ID NO: 8
318	1356 Sel1 repeat :: PF08238		SEQ ID NO: 9	SEQ ID NO: 10
319	1360 Sel1 repeat :: PF08238		SEQ ID NO: 11	SEQ ID NO: 12
323	1414 Sel1 repeat :: PF08238		SEQ ID NO: 13	SEQ ID NO: 14
317	1343 Sel1 repeat :: PF08238		SEQ ID NO: 15	SEQ ID NO: 16
324	1548 Sel1 repeat :: PF08238		SEQ ID NO: 17	SEQ ID NO: 18
325	1549 Sel1 repeat :: PF08238		SEQ ID NO: 19	SEQ ID NO: 20
310	1344 Sel1 repeat :: PF08238		SEQ ID NO: 21	SEQ ID NO: 22
309	1525 Peptidase family S49 :: PF01343		SEQ ID NO: 91	SEQ ID NO: 92
315	272 Outer membrane efflux protein :: PF02321		SEQ ID NO: 93	SEQ ID NO: 94
316	1285 Outer membrane efflux protein :: PF02321		SEQ ID NO: 95	SEQ ID NO: 96
314	8 LD-carboxypeptidase :: PF02016		SEQ ID NO: 89	SEQ ID NO: 90
322	1364 Sel1 repeat :: PF08238		SEQ ID NO: 23	SEQ ID NO: 24
305	1524 haloacid dehalogenase-like hydrolase :: PF00702		SEQ ID NO: 98	SEQ ID NO: 101
308	769 EAL domain :: PF00563		SEQ ID NO: 99	SEQ ID NO: 102
311	1847 MotA/TolQ/ExbB proton channel family :: PF01618			

the recombinant purified proteins were prepared. 12 Sel1 proteins and 7 additional Of-derived factors were purified (Table 2) and the effects of different concentrations and incubation periods of the purified proteins on oxalate transport by C2 cells were evaluated. Experiments conducted during development of embodiments herein demonstrate that Sel1 proteins (# s 194, 1356, 1361, and 1362), significantly stimulate (up to 2.5-fold) oxalate transport by C2 cells. In addition to the Sel1 proteins, LD-carboxypeptidase (#8), a peptidase (#1525), haloacid dehalogenase-like hydrolase (1524=305), and guanylate cyclase (769=308) also stimulated (~1.4-1.6-fold) oxalate transport by C2 in

[0053] Experiments conducted during development of embodiments herein demonstrate the capacity for Of-derived media, Of-derived factors, SLR proteins (e.g., Sel1 proteins), SLR peptides (e.g., Sel1 peptides), SLR variant polypeptides (e.g., Sel1 variant polypeptides), etc. to reduce oxalate concentrations and thereby treat and/or prevent conditions related to excess oxalate in the blood, urine, etc.

[0054] In some embodiments, Of-derived factors that find use in embodiments herein are derived from any suitable species or strain of *Oxalobacter formigenes*, such as, for example, HC-1, Va3, OxK, OxB, OXCC13, BA1, HOxBLs, HOxRW, POxC, etc. In some embodiments, variants, frag-

ments, and/or peptidomimetics of bioactive factors derived from suitable Of strains are provided.

[0055] In some embodiments, media conditioned by the growth of *Oxalobacter formigenes* (CM) is provided. In some embodiments, Of CM is provided in a pharmaceutical composition for the treatment/prevention of conditions related to excess oxalate. In some embodiments, Of CM is processed (e.g., fractionated, purified, concentrated, diluted, filtered, etc.) prior to administration to a subject. In some embodiments, Of CM is formulated for administration by any suitable techniques described herein or known in the field.

[0056] In some embodiments, bioactive Of-derived factors are provided for the treatment/prevention of conditions related to excess oxalate. In some embodiments, such factors are obtained, isolated, and/or purified from Of CM. In some embodiments, such factors are obtained, isolated, and/or purified from Of cultures. In some embodiments, are prepared recombinantly and/or synthetically. In some embodiments, bioactive Of-derived factors are SLR proteins (e.g., Sell proteins), such as Sell, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, MerG, and variants, fragment, and peptidomimetics thereof.

[0057] Provided herein are compositions (e.g., Of conditioned media, Of-derived factors, SLR proteins (e.g., Sell proteins), SLR peptides (e.g., Sell peptides), SLR variant polypeptides (e.g., Sell variant polypeptides), etc.) which stimulate the clearance of oxalate (e.g., activate oxalate transport) from a biological environment (e.g., blood, urine, etc.). In some embodiments, compositions significantly reduce oxalate concentrations (e.g., urine oxalate, blood oxalate, etc.) for example, by at least 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or ranges therebetween. In some embodiments, compositions herein stimulate oxalate transport, thereby reducing in vivo oxalate levels in the blood (e.g., plasma oxalate levels), urine, etc., through mechanisms such as, for example, PKA activation and increased activity of SLC26 family members (e.g., SLC26A6) or other transporter(s); although embodiments herein are not limited to any particular mechanism of action and an understanding of the mechanism of action is not necessary to practice such embodiments.

[0058] In some embodiments, provided herein are compositions, kits, systems, and/or methods to treat, prevent, reduce the likelihood, treat/prevent a side effect of one or more of: hyperoxalemia, hyperoxaluria, nephrolithiasis, chronic kidney disease, end stage renal disease, calcium oxalate kidney stones, nephrocalcinosis, oxalate nephropathy, primary hyperoxaluria (PH), enteric hyperoxaluria (seen for example in IBD, following small bowel surgery or bariatric surgery, obesity, and celiac disease) and systemic oxalosis. In some embodiments, the reduction in oxalate levels and/or activation of oxalate transport is activated by compositions and methods described herein. In some embodiments, oxalate transport pathways are activated by the compositions and methods described herein. In some embodiments, compositions and methods are utilized in the treatment and/or prevention of hyperoxalemia, hyperoxaluria, and/or related diseases and conditions. In some embodiments, compositions and methods are utilized in screening for peptides and polypeptides useful in the treatment and/or prevention of hyperoxalemia, hyperoxaluria, and/or related diseases and conditions.

[0059] In some embodiments, provided herein are pharmaceutical compositions, Of CM, Of-derived factors, SLR peptides (e.g., Sell peptides), SLR proteins (e.g., Sell proteins), SLR polypeptides (e.g., Sell polypeptides), nucleic acids encoding peptides, proteins and polypeptides, molecular complexes of the foregoing, etc. for the treatment or prevention of hyperoxalemia, hyperoxaluria, and/or related diseases and conditions. In some embodiments, provided herein are SLR-peptides and SLR-polypeptides (e.g., comprising less than 100% sequence identity with full length native SLR proteins (e.g., SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, Sell, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, MerG), or a fragment of full length SLR protein or variant thereof. In some embodiments, a peptide/polypeptide is artificial. In some embodiments, a polypeptide or peptide described herein is prepared by methods known to those of ordinary skill in the art. For example, the peptide or polypeptide can be synthesized using solid phase polypeptide synthesis techniques (e.g. Fmoc or Boc chemistry). Alternatively, the peptide or polypeptide can be produced using recombinant DNA technology (e.g., using bacterial or eukaryotic expression systems). Further, a peptide or polypeptide may be expressed within a subject (e.g., following administration of an appropriate vector). Accordingly, to facilitate such methods, provided herein are genetic vectors (e.g., plasmids, viral vectors (e.g. AAV), etc.) comprising a sequence encoding the polypeptide, as well as host cells comprising such vectors. Furthermore, provided herein are the peptides and polypeptides produced via such methods.

[0060] In some embodiments, the administration of Of-derived factors (e.g., peptides and polypeptides) and compositions related thereto (e.g. variants and mimetics of Of-derived factors, nucleic acids encoding Of-derived factors, etc.) is provided. In some embodiments, provided herein is the administration of bioactive agents which reduce oxalate levels in vivo, or are otherwise described herein. Examples of such peptides and polypeptides include those selected from the group consisting of SEQ ID NOS: 2, 4, 6, 8, 10, 12, 14, 16, 90, 92, 94, 96, 101, and 102. Other examples include SLR proteins (e.g., Sell, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, MerG) derived from various Of strains, and variants and fragments thereof.

[0061] In some embodiments, a peptide or polypeptide is provided comprising or consisting of all or a portion of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, and SEQ ID NO: 102. In some embodiments, a peptide or polypeptide is provided comprising at least 50% sequence identity to one of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, and SEQ ID NO: 102 (e.g. at least 60% sequence identity, at least 70% sequence identity, at least 80% sequence identity, at least 90% sequence identity, at least 95% sequence identity, etc.). In some embodiments, peptide and polypeptides comprise at least one mutation from a wild-type sequence (e.g., SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID

NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, and SEQ ID NO: 102).

[0062] In some embodiments, a peptide/polypeptide is provided that is a fragment of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, and SEQ ID NO: 102. In some embodiments, a peptide/polypeptide is provided that comprises one or more substitutions (e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, or more) compared to a fragment of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, and SEQ ID NO: 102.

[0063] In some embodiments, a peptide or polypeptide is provided comprising or consisting of all or a portion of an Of-derived SLR protein (e.g., Sel1 protein) that facilitates reduction of in vivo oxalate levels, such as an SLR protein (e.g., Sel1, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, MerG) derived from an Of strains, and variants and fragments thereof. In some embodiments, a peptide or polypeptide is provided comprising at least 50% sequence identity to an SLR protein (e.g., Sel1, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, MerG) derived from an Of strains (e.g. at least 60% sequence identity, at least 70% sequence identity, at least 80% sequence identity, at least 90% sequence identity, at least 95% sequence identity, etc.). In some embodiments, peptide and polypeptides comprise at least one mutation from a wild-type sequence (e.g., Sel1, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, MerG).

[0064] In some embodiments, a peptide/polypeptide is provided that is a fragment of an SLR protein (e.g., Sel1, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, MerG) derived from an Of strain. In some embodiments, a peptide/polypeptide is provided that comprises one or more substitutions (e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, or more) compared to a fragment of an SLR protein (e.g., Sel1, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, MerG) derived from an Of strain.

[0065] Embodiments are not limited to the specific sequences listed herein. In some embodiments, peptides/polypeptides meeting limitations described herein (e.g., Of-derived, reduce oxalate in vivo, biostable, bioavailable, biocompatible, etc.), and having substitutions not explicitly described are within the scope of embodiments here. In some embodiments, the peptides/polypeptides described herein are further modified (e.g., substitution, deletion, or addition of standard amino acids; chemical modification; etc.). Modifications that are understood in the field include N-terminal modification, C-terminal modification (which protects the peptide from proteolytic degradation), alkylation of amide groups, hydrocarbon "stapling" (e.g., to stabilize conformations). In some embodiments, the peptides/polypeptides described herein may be modified by conservative residue substitutions, for example, of the charged residues (K to R, R to K, D to E and E to D). Modifications of the terminal carboxy group include, without limitation, the amide, lower alkyl amide, constrained

alkyls (e.g. branched, cyclic, fused, adamantyl) alkyl, dialkyl amide, and lower alkyl ester modifications. Lower alkyl is C1-C4 alkyl. Furthermore, one or more side groups, or terminal groups, may be protected by protective groups known to the ordinarily-skilled peptide chemist. The α -carbon of an amino acid may be mono- or dimethylated.

[0066] In some embodiments, any embodiments described herein may comprise mimetics corresponding to all or a portion of the Of-derived factors and/or SLR proteins (e.g., Sel1 proteins) described herein, with various modifications that are understood in the field. In some embodiments, residues in the peptide sequences described herein may be substituted with amino acids having similar characteristics (e.g., hydrophobic to hydrophobic, neutral to neutral, etc.) or having other desired characteristics (e.g., more acidic, more hydrophobic, less bulky, more bulky, etc.). In some embodiments, non-natural amino acids (or naturally-occurring amino acids other than the standard 20 amino acids) are substituted in order to achieve desired properties.

[0067] In some embodiments, residues having a side chain that is positively charged under physiological conditions, or residues where a positively-charged side chain is desired, are substituted with a residue including, but not limited to: lysine, homolysine, δ -hydroxylysine, homoarginine, 2,4-diaminobutyric acid, 3-homoarginine, D-arginine, arginal (—COOH in arginine is replaced by —CHO), 2-amino-3-guanidinopropionic acid, nitroarginine (N(G)-nitroarginine), nitrosoarginine (N(G)-nitrosoarginine), methylarginine (N-methyl-arginine), ϵ -N-methyllysine, allo-hydroxylysine, 2,3-diaminopropionic acid, 2,2'-diaminopimelic acid, ornithine, sym-dimethylarginine, asym-dimethyl arginine, 2,6-diaminohexinic acid, p-aminobenzoic acid and 3-aminotyrosine and, histidine, 1-methylhistidine, and 3-methylhistidine.

[0068] A neutral residue is a residue having a side chain that is uncharged under physiological conditions. A polar residue preferably has at least one polar group in the side chain. In some embodiments, polar groups are selected from hydroxyl, sulfhydryl, amine, amide and ester groups or other groups which permit the formation of hydrogen bridges.

[0069] In some embodiments, residues having a side chain that is neutral/polar under physiological conditions, or residues where a neutral side chain is desired, are substituted with a residue including, but not limited to: asparagine, cysteine, glutamine, serine, threonine, tyrosine, citrulline, N-methylserine, homoserine, allo-threonine and 3,5-dinitrotyrosine, and β -homoserine.

[0070] Residues having a non-polar, hydrophobic side chain are residues that are uncharged under physiological conditions, preferably with a hydrophathy index above 0, particularly above 3. In some embodiments, non-polar, hydrophobic side chains are selected from alkyl, alkylene, alkoxy, alkenoxy, alkylsulfanyl and alkenylsulfanyl residues having from 1 to 10, preferably from 2 to 6, carbon atoms, or aryl residues having from 5 to 12 carbon atoms. In some embodiments, residues having a non-polar, hydrophobic side chain are, or residues where a non-polar, hydrophobic side chain is desired, are substituted with a residue including, but not limited to: leucine, isoleucine, valine, methionine, alanine, phenylalanine, N-methylleucine, tert-butylglycine, octylglycine, cyclohexylalanine, β -alanine, 1-aminocyclohexylcarboxylic acid, N-methylisoleucine, norleucine, norvaline, and N-methylvaline.

[0071] In some embodiments, peptide and polypeptides are isolated and/or purified (or substantially isolated and/or substantially purified). Accordingly, in such embodiments, peptides and/or polypeptides are provided in substantially isolated form. In some embodiments, peptides and/or polypeptides are isolated from other peptides and/or polypeptides as a result of solid phase peptide synthesis, for example. Alternatively, peptides and/or polypeptides can be substantially isolated from other proteins after cell lysis from recombinant production. Standard methods of protein purification (e.g., HPLC) can be employed to substantially purify peptides and/or polypeptides. In some embodiments, the present invention provides a preparation of peptides and/or polypeptides in a number of formulations, depending on the desired use. For example, where the polypeptide is substantially isolated (or even nearly completely isolated from other proteins), it can be formulated in a suitable medium solution for storage (e.g., under refrigerated conditions or under frozen conditions). Such preparations may contain protective agents, such as buffers, preservatives, cryoprotectants (e.g., sugars such as trehalose), etc. The form of such preparations can be solutions, gels, etc. In some embodiments, peptides and/or polypeptides are prepared in lyophilized form. Moreover, such preparations can include other desired agents, such as small molecules or other peptides, polypeptides or proteins. Indeed, such a preparation comprising a mixture of different embodiments of the peptides and/or polypeptides described here may be provided.

[0072] In some embodiments, provided herein are peptidomimetic versions of the peptide sequences described herein or variants thereof. In some embodiments, a peptidomimetic is characterized by an entity that retains the polarity (or non-polarity, hydrophobicity, etc.), three-dimensional size, and functionality (bioactivity) of its peptide equivalent but wherein all or a portion of the peptide bonds have been replaced (e.g., by more stable linkages). In some embodiments, 'stable' refers to being more resistant to chemical degradation or enzymatic degradation by hydrolytic enzymes. In some embodiments, the bond which replaces the amide bond (e.g., amide bond surrogate) conserves some properties of the amide bond (e.g., conformation, steric bulk, electrostatic character, capacity for hydrogen bonding, etc.). Chapter 14 of "Drug Design and Development", Krogsgaard, Larsen, Liljefors and Madsen (Eds) 1996, Horwood Acad. Publishers provides a general discussion of techniques for the design and synthesis of peptidomimetics and is herein incorporated by reference in its entirety. Suitable amide bond surrogates include, but are not limited to: N-alkylation (Schmidt, R. et al., *Int. J. Peptide Protein Res.*, 1995, 46,47; herein incorporated by reference in its entirety), retro-inverse amide (Chorev, M. and Goodman, M., *Acc. Chem. Res.*, 1993, 26, 266; herein incorporated by reference in its entirety), thioamide (Sherman D. B. and Spatola, A. F. *J. Am. Chem. Soc.*, 1990, 112, 433; herein incorporated by reference in its entirety), thioester, phosphonate, ketomethylene (Hoffman, R. V. and Kim, H. O. *J. Org. Chem.*, 1995, 60, 5107; herein incorporated by reference in its entirety), hydroxymethylene, fluorovinyl (Allmendinger, T. et al., *Tetrahydron Lett.*, 1990, 31, 7297; herein incorporated by reference in its entirety), vinyl, methyleneamino (Sasaki, Y and Abe, *J. Chem. Pharm. Bull.* 1997 45, 13; herein incorporated by reference in its entirety), methylenethio (Spatola, A. F., *Methods Neurosci*, 1993, 13, 19; herein incorporated

by reference in its entirety), alkane (Lavielle, S. et. al., *Int. J. Peptide Protein Res.*, 1993, 42, 270; herein incorporated by reference in its entirety) and sulfonamido (Luisi, G. et al. *Tetrahydron Lett.* 1993, 34, 2391; herein incorporated by reference in its entirety).

[0073] As well as replacement of amide bonds, peptidomimetics may involve the replacement of larger structural moieties with di- or tripeptidomimetic structures and in this case, mimetic moieties involving the peptide bond, such as azole-derived mimetics may be used as dipeptide replacements. Suitable peptidomimetics include reduced peptides where the amide bond has been reduced to a methylene amine by treatment with a reducing agent (e.g. borane or a hydride reagent such as lithium aluminum-hydride); such a reduction has the added advantage of increasing the overall cationicity of the molecule.

[0074] Other peptidomimetics include peptoids formed, for example, by the stepwise synthesis of amide-functionalised polyglycines. Some peptidomimetic backbones will be readily available from their peptide precursors, such as peptides which have been permethylated, suitable methods are described by Ostresh, J. M. et al. in *Proc. Natl. Acad. Sci. USA* (1994) 91, 11138-11142; herein incorporated by reference in its entirety.

[0075] In some embodiments, the peptides/polypeptides described herein are provided as fusions with other peptides or polypeptides. Such fusions may be expressed from a recombinant DNA which encodes the SLR and/or Of-derived peptide/polypeptide and the additional peptide/polypeptide or may be formed by chemical synthesis. For instance, the fusion may comprise a SLR and/or Of-derived peptide/polypeptide and an enzyme of interest, a luciferase, RNasin or RNase, and/or a channel protein (e.g., ion channel protein), a receptor, a membrane protein, a cytosolic protein, a nuclear protein, a structural protein, a phosphoprotein, a kinase, a signaling protein, a metabolic protein, a mitochondrial protein, a receptor associated protein, a fluorescent protein, an enzyme substrate, a transcription factor, selectable marker protein, nucleic acid binding protein, extracellular matrix protein, secreted protein, receptor ligand, serum protein, a protein with reactive cysteines, a transporter protein, a targeting sequence (e.g., a myristylation sequence), a mitochondrial localization sequence, or a nuclear localization sequence. The additional peptide/polypeptide may be fused to the N-terminus and/or the C-terminus of the SLR and/or Of-derived peptide/polypeptide. In one embodiment, the fusion protein comprises a first peptide/polypeptide at the N-terminus and another (different) peptide/polypeptide at the C-terminus of the SLR and/or Of-derived peptide/polypeptide. Optionally, the elements in the fusion are separated by a connector sequence, e.g., preferably one having at least 2 amino acid residues, such as one having 13 and up to 40 or 50 amino acid residues. The presence of a connector sequence in a fusion protein of the invention does not substantially alter the function of either element (e.g., the SLR and/or Of-derived peptide/polypeptide) in the fusion relative to the function of each individual element, likely due to the connector sequence providing flexibility (autonomy) for each element in the fusion. In certain embodiment, the connector sequence is a sequence recognized by an enzyme or is photocleavable. For example, the connector sequence may include a protease recognition site.

[0076] In some embodiments, provided herein are pharmaceutical compositions comprising of one or more SLR and/or Of-derived peptide/polypeptide described herein and a pharmaceutically acceptable carrier. Any carrier which can supply an active peptide or polypeptide (e.g., without destroying the peptide or polypeptide within the carrier) is a suitable carrier, and such carriers are well known in the art. In some embodiments, compositions are formulated for administration by any suitable route, including but not limited to, orally (e.g., such as in the form of tablets, capsules, granules or powders), sublingually, buccally, parenterally (such as by subcutaneous, intravenous, intramuscular, intradermal, or intracisternal injection or infusion (e.g., as sterile injectable aqueous or non-aqueous solutions or suspensions, etc.)), nasally (including administration to the nasal membranes, such as by inhalation spray), topically (such as in the form of a cream or ointment), transdermally (such as by transdermal patch), rectally (such as in the form of suppositories), etc.

[0077] In some embodiments, provided herein are methods for treating patients suffering from (or at risk of) hyperoxaluria, hyperoxalemia, and/or in need of treatment (or preventative therapy). In some embodiments, a pharmaceutical composition comprising at least one Of-derived and/or SLR peptide/polypeptide described herein is delivered to such a patient in an amount and at a location sufficient to treat the condition. In some embodiments, peptides and/or polypeptides (or pharmaceutical composition comprising such) can be delivered to the patient systemically or locally, and it will be within the ordinary skill of the medical professional treating such patient to ascertain the most appropriate delivery route, time course, and dosage for treatment. It will be appreciated that application methods of treating a patient most preferably substantially alleviates or even eliminates such symptoms; however, as with many medical treatments, application of the inventive method is deemed successful if, during, following, or otherwise as a result of the inventive method, the symptoms of the disease or disorder in the patient subside to an ascertainable degree.

[0078] A pharmaceutical composition may be administered in the form which is formulated with a pharmaceutically acceptable carrier and optional excipients, adjuvants, etc. in accordance with good pharmaceutical practice. The Of-derived and/or SLR peptide/polypeptide pharmaceutical composition may be in the form of a solid, semi-solid or liquid dosage form: such as powder, solution, elixir, syrup, suspension, cream, drops, paste and spray. As those skilled in the art would recognize, depending on the chosen route of administration (e.g. pill, injection, etc.), the composition form is determined. In general, it is preferred to use a unit dosage form in order to achieve an easy and accurate administration of the active pharmaceutical peptide or polypeptide. In general, the therapeutically effective pharmaceutical compound is present in such a dosage form at a concentration level ranging from about 0.5% to about 99% by weight of the total composition, e.g., in an amount sufficient to provide the desired unit dose. In some embodiments, the pharmaceutical composition may be administered in single or multiple doses. The particular route of administration and the dosage regimen will be determined by one of skill in keeping with the condition of the individual to be treated and said individual's response to the treatment. In some embodiments, an Of-derived and/or SLR peptide/polypeptide pharmaceutical composition is provided in a

unit dosage form for administration to a subject, comprising one or more nontoxic pharmaceutically acceptable carriers, adjuvants or vehicles. The amount of the active ingredient that may be combined with such materials to produce a single dosage form will vary depending upon various factors, as indicated above. A variety of materials can be used as carriers, adjuvants and vehicles in the composition of the invention, as available in the pharmaceutical art. Injectable preparations, such as oleaginous solutions, suspensions or emulsions, may be formulated as known in the art, using suitable dispersing or wetting agents and suspending agents, as needed. The sterile injectable preparation may employ a nontoxic parenterally acceptable diluent or solvent such as sterile nonpyrogenic water or 1,3-butanediol. Among the other acceptable vehicles and solvents that may be employed are 5% dextrose injection, Ringer's injection and isotonic sodium chloride injection (as described in the USP/NF). In addition, sterile, fixed oils may be conventionally employed as solvents or suspending media. For this purpose, any bland fixed oil may be used, including synthetic mono-, di- or triglycerides. Fatty acids such as oleic acid can also be used in the preparation of injectable compositions.

[0079] In various embodiments, the peptides and polypeptides disclosed herein are derivatized by conjugation to one or more polymers or small molecule substituents.

[0080] In certain of these embodiments, the peptides and polypeptides described herein are derivatized by coupling to polyethylene glycol (PEG). Coupling may be performed using known processes. See, *Int. J. Hematology*, 68:1 (1998); *Bioconjugate Chem.*, 6:150 (1995); and *Crit. Rev. Therap. Drug Carrier Sys.*, 9:249 (1992) all of which are incorporated herein by reference in their entirety. Those skilled in the art, therefore, will be able to utilize such well-known techniques for linking one or more polyethylene glycol polymers to the peptides and polypeptides described herein. Suitable polyethylene glycol polymers typically are commercially available or may be made by techniques well known to those skilled in the art. The polyethylene glycol polymers preferably have molecular weights between 500 and 20,000 and may be branched or straight chain polymers.

[0081] The attachment of a PEG to a peptide or polypeptide described herein can be accomplished by coupling to amino, carboxyl or thiol groups. These groups will typically be the N- and C-termini and on the side chains of such naturally occurring amino acids as lysine, aspartic acid, glutamic acid and cysteine. Since the peptides and polypeptides of the present disclosure can be prepared by solid phase peptide chemistry techniques, a variety of moieties containing diamino and dicarboxylic groups with orthogonal protecting groups can be introduced for conjugation to PEG.

[0082] The present disclosure also provides for conjugation of the peptides and polypeptides described herein to one or more polymers other than polyethylene glycol.

[0083] In some embodiments, the peptides and polypeptides described herein are derivatized by conjugation or linkage to, or attachment of, polyamino acids (e.g., poly-his, poly-arg, poly-lys, etc.) and/or fatty acid chains of various lengths to the N- or C-terminus or amino acid residue side chains. In certain embodiments, the peptides and polypeptides described herein are derivatized by the addition of polyamide chains, particularly polyamide chains of precise lengths, as described in U.S. Pat. No. 6,552,167, which is incorporated by reference in its entirety. In yet other embodiments, the peptides and polypeptides are modified by the

addition of alkylPEG moieties as described in U.S. Pat. Nos. 5,359,030 and 5,681,811, which are incorporated by reference in their entireties.

[0084] In select embodiments, the peptides and polypeptides disclosed herein are derivatized by conjugation to polymers that include albumin and gelatin. See, Gombotz and Pettit, *Bioconjugate Chem.*, 6:332-351, 1995, which is incorporated herein by reference in its entirety.

[0085] In further embodiments, the peptides and polypeptides disclosed herein are conjugated or fused to immunoglobulins or immunoglobulin fragments, such as antibody Fc regions.

[0086] In some embodiments, the pharmaceutical compositions described herein (e.g., comprising SLR proteins (e.g., Sell proteins), Of-derived factors, and/or variants and fragments thereof) find use in the treatment and/or prevention of hyperoxaluria, hyperoxalemia, and related conditions. In some embodiments, the compositions are administered to a subject. In certain embodiments, the patient is an adult. In other embodiments, the patient is a child.

[0087] In various embodiments, the peptide/polypeptide is administered in an amount, on a schedule, and for a duration sufficient to decrease triglyceride levels by at least 5%, 10%, 15%, 20% or 25% or more as compared to levels just prior to initiation of treatment. In some embodiments, the peptide/polypeptide is administered in an amount, on a dosage schedule, and for a duration sufficient to decrease oxalate levels (e.g., in urine, in plasma) by at least 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45% or 50%. In particular embodiments, the peptide/polypeptide is administered in an amount, on a schedule, and for a time sufficient to decrease oxalate levels (e.g., in urine, in plasma) by at least 55%, 60%, 65%, even at least about 70% or more.

[0088] In certain embodiments, the peptide/polypeptide is administered in an amount, expressed as a daily equivalent dose regardless of dosing frequency, of 50 micrograms ("mcg") per day, 60 mcg per day, 70 mcg per day, 75 mcg per day, 100 mcg per day, 150 mcg per day, 200 mcg per day, or 250 mcg per day. In some embodiments, the polypeptide is administered in an amount of 500 mcg per day, 750 mcg per day, or 1 milligram ("mg") per day. In yet further embodiments, the polypeptide is administered in an amount, expressed as a daily equivalent dose regardless of dosing frequency, of 1-10 mg per day, including 1 mg per day, 1.5 mg per day, 1.75 mg per day, 2 mg per day, 2.5 mg per day, 3 mg per day, 3.5 mg per day, 4 mg per day, 4.5 mg per day, 5 mg per day, 5.5 mg per day, 6 mg per day, 6.5 mg per day, 7 mg per day, 7.5 mg per day, 8 mg per day, 8.5 mg per day, 9 mg per day, 9.5 mg per day, or 10 mg per day.

[0089] In various embodiments, the peptide/polypeptide is administered on a monthly dosage schedule. In other embodiments, the peptide/polypeptide is administered biweekly. In yet other embodiments, the polypeptide is administered weekly. In certain embodiments, the peptide/polypeptide is administered daily ("QD"). In select embodiments, the polypeptide is administered twice a day ("BID").

[0090] In typical embodiments, the peptide/polypeptide is administered for at least 3 months, at least 6 months, at least 12 months, or more. In some embodiments, the peptide/polypeptide is administered for at least 18 months, 2 years, 3 years, or more.

EXPERIMENTAL

Example 1

[0091] Of interacts with colonic epithelium and induces distal colonic oxalate secretion, leading to reduced urinary excretion (ref. 14; incorporated by reference in its entirety). Of whole cells, cell membranes, and lysates have been tested on oxalate transport across rat distal colonic tissues mounted in Ussing chambers, and were found to have no effect (ref. 14; incorporated by reference in its entirety). Both the whole cells and lysates caused significant degradation of oxalate in the chamber, necessitating heat treatment of samples to eradicate this inherent enzymatic activity. To further understand this phenomenon, Of was obtained from ATCC (strain OxB) and was grown in an anaerobic chamber (ref. 2; incorporated by reference in its entirety). Of cultures were centrifuged (3,000 g at 4° C. for 10 min) and the supernatant (conditioned medium=CM) was filtered through a 0.22 µm filter to sterilize and remove all bacterial cells and stored at -80° C. (ref. 5; incorporated by reference in its entirety). C2 cells were used as a model to evaluate the effects of Of CM on intestinal oxalate transport. Apical ¹⁴C-oxalate flux studies in C2 cells were performed. Apical oxalate uptake by C2 cells was assessed by imposing an outward Cl gradient by removing extracellular Cl [Cl_i>Cl_o] and measuring DIDS (anion exchange inhibitor)-sensitive influx of radioactive ¹⁴C-oxalate in exchange for intracellular Cl [i.e. apical Cl-oxalate exchange activity, ≥50% of which is mediated by A6 in C2 cells (ref. 12; incorporated by reference in its entirety)]. A6 operates in the direction of exchanging intracellular oxalate for mucosal Cl during the process of transepithelial intestinal oxalate secretion. However, A6 can operate in either direction (ref. 21; incorporated by reference in its entirety), and therefore its activity was measured by the more convenient assay of cellular oxalate uptake. C2 cells grown on TRANSWELLS were treated apically with Of culture medium (OM) or CM (1:50 dilution×24 h) before measuring ¹⁴C-oxalate uptake.

[0092] Compared to untreated (UT) and OM, the CM significantly stimulated (>2.4-fold) oxalate transport by C2 cells (ref 4; incorporated by reference in its entirety). Similar effects were also observed with 6- and 16-h incubations, but no effect was seen at 1 h. CM or OM did not affect the medium pH. In addition, OM or CM had no significant effect on the transepithelial resistance, indicating that the OM or the CM does not affect the paracellular permeability. These results indicate that secreted bioactive factors in Of CM are responsible for the observed stimulatory regulation by modulating the activity of the involved anion exchanger(s) (A1, A2, and/or A6). *Lactobacillus acidophilus* (La) degrades intraluminal oxalate (19), but it is unknown whether La similarly interacts with enterocytes and modulates intestinal oxalate transport as Of (ref. 14; incorporated by reference in its entirety). Therefore, to ensure specificity, experiments similarly evaluated the effect of La CM. It was observed that La CM (1:25 dilution×24 h) had no effect on oxalate transport by C2 cells compared to UT cells and cells treated with the control medium (ref. 4; incorporated by reference in its entirety). These results indicate that Of CM-induced stimulation of oxalate transport is specific and is possibly mediated by one or more of Of-derived secreted bioactive factor(s).

[0093] To have an idea about the nature of Of-derived secreted bioactive factor(s), CM was subjected to heat

treatment (boiling at 100° C. for 20-30 min). Heat treatment completely abolished the CM-induced stimulation, indicating that the secreted factor(s) are likely to be protein(s) or peptide(s), rather than, for example, small molecules. Pretreatment of the CM with pepsin or trypsin also completely abolished the stimulatory effect, providing further evidence that the secreted factor(s) is/are proteins or peptides. Selective ultrafiltration revealed that the secreted factors have molecular masses (MM) between 10-30 kDa (ref. 4; incorporated by reference in its entirety). Pretreatment with the PKA inhibitor H89 completely blocked the CM-induced stimulation of oxalate transport by C2 cells, indicating that the observed stimulation is mediated by PKA activation. The observed stimulation is also completely blocked by DIDDS, indicating that it is due to active anion exchange-mediated transport process. siRNA A6 knockdown greatly reduced the observed stimulation, indicating that a significant component of CM-induced oxalate transport is A6-mediated. The CM significantly increased (>2-fold) the V_{max} (e.g., greater transport capacity) and reduced (>3.4-fold) the K_m (e.g., greater affinity for oxalate) of the involved transporter(s). The CM did not affect A6 mRNA and total/surface protein expression, and in view of the reduced K_m (reflecting greater A6 affinity for oxalate), indicating that the observed stimulation is due to mechanisms including CM-induced enhanced A6 transport activity (e.g., resulting from an increase in the intrinsic activity of the preexisting A6 membrane transporters), although embodiments herein are not limited to any particular mechanism of action and an understanding of the mechanism of action is not necessary to practice such embodiments.

[0094] To evaluate the in vivo effects of the CM on overall oxalate homeostasis, CM or OM was given rectally as enemas (100 μ l twice daily \times 21 days) to PH1 mice (a model of PH type 1 due to deficiency in the liver enzyme AGAT and they have significant hyperoxalemia and hyperoxaluria) (ref. 28; incorporated by reference in its entirety). It was confirmed that the PH1 mice have significant hyperoxaluria (>2.7-fold) compared to their controls. The CM significantly reduced urinary oxalate excretion by >32.5%, while the OM had no significant effect (ref. 4; incorporated by reference in its entirety). To determine whether the Of-derived bioactive factor(s) interact(s) with the colonocyte and induce(s) oxalate secretion in vivo, leading to the observed reduction in urinary oxalate excretion in PH1 mice, distal colonic tissues were isolated and mounted in Ussing chambers at the end of the treatment period. While a small net oxalate secretory flux (-4.7 \pm 2.4) was observed in distal colonic tissues from OM-treated PH1 mice, a >4.2-fold higher net oxalate secretory flux (-20.0 \pm 5.5) was seen in distal colonic tissues from CM-treated PH1 mice (pmol/cm²/h: OM: J_{MS} (absorptive flux)=37.8 \pm 5.9, J_{SM} (secretory flux)=42.6 \pm 5.5; CM: J_{MS} =40.6 \pm 3.7, J_{SM} =60.6 \pm 3.1), which is due to significantly increased (>42%) J_{SM} (4). CM or OM had no significant effect on J_{MS} . These results indicate that the Of-derived factors reduce urinary oxalate excretion in PH1 mice through mechanisms including enhanced net distal colonic oxalate secretion, as well as they retain their biological activity in vivo, thus indicating their significant therapeutic value.

[0095] An important factor in purifying the bioactive protein(s) is their presence in a high concentration in the CM. Since Of might secrete these factors as a survival strategy when oxalate is limited (ref. 14; incorporated by

reference in its entirety), it was tested whether lower growth medium oxalate concentration lead to more secreted factor(s), and therefore, to a CM with higher bioactivity, by reducing oxalate concentration from 37.5 mM to 18.8 & 9.4 mM. The bioactivity of the CM is significantly higher (>2-fold) with CM9.4 compared to CM37.5 (ref. 4; incorporated by reference in its entirety), demonstrating that the secretion of these factors is inducible, which facilitates their characterization. As a first step in characterizing these factors, secreted proteins in Of CM were purified utilizing column chromatography, using a commercially available kit for rapid screening of a suitable column. 1 ml of the CM was loaded onto different columns (e.g. anion and cation exchange columns) and the flow-through fractions were collected and their effects on oxalate influx into C2 cells were evaluated. The fractions from the weak cationic column (WCC) were found to have no stimulatory effect compared with the parent CM, indicating that this is due to sticking of the potential factor(s) to this column. The factor(s) were then eluted from this column using high salt (0.5-1 M NaCl). Following desalting and concentration of the eluted fraction using a desalting/concentrating column, the eluted fraction is found to have an activity similar to the parent CM (pmoles/cm²/min: UT=1.56 \pm 0.30; CM=8.47 \pm 0.86; Eluted fraction=7.36 \pm 0.69). These findings indicate that the purification process has led to enrichment of the factor(s) in the eluted fraction and the WCC was utilized in subsequent purification studies using FPLC.

[0096] The experiments demonstrate that small molecular weight protein(s) and/or peptide(s) in Of CM significantly stimulate(s) oxalate transport (>2.4-fold) by human intestinal Caco2-BBE cells (FIG. 1, FIG. 2, and FIG. 4). Importantly, Of CM also significantly reduced (>32.5%) urinary oxalate excretion in a mouse model of primary hyperoxaluria (PH1), by stimulating (>42%) distal colonic oxalate secretion, reflecting the in vivo retention of biologic activity and the therapeutic value of these factors.

[0097] The high specific activity elution fraction was run in SDS-PAGE and then stained by Coomassie blue. Several bands in the molecular weight range from 10-30 kDa, as well as bands ~52 and 68 kDa were noted. Tandem mass spectrometry was utilized as a primary strategy to identify candidates for the mediators. The gel sections containing the bands were cut and in-gel trypsin proteolysis was performed. Isolated peptides were subjected to LTQ Orbitrap ESI LC-MS/MS analysis, using standard conditions. The data were analyzed by Mascot and X! Tandem database search against the predicted Of proteome, combined with a reverse decoy database to estimate false discovery rate. The results were then validated and visualized using Scaffold. On the basis of 16S rRNA sequence similarities and lipid content, Of strains are divided into two groups: Group I is represented by strain OxB (the strain used in the above experiments) and human strain OxCC13, while Group II is represented by human strain HOxBLS. OXCC13 and HOxBLS genomes were published. Searching against the OXCC13 genome yielded 52 candidates Of protein CM stimulatory factors. A more stringent search (Maxquant: 20 ppm, with additional filtering at 1% FDR) yielded 6 candidate proteins which are the following: transcriptional regulator, LuxR family (ID # C3X886), Raf-like protein (ID # C3XB51), Tryptophan synthase alpha chain (ID # C3XAU0), Sell repeat protein (C3X8T9), Uncharacterized protein (ID C3X964), and Uncharacterized protein (ID #

C3XC9Y9). CM from OXCC13 is not commercially available to be cultured. Therefore, the OxB whole genome was sequenced and assembled. The data generated by Mass-Spect was re-analyzed against the OxB genome and the amino acid sequences for the corresponding proteins identified above by searching the OXCC13 genome were obtained.

Example 2

[0098] Several of the identified Of proteins were of interest, especially the Sell proteins. Sell-like repeat (SLR) proteins (e.g. Sell1, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, and MerG) are involved in signal transduction pathways (ref 24; incorporated by reference in its entirety). It was found that Of CM signals through PKA to stimulate oxalate transport by C2 cells. SLR proteins have repeat units and most repeats consist of 5 to 40 amino acids (but some could be much larger) that fold into two to four secondary structural elements. All SLR proteins seem to serve as adaptor proteins for the assembly of membrane-bound macromolecular complexes (ref. 24; incorporated by reference in its entirety). Several bacterial and eukaryotic SLR proteins (e.g. Sell1 & Hrd3) are activated upon cellular stress, which is of interest since Of Sell1 proteins might be activated when oxalate is low in the culture medium given the observation of a CM of higher (>2-fold) bioactivity under this condition. Bacterial LpnE, EnhC, HcpA, ExoR, and AlgK proteins mediate the interactions between bacterial and eukaryotic host cells. The SLR motif establishes a link between signal transduction pathways from eukaryotes and bacteria. Many SLR proteins contain leader sequences and even SLR proteins without leader sequences, such as PodJ for example, are active in the periplasmic space. In addition, other bacterial SLR proteins, such as HcpA, ExoR, EnhC and LpnE seem to be responsible for the adaptation of bacteria to different eukaryotic hosts (ref 24; incorporated by reference in its entirety).

[0099] Of has about 44 Sell1 proteins, with many having molecular masses between 10-30 kDa. In addition, several other Sell1 proteins have molecular masses between 32-68 kDa. Most of the Sell1 proteins are predicted to have signal peptides and therefore are secreted proteins. Moreover, Sell1 proteins with molecular masses of 25 (#1361), 33 (#1414), and 68 (#1362) kDa blast in common with another Sell1 protein having a molecular mass of 59 (#1344) kDa. Five Sell1 proteins (#s 1360-1364) resides in an operon, with carbon starvation protein CstA located immediately upstream of this operon. Sell1 proteins #1343, 1344, and 1356 are also located near this operon. Other interesting candidates include diguanylate cyclase, a peptidase, putative outer membrane efflux protein OprC, outer membrane efflux protein OprM, LD-carboxypeptidase, stringent starvation protein B, ribonucleotide reductase, and Hsp20. Although most of the CM stimulatory activity is mediated by factors with masses between 10-30 kDa, the results of selective ultrafiltration using 30 kDa cutoff column indicate that the factors might exist as a multifunctional complex requiring a bacterial product of >30 kDa for optimal functioning. The fact that Sell1 proteins seem to serve as adaptor proteins for the assembly of membrane-bound macromolecular complexes might be of particular interest here. To evaluate whether one or more of these candidate proteins are the Of-derived bioactive factor(s) mediating the CM-induced stimulation of oxalate transport, the proteins (starting with

the target Sell1 proteins) were cloned and overexpressed in *E. coli* and the recombinant purified proteins were obtained (Table 2).

[0100] The effects of different concentrations (including different incubation periods) of the purified proteins on oxalate transport by C2 cells is similarly tested, as well as examining the effects of different combinations of the purified proteins. Testing the combined effects of the different proteins is important even if one candidate is found to have a significant stimulatory effect, giving the possibility that this might lead to a much higher stimulation (FIG. 3 and FIG. 5).

[0101] To evaluate the in vivo effects of the Sell1 proteins, LD-carboxypeptidase, and the peptidase (including any peptidomimetics versions of them or other embodiments), the proteins (individually or in different combinations) are administered (rectally, orally, both oral and rectal, or any other feasible route) to PH1 mice for up to 28 days. The PH1 mice are placed in metabolic cages and urine, plasma, and feces is collected on days 0, 7, 14, 21, and 28. When a factor is found to normalize or significantly reduce serum and urinary oxalate levels in PH1 mice, it is then examined whether the observed changes are due to factor-mediated enhanced net intestinal (small and/or large intestine depending on the route of administration) oxalate secretion. To this end, intestinal tissues (jejunum, ileum, cecum, proximal & distal colon) are isolated and mounted in Ussing chambers at the end of the treatment period found to be associated with the maximum reduction in serum and/or urine oxalate levels. Compared to vehicle-treated PH1 mice, if net basal cecal oxalate flux is converted from absorption to secretion, or significantly higher net basal secretory flux(s) is/re observed in the proximal and/or the distal colon of treated PH1 mice (due to significantly increased secretory flux and/or reduced absorptive flux), then such findings demonstrate that the factor reduces serum and/or urine oxalate levels by enhancing intestinal oxalate secretion. As a result of enhanced intestinal oxalate secretion/excretion, fecal oxalate is higher in the factor-treated mice, which is confirmed. Collectively, such findings provide a molecular basis for therapeutic application of a factor for prevention and/or treatment of hyperoxaluria, hyperoxalemia, and related COKS.

[0102] An alternative approach is to use differential proteomics. As described above, lowering culture medium oxalate from 37.5 mM (5 g/L) to 9.4 mM (1 g/L) led to a CM with >2-fold higher bioactivity, while increasing oxalate to 187.5 (25 g/L) led to a CM with reduced activity by ~50-70%. Further lowering of oxalate (0.5, 0.25, 0. g/L, as well as oxalate free culture medium) may lead to a CM with much higher bioactivity compared with 9.4 mM. If 0.25 g/L is found to be associated with the highest bioactivity, CMs are then prepared from Of cells grown in the low (will designate CM-low and its corresponding control medium will be OM-low) and high (will designate CM-high and its corresponding control medium will be OM-high) oxalate culture media (6 per condition). Label-free differential proteomics is used to analyze the samples. After excluding the proteins showing up in OM-low & high, the abundance of proteins in CM-low & high are compared. Observing proteins with significantly higher abundance in CM-low, while the same proteins are noted to be low or absent in CM-high, indicates that one or more of these proteins is an Of-derived bioactive factor(s). Promising candidate proteins are then be

cloned and overexpressed in *E. coli* and the effects of the recombinant protein(s) are tested as above.

[0103] An alternative strategy is to examine the Of transcriptome under low and high oxalate concentrations in the culture medium. To this end, Of cells grown in the low and high oxalate culture media are isolated and RNA for transcriptome analysis is prepared. RNA is extracted from replicate cultures (e.g., 3-5 per condition) grown under the two oxalate concentrations. Ribosomal RNA is subtracted and mRNA is used to generate cDNA libraries for sequencing. Transcript data is mapped to the assembled Of genome sequence using Bowtie (ref. 22; incorporated by reference in its entirety), and data normalization and differential expression analysis is done using the methods implemented in the DESeq2 R package (ref 23; incorporated by reference in its entirety). This approach facilitates the identification of genes (and/or operons) that are differentially regulated under these conditions. If, for example, certain genes are noted to be upregulated under low oxalate, but downregulated under the high oxalate, the protein products of these genes are then overexpressed in *E. coli* and the effects of the recombinant protein(s) are tested.

Example 3

[0104] The CM significantly reduced (>32.5%) urinary oxalate excretion in PH1 mice, indicating that the Of-derived factors retain their biological activity in vivo. To test whether the Of-derived factor(s) also decrease urinary oxalate excretion in enteric hyperoxaluria (EH) (e.g. IBD—and obesity-associated hyperoxaluria), the purified factor(s) are similarly given to SAPM1/YitFc (SAM) and ob/ob (ob) mice. SAM is a mouse model with remarkable similarities to human Crohn's disease developed as an ideal model for the IBD-associated hyperoxaluria (>2-fold hyperoxaluria compared to their controls). ob is an obesity model developed as an ideal model for the obesity associated hyperoxaluria (>3.3-fold hyperoxaluria compared to their controls). SAM and ob mice are treated with the purified factors as described above. SAM and ob mice are placed in metabolic cages and urine & feces are collected (including baseline collection). Observing normalization or significantly reduced urinary oxalate levels in factor-treated SAM and/or ob mice compared with vehicle-treated mice indicates that the factor has therapeutic potential not only for PH but also for EH and related COKS. Experiments then examine whether any observed reduction in urinary oxalate excretion in SAM and/or ob mice is due to factor-mediated enhanced net intestinal oxalate secretion.

[0105] Experiments are also conducted to evaluate the effects of the CM on the hyperoxaluria observed in the SAM and ob mice. Given the unavailability of any specific therapy that effectively reduces urinary oxalate excretion, the CM provides a therapeutic option under conditions in which it significantly reduces urine oxalate levels in these mouse models. Since Of has been given (as a frozen cell paste or a lyophilized Of formulated in enteric coated capsules) to PH patients (ref. 18; incorporated by reference in its entirety), it is reasonable to give a product of this bacterium to PH patients. Of CM or OM will be similarly (as described above with PH1 mice) given to SAM and ob mice and the results will be similarly interpreted.

[0106] Artificial colonization of wild-type mice with the human Of strain HC-1 led to the presence of the bacteria not only in the colon, but unexpectedly also in the small

intestine for some time (ref 15; incorporated by reference in its entirety). This was associated with a significant net oxalate secretion in the distal ileum, cecum, and distal colon, as well as significantly reduced urinary oxalate excretion. In view of these findings, it is possible that oral administration of Of CM (or the purified factor(s)) might promote significant stimulation of net oxalate secretion in both small and large intestines, thereby potentially leading to normalization or greater reduction in plasma and urinary oxalate levels in PH1 mice compared to rectal CM. Therefore, gelatin capsules containing freeze-dried Of CM (or the purified factor(s)) are prepared for oral administration. To protect the contents of the capsules during passage through the acidic upper gastrointestinal tract, the capsules are coated with Eudragit L 100-55 (ref. 14), which will also protect their contents from protease digestion. Placebo-treated mice will receive Eudragit-coated empty capsules. This approach was used for making encapsulated freeze-dried preparation of Of lysate which was then given orally to hyperoxaluric rats and led to induction of distal colonic oxalate secretion. A 2x2 factorial design is followed with the following 4 groups: placebo-placebo, oral CM, rectal CM, and oral+rectal CM. The PH1, SAM, and ob mice receive the capsules by an intragastric needle twice daily for 5-20 days, and urine, plasma, & feces is collected (including baseline collection), as well as intestinal tissues (jejunum, ileum, cecum, proximal & distal colon) are isolated and mounted in Ussing chambers at the end of the treatment period. The data are analyzed by repeated measures ANOVA with treatment group as the between-subjects factor and time as the within-subjects repeated factor. The treatment group is tested by time interaction to determine whether the change with treatment varies based on treatment received. Observing normalization or a greater reduction in plasma and urinary oxalate levels in PH1 mice with oral CM compared to rectal CM, due to significant stimulation of net oxalate secretion in the small and/or large intestine(s), indicates that the oral route is better. Finding that oral+rectal CM is associated with a much higher reduction in plasma and urinary oxalate levels compared with either oral CM or rectal CM alone, has significant therapeutic implications. Of note is that the CM or OM was given as rectal enemas through special tips that were introduced to ~2 cm from the anus, and the mice were then held from their tails with the heads down and kept in that position for ~one minute.

Example 4

[0107] The nucleotide sequences of the proteins in Table 2 from the OxB *oxalobacter* strain were sequenced compared to the human *oxalobacter* strain OXCC13. The OxB, OXC, or other Of-derived nucleotide sequences, and variants thereof, may find use in embodiments described herein. In some embodiments, nucleic acids and/or polypeptides comprising such sequences find use in the embodiments described herein.

Example 5

[0108] Experiments were conducted during development of embodiments herein to determine the molecular weight(s) (MWs) of the secreted factor(s), the conditioned medium (CM) was subjected to selective ultrafiltration using 10- and 30-kDa cutoff spin columns. C2 cells were untreated (UT) or treated with the CM, filtrate (F), retentate (R), or F+R. Using

a 10-kDa column, the R and the F+R significantly stimulated oxalate uptake by C2 cells, while the F has no effect (FIG. 6A). Using a 30-kDa column, the F and the F+R significantly stimulated oxalate uptake by C2 cells, while the R has no effect (FIG. 6B). Collectively, these results indicate that the MWs of the factors are largely between 10-30 kDa. Since F+R have a better stimulatory effect than F, while R has no effect, such data suggest the possibility that these factors might exist as a multifunctional complex requiring a bacterial product of >30 kDa for optimal functioning. Therefore, evaluating the effects of different combinations of the Sell1 proteins and the other proteins (305, 308, 309, & 314) is critical to reach to a level of stimulation similar to the CM-induced stimulation (~3-fold) (FIG. 5 and FIG. 6). Sell1 proteins 304, 317, 319, 322, 323, 324, & 325 stimulate oxalate transport by ~1.2-1.6-fold in preliminary studies.

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gtcgacaagg cggatgtgaa tgattcgacg gtgtttaagg cgagacagga gatgaactga 1920

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<210> SEQ ID NO 6

<211> LENGTH: 639

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 6

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Met Lys Ser Val Ser His Met Gly Phe Tyr Arg Phe Lys Ser Phe Val
1           5           10           15
Phe Ser Val Val Val Phe Ala Ala Gly Cys Ala Trp Val Pro Ser Val
20           25           30
Ser Phe Ala Gly Gln Lys Asp Asp Cys Ala Ser Val Gly Cys Ser Asp
35           40           45
Thr Thr Val Gln Lys Asp Gly Asp Leu Glu Lys Ile Arg Gln Lys Ala
50           55           60
Gln Gln Gly Asp Ala Asn Ala Gln Met Ala Leu Gly Leu Arg Tyr Leu
65           70           75           80
Met Gly Asn Gly Leu Ala Ala Asp Glu Val Leu Ala Gln Glu Trp Phe
85           90           95
Leu Lys Ser Ala Gln Gln Asn Asn Val Val Ala Gln Val Ala Leu Ala
100          105          110
Thr Met Leu Ala Phe Glu Ser Asp Arg Gln Asp Leu Pro Ala Ala Ala
115          120          125

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-continued

Met Trp Phe Ser Lys Ala Ala Asp Ala Gly Asn Val Gln Ala Met Ser
 130 135 140

Glu Leu Val Arg Leu Tyr Glu Thr Gly Ser Gly Val Thr Arg Asp Met
 145 150 155 160

Ala Lys Ala Glu Glu Trp Arg Val Arg Ala Lys Met Arg Ser Asp Ala
 165 170 175

Val Lys Leu Glu Arg Val Trp Lys Ile Ala Leu Ala Asp Lys Ala Arg
 180 185 190

Trp Met Arg Lys Thr Ser Arg Pro Val Glu Leu Ala Ala Gln Asp Gly
 195 200 205

Val Ala Thr Thr Ala Ala Arg Ile Asp Val Val Ala Leu Lys Lys Ala
 210 215 220

Ala Glu Asn Gly Asp Asp His Ala Gln Thr Leu Leu Gly Ala Leu Leu
 225 230 235 240

Ala Thr Gly Asp Gly Val Lys Lys Asp Glu Lys Ala Ala Ile Gly Trp
 245 250 255

Phe Glu Lys Ala Ala Asp Ser Gly Asn Thr Gln Ala Gln Ala Val Leu
 260 265 270

Gly Glu Leu Tyr Gly Leu Gly Trp Gly Gly Leu Lys Lys Asp Glu Ala
 275 280 285

Lys Ala Ala Lys Trp Met Glu Lys Ala Ala Leu Gly Gly Leu Val Ala
 290 295 300

Ala Lys Ala Ala Trp Gly Ser Met Leu Ser Gln Gly Lys Gly Val Glu
 305 310 315 320

Lys Asp Pro Lys Lys Gly Leu Glu Trp Phe Val Gln Ala Gly Gln Asp
 325 330 335

Gly Asp Gly Arg Thr Arg Leu Leu Met Gly Met Met Leu Ile His Gln
 340 345 350

Asn Arg Asp Ala Ala Ala Gln Trp Phe Tyr Gly Ala Ala Glu Val Gly
 355 360 365

Asp Glu Glu Val Leu Ser Ala Leu Gly Thr Phe Tyr Gly Trp Gly Asn
 370 375 380

Gly Pro Val Leu Asp Glu Ser Glu Lys Leu Ser Glu Val Arg Arg Tyr
 385 390 395 400

Ala Gln Arg Asp Glu Ser Glu Ala Gln Gln Met Met Gly Phe Leu Tyr
 405 410 415

Gly Glu Gly Trp Gly Ala Lys Arg Asp Pro Val Lys Ala Glu Tyr Trp
 420 425 430

Phe Asp Lys Ala Ala Ala Ser Gly Asp Val Glu Val Trp Leu Pro Leu
 435 440 445

Gly Leu Leu Tyr Ala Glu Thr Gly Arg Asp Asp Met Ala Ala Ala Ala
 450 455 460

Phe Ala Lys Ala Val Ala Ser Gly Gly Phe Gly Leu Ala Asn Asp Gly
 465 470 475 480

Glu Leu Leu Gln Leu Ile Phe Val Asp Ser Glu Lys Met Pro Asp Val
 485 490 495

Gly Asn Ala Leu Lys Arg Pro Ala Ser Ala Lys Lys Thr Ala Ser Gly
 500 505 510

Ala Lys Glu Pro Asn Gly Gly Lys Lys Ala Gly Val Asn Asp Lys Asp
 515 520 525

-continued

Gly Val Ala Ser Asp Glu Arg Leu Val Arg Val Ala Lys Lys Arg Ala
 530 535 540

Phe Val Leu Ala Glu Ala Lys Lys Gly Asn Pro Ala Ala Gln Leu Met
 545 550 555 560

Met Ala Arg Ile Leu Lys Glu Gly Trp Gly Val Lys Lys Asp Glu Glu
 565 570 575

Ala Ala Ala Ser Leu Arg Ala Asp Gly Ile Arg Gly Met Cys Ala Ala
 580 585 590

Leu Lys Asp Lys Ala Glu Glu Glu Pro Leu Cys Ser Asn Glu Gly Asp
 595 600 605

Gly Gly Asn Gly Gly Val Leu Pro Asp Ala Ala Ser Val Asp Lys Ala
 610 615 620

Asp Val Asn Asp Ser Thr Val Phe Lys Ala Arg Gln Glu Met Asn
 625 630 635

<210> SEQ ID NO 7
 <211> LENGTH: 456
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 7

```

atggctttgg cggatggggc ggctgctccg gtggcagtga cgtcttatgc gcagcagccg      60
ttgaagctgg tgcaggaaaa ggcttctgac ggggatgggt ctgccgaact tgaactgggt      120
ttgcggtatg ttttcggctc tgacggggtc aaaaatgtgc cgctcggggg ttcttgatc      180
aataaggcgg ctctaaaggg tattccgcag gcggagcatg agatggggtc gctgtatctg      240
atgggggattg gcgttgccca aagcaatgtg atggctgtgg cctggtacag gaaggcggca      300
attcagggtt acgccccgtc gcaaacggcg atggggatg cgtatgaaga aggggcccgg      360
gtgccacagg atcgggatct ggcccgttac tggtttgaca aggcggcggc acagggaaat      420
ggtattgccc tggaaagcct tgaaggaggg atgtag                                     456
    
```

<210> SEQ ID NO 8
 <211> LENGTH: 151
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 8

Met Ala Leu Ala Asp Gly Ala Ala Ala Pro Val Ala Val Thr Ser Tyr
 1 5 10 15

Ala Gln Gln Pro Leu Lys Leu Val Gln Glu Lys Ala Ser Asp Gly Asp
 20 25 30

Gly Ser Ala Glu Leu Glu Leu Gly Leu Arg Tyr Val Phe Gly Ser Asp
 35 40 45

Gly Val Lys Asn Val Pro Leu Gly Val Ser Trp Ile Asn Lys Ala Ala
 50 55 60

Leu Lys Gly Ile Pro Gln Ala Glu His Glu Met Gly Ser Leu Tyr Leu
 65 70 75 80

Met Gly Ile Gly Val Ala Gln Ser Asn Val Met Ala Val Ala Trp Tyr
 85 90 95

Arg Lys Ala Ala Ile Gln Gly Tyr Ala Pro Ser Gln Thr Ala Met Gly
 100 105 110

Tyr Ala Tyr Glu Glu Gly Ala Gly Val Pro Gln Asp Ala Asp Leu Ala
 115 120 125

-continued

Arg Tyr Trp Phe Asp Lys Ala Ala Ala Gln Gly Asn Gly Ile Ala Val
130 135 140

Glu Ser Leu Glu Gly Gly Met
145 150

<210> SEQ ID NO 9
<211> LENGTH: 1470
<212> TYPE: DNA
<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 9

```

atgaaaagac gggttttgta tgggtgtgctg tttgcgggtgc tggcgggggtg ttctgccggt      60
gcgttggcgt cctgtgtccgg cttgtcggat gggctggcgc agcttgaacg gaggcagttc      120
gaatcggtt atgcgacgt catgccggag gcggagaagg ggaatccccg tgcggcgctg      180
gaggtgggga agctgctgtt gacggggcgc ggggtggcaa aggatgaagc ggcggcggtg      240
aagtggcttc tgggtggcgc ggtatgtggc aaccgggatg cgcagtatat gctgggggcg      300
atgtcgggtg aggggatcgg tctgccgaag gattctcagg tggcgttgac ctggctgtcg      360
aaggcggcgg cgcaggggga tgcgctgctg aagacggctt tggggattct gatgcagtcc      420
gccgggccgg gatcgcagca tacggaacag gcagcccgtt ggttcgagag ggcggcgcg      480
tcaggggaac cggaggcga acggcgtcgg gcgctgatgc tggcgtctgg ccgggggggtg      540
gccaaaaatg agggtagagg actgaaatgg ttaagaagg cggctgtcgc ggggatgtg      600
gaagcgcagc gcaatctggg gattatgttc tcgacgggaa agggggtgac ggcggcgaag      660
ccggattttg cggaaagcgc acgctggtac ggcctggcag cgaagaaggg ggatgcgaag      720
gcgcagtatg ggttgggcat tttgatgctg aaggggcagg ggggtggcgc cgatcaggaa      780
aaggcgtgta ttctgtatcg catggcggcg actcaggggc tggcgaaggc ggagtatgcc      840
gtcgggctgg cgtatgcgta tggacggggg acggcacaaa atgatgtgaa ggcggcccag      900
tggttcgagg cggcggcgca gcaggggggtg gtgcgtgcgc aatataatct cgctctgatg      960
ctggaggcgg gtcgcggtcg gcctgtggat acggtggcgg cgagcaagtg gtttttgatg     1020
gcggcggaga agggcttgcg ggaggcacia tacaatatgg ggtatcacta tgccgagggg     1080
aaaggggtgc cacgcgatca gggcaaggcg gtgttctggt atgaaaaggc ggcggctgcc     1140
gggatgtgta agcccagta caatctgggg atgctgtatc tgaacggggg taatggcaag     1200
gcgatgacg aaaaggcggc tttttctac cggatggcgg ccggggcggg atatggcccg     1260
gcgatgtacc ggctggcggg gttgtatgag gaagcccggt gggtaaagca gagttatcag     1320
ctggcagggg aatggatga gcgggaggat ctggcccgca aagtgaagat tgacgaggcg     1380
atgaaaaaga atccgcaccc gtttgtgcaa cggactctgc aggtgccgga tgatttgaat     1440
caatcttcag ataaattggc gggacattag                                     1470

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<210> SEQ ID NO 10
<211> LENGTH: 489
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 10

Met Lys Arg Arg Val Leu Tyr Gly Val Leu Phe Ala Val Leu Ala Gly
1 5 10 15

-continued

Cys Ser Ala Gly Ala Leu Ala Ser Val Ser Gly Leu Ser Asp Gly Leu
 20 25 30

Ala Gln Leu Glu Arg Arg Gln Phe Glu Ser Ala Tyr Ala Thr Leu Met
 35 40 45

Pro Glu Ala Glu Lys Gly Asn Pro Arg Ala Ala Leu Glu Val Gly Lys
 50 55 60

Leu Leu Leu Thr Gly Arg Gly Val Ala Lys Asp Glu Ala Ala Ala Val
 65 85 70 90 75 110 80

Lys Trp Leu Leu Val Ala Ala Asp Ser Gly Asn Arg Asp Ala Gln Tyr
 85 90 95

Met Leu Gly Ala Met Ser Val Glu Gly Ile Gly Leu Pro Lys Asp Ser
 100 105 110

Gln Val Ala Leu Thr Trp Leu Ser Lys Ala Ala Ala Gln Gly Asp Ala
 115 120 125

Arg Ala Lys Thr Ala Leu Gly Ile Leu Met Gln Ser Ala Gly Pro Gly
 130 135 140

Ser Gln His Thr Glu Gln Ala Ala Arg Trp Phe Glu Arg Ala Ala Ala
 145 150 155 160

Ser Gly Glu Pro Glu Ala Gln Arg Arg Trp Ala Leu Met Leu Ala Ser
 165 170 175

Gly Arg Gly Val Ala Lys Asn Glu Gly Glu Ala Leu Lys Trp Phe Lys
 180 185 190

Lys Ala Ala Val Ala Gly Asp Val Glu Ala Gln Arg Asn Leu Gly Ile
 195 200 205

Met Leu Ser Thr Gly Lys Gly Val Thr Gly Gly Lys Pro Asp Phe Ala
 210 215 220

Glu Ala Ala Arg Trp Tyr Gly Leu Ala Ala Lys Lys Gly Asp Ala Lys
 225 230 235 240

Ala Gln Tyr Gly Leu Gly Ile Leu Tyr Ala Lys Gly Gln Gly Val Ala
 245 250 255

Pro Asp Gln Glu Lys Ala Leu Ile Leu Tyr Arg Met Ala Ala Thr Gln
 260 265 270

Gly Leu Ala Thr Ala Glu Tyr Ala Val Gly Leu Ala Tyr Ala Tyr Gly
 275 280 285

Arg Gly Thr Ala Gln Asn Asp Val Lys Ala Ala Asp Trp Phe Glu Ala
 290 295 300

Ala Ala Gln Gln Gly Val Val Arg Ala Gln Tyr Asn Leu Ala Leu Met
 305 310 315 320

Leu Glu Ala Gly Arg Gly Arg Pro Val Asp Thr Val Ala Ala Ser Lys
 325 330 335

Trp Phe Leu Met Ala Ala Glu Lys Gly Leu Arg Glu Ala Gln Tyr Asn
 340 345 350

Met Gly Tyr His Tyr Ala Glu Gly Lys Gly Val Pro Arg Asp Gln Gly
 355 360 365

Lys Ala Val Phe Trp Tyr Glu Lys Ala Ala Ala Ala Gly Asp Val Lys
 370 375 380

Ala Gln Tyr Asn Leu Gly Met Leu Tyr Leu Asn Gly Val Asn Gly Lys
 385 390 395 400

Ala Asp Asp Glu Lys Ala Ala Phe Phe Tyr Arg Met Ala Ala Gly Ala
 405 410 415

Gly Tyr Gly Pro Ala Met Tyr Arg Leu Ala Val Leu Tyr Glu Glu Gly

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420	425	430
Arg Gly Val Lys Gln Ser Tyr Gln Leu Ala Gly Glu Trp Tyr Glu Arg		
435	440	445
Ala Asp Leu Ala Ala Lys Val Lys Ile Asp Glu Ala Met Lys Lys Asn		
450	455	460
Pro His Pro Phe Val Gln Arg Thr Leu Gln Val Pro Asp Asp Leu Asn		
465	470	475
480		
Gln Ser Ser Asp Lys Leu Ala Gly His		
485		

<210> SEQ ID NO 11
 <211> LENGTH: 1206
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 11

```

atgaaaaaag gggttgctt cgcgctggg gctcttggga gcatactggg ttgtacggcc    60
ttggccggaa gcaaaaagaa actggacgcc ttgctggcga tgccggtgaa agagacaaaag    120
gtttttgtcg aatcgaatga agagccgttg tttgtgatgt tgaaaagctc gttgtccgaa    180
gagacgggatg aaactgtcgg gaagccggtt gtttcggaaa agacggcaga gccgggacaa    240
gacaatgacg gtgaaacagg gggagcctgg atgcagcagc tgcggcatca ggcggatcag    300
ggggatgcga aatccgcttt ctggctgggc cggttcacgg ttgaggacag ccgggacggg    360
aaaacgattg acgagggcat cgtctgatc cggcgttctg ccgaaggggg attcgtgctg    420
gcacaattgt atctcggcac gctgtatgcc aatgggactc atgtgaaggc tgaccgcac    480
gaggcggaaa aatggctttc cagggccgca gggcaaggtt ctccgatggt tcagctttat    540
ctcggcctga tgtatggtca tggcaagggt gttccccgtg acttgaacaa gtcgcttttc    600
tgggtgaaaa aggcggcgga caggggcttg ccgcatgcgc agctggcgcg tgggcttttt    660
gcgtcgtttt cccattatta tccccgggat gatgaaaagg ccgtgctgta tctgacgaaa    720
gcccgaaaag aggggatgcc gatggcccag ttttatctgg cgctgatgta tcagcgtggc    780
cggggtgtcg aacagagtaa cgagcaggcc ttgcaactgga atatgctggc ggcggaacag    840
ggctatccgg atgccagta tgcgatgtcg cggatggcgg aactcggtat cggggtgacg    900
gccgataaag catggagcat gatgtggctg gatcgtgccg cccatcacgg gatgccgctg    960
gcgcaatatt tgatgggcat ggcctatctg gaagaaaat cggccccgca ggacttgcc    1020
gttgcggcgg catggtttta caaggcggcg atgcaggaa atgccgatgc ccagttgca    1080
ctcggttata tgtatgccag ggaatcggg gttcctgtgg acaagccgaa ggcgggttgc    1140
tggcttgaag aggccgcttc ggctggcaat acggtggccg ggcagtggct gaaacaactg    1200
gattga                                           1206
    
```

<210> SEQ ID NO 12
 <211> LENGTH: 401
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 12

Met Lys Lys Gly Val Ala Phe Ala Leu Val Ala Leu Gly Ser Ile Leu
1 5 10 15
Val Cys Thr Ala Leu Ala Gly Ser Glu Lys Lys Leu Asp Ala Leu Leu

-continued

20					25					30					
Ala	Met	Pro	Val	Lys	Glu	Thr	Lys	Val	Phe	Val	Glu	Ser	Asn	Glu	Glu
	35						40					45			
Pro	Leu	Phe	Val	Met	Leu	Lys	Ser	Ser	Leu	Ser	Glu	Glu	Thr	Asp	Glu
	50					55					60				
Thr	Val	Gly	Lys	Pro	Val	Val	Ser	Glu	Lys	Thr	Ala	Glu	Pro	Gly	Gln
	65					70					75				80
Asp	Asn	Asp	Gly	Glu	Thr	Gly	Gly	Ala	Trp	Met	Gln	Gln	Leu	Arg	His
			85						90					95	
Gln	Ala	Asp	Gln	Gly	Asp	Ala	Lys	Ser	Ala	Phe	Trp	Leu	Gly	Arg	Phe
			100						105					110	
Thr	Val	Glu	Asp	Ser	Arg	Asp	Gly	Lys	Thr	Ile	Asp	Glu	Gly	Ile	Arg
		115					120					125			
Leu	Ile	Arg	Arg	Ser	Ala	Glu	Gly	Gly	Phe	Val	Arg	Ala	Gln	Leu	Tyr
		130				135					140				
Leu	Gly	Thr	Leu	Tyr	Ala	Asn	Gly	Thr	His	Val	Lys	Ala	Asp	Pro	His
		145				150					155				160
Glu	Ala	Glu	Lys	Trp	Leu	Ser	Arg	Ala	Ala	Gly	Gln	Gly	Ser	Pro	Met
			165						170					175	
Val	Gln	Leu	Tyr	Leu	Gly	Leu	Met	Tyr	Gly	His	Gly	Lys	Gly	Val	Pro
			180					185						190	
Arg	Asp	Leu	Asn	Lys	Ser	Leu	Phe	Trp	Val	Glu	Lys	Ala	Ala	Asp	Arg
		195					200					205			
Gly	Leu	Pro	His	Ala	Gln	Leu	Ala	Arg	Gly	Leu	Phe	Ala	Ser	Phe	Ser
		210				215					220				
His	Tyr	Tyr	Pro	Arg	Asp	Asp	Glu	Lys	Ala	Val	Leu	Tyr	Leu	Thr	Lys
			225			230					235				240
Ala	Ala	Lys	Gln	Gly	Met	Pro	Met	Ala	Gln	Phe	Tyr	Leu	Ala	Leu	Met
			245						250					255	
Tyr	Gln	Arg	Gly	Arg	Gly	Val	Glu	Gln	Ser	Asn	Glu	Gln	Ala	Leu	His
			260					265						270	
Trp	Asn	Met	Leu	Ala	Ala	Glu	Gln	Gly	Tyr	Pro	Asp	Ala	Glu	Tyr	Ala
		275					280					285			
Met	Ser	Arg	Met	Ala	Glu	Leu	Gly	Ile	Gly	Val	Thr	Ala	Asp	Lys	Ala
		290				295					300				
Trp	Ser	Met	Met	Trp	Leu	Asp	Arg	Ala	Ala	His	His	Gly	Met	Pro	Leu
		305				310					315				320
Ala	Gln	Tyr	Leu	Met	Gly	Met	Ala	Tyr	Leu	Glu	Gly	Lys	Ser	Val	Pro
			325						330					335	
Gln	Asp	Leu	Pro	Val	Ala	Ala	Ala	Trp	Phe	Tyr	Lys	Ala	Ala	Met	Gln
			340					345						350	
Gly	Asn	Ala	Asp	Ala	Gln	Leu	Arg	Leu	Gly	Tyr	Met	Tyr	Ala	Arg	Gly
		355					360					365			
Ile	Gly	Val	Pro	Val	Asp	Lys	Pro	Lys	Ala	Val	Ala	Trp	Leu	Glu	Lys
		370				375					380				
Ala	Ala	Ser	Ala	Gly	Asn	Thr	Val	Ala	Gly	Gln	Trp	Leu	Lys	Gln	Leu
			385			390					395				400

Asp

<210> SEQ ID NO 13

<211> LENGTH: 900

-continued

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 13

```

atgaagaaaa ccatctatca attcgcactt atttttatta ccttaagttt tttacggcg      60
tgtacaggca atattgatc  ggaagcgaa aataaaggta ttcaacatta taaaaatggg      120
gaatatcaaa aagcaattcc cttattggag aaagctgaag atggcggcag ttccagtgcc      180
gcattctatc ttggagaaat atttcgtaaa ggtgaaggcg ttaatcaaga ttttgggaga      240
tcatgtacgc attatataaa atcagcaaaa ggtggaata acaatgccta tttactggcc      300
ggttcgtggt tcgtaatggg aaaaggtgtg aagcaggatt ttgcagaagc attgaaatgg      360
tttaaaaaag cgctccgatg aagtgaaaaa actgatttga ctgaatctga caagaaatat      420
ttaacgcgat cattggccac aatgtactat tccgaaaaag gaacctgca agacttcagc      480
gaagccgcaa agtgggcgga aaaagcggct gaactgggtg atgcgaattc acaggcggtt      540
atggcggtcc tgctttatac ggggcagggc gttctggctg acaggaaggc tgcccggata      600
tgggcgcaaa aatcagcgga tcagggaaac gatctggggg aagtctgat ggggtgattc      660
aatcaatatg ctgattcccc ggacatgaaa gcggcctttg actggtatga gaaatcggca      720
aaacagggca acccggccgc gcagtatcaa ttgggaacgt tttatgaaga aggcattatt      780
gttctgaag acattgaaaa agcccagct tgttataaac aggcggccga tagcaaaaag      840
tcggatactc tggcaaggc tctgatggat ttcgaggcgc gacagaaaaa acagaaatga      900

```

<210> SEQ ID NO 14

<211> LENGTH: 299

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 14

```

Met Lys Lys Thr Ile Tyr Gln Phe Ala Leu Ile Phe Ile Thr Leu Ser
 1             5             10             15
Phe Leu Thr Ala Cys Thr Gly Asn Ile Asp Ser Glu Ser Glu Asn Lys
          20             25             30
Gly Ile Gln His Tyr Lys Asn Gly Glu Tyr Gln Lys Ala Ile Pro Leu
          35             40             45
Leu Glu Lys Ala Glu Asp Gly Gly Ser Ser Ser Ala Ala Phe Tyr Leu
          50             55             60
Gly Glu Ile Phe Arg Lys Gly Glu Gly Val Asn Gln Asp Phe Gly Arg
          65             70             75             80
Ser Cys Thr His Tyr Ile Lys Ser Ala Lys Gly Gly Asn Asn Asn Ala
          85             90             95
Tyr Leu Leu Ala Gly Ser Cys Phe Val Met Gly Lys Gly Val Lys Gln
          100            105            110
Asp Phe Ala Glu Ala Leu Lys Trp Phe Lys Lys Ala Ser Asp Glu Ser
          115            120            125
Glu Lys Thr Asp Leu Thr Glu Ser Asp Lys Lys Tyr Leu Thr Arg Ser
          130            135            140
Leu Ala Thr Met Tyr Tyr Ser Gly Lys Gly Thr Leu Gln Asp Phe Ser
          145            150            155            160
Glu Ala Ala Lys Trp Ala Glu Lys Ala Ala Glu Leu Gly Asp Ala Asn
          165            170            175

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-continued

Ser Gln Ala Val Met Ala Phe Leu Leu Tyr Thr Gly Gln Gly Val Leu
 180 185 190

Ala Asp Arg Lys Ala Ala Arg Ile Trp Ala Gln Lys Ser Ala Asp Gln
 195 200 205

Gly Asn Asp Leu Gly Glu Val Leu Met Gly Val Phe Asn Gln Tyr Ala
 210 215 220

Asp Ser Pro Asp Met Lys Ala Ala Phe Asp Trp Tyr Glu Lys Ser Ala
 225 230 235 240

Lys Gln Gly Asn Pro Ala Ala Gln Tyr Gln Leu Gly Thr Phe Tyr Glu
 245 250 255

Glu Gly Ile Ile Val Pro Glu Asp Ile Glu Lys Ala His Ala Cys Tyr
 260 265 270

Lys Gln Ala Ala Asp Ser Lys Lys Ser Asp Thr Leu Val Lys Ala Leu
 275 280 285

Met Asp Phe Glu Ala Arg Gln Lys Lys Gln Lys
 290 295

<210> SEQ ID NO 15
 <211> LENGTH: 918
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 15

```

atgtcaaac agggatgacag gagaaggaca gggatgaaga tgtcgggttt cagatttgcg      60
ttgggggtgt cggtatgtct gttgacggta tgggttccag cgacgtctat ggcaggaaac      120
gtgtcggcgt cggcatatca gatgtctcaa ggcgaagacg aagcatggac gttttatcgt      180
accggacagt acgggaaggc actggccgct ttcagggggc tggagacgaa aggggatggt      240
gccgccttgt atggactggg tgtgatggcg acgaatggtc tccggtatgcc ccgtaatgat      300
gaaaaggcgc tggataggtt tagggaaggg gcggcaaaag gctcgcgtga ggcgcaattc      360
gggctggggg cgatgtacga tttgagtcgc ggtgtccggc aggatatgac gctggcgatc      420
gactggtatg aaaagtggc gagagcggga tatgcccgg ctctgacgcg gctgggcagg      480
atgaatttgc tggggagggg gatgtcccgc aattatggga aggcgttccg gtttttcaaa      540
cagtcggcac agcgtgggga tagggacggt gagttttatt tccgggatgat gtttatccgt      600
ggctggggga cgaagcggga tgtggaagag gcggctggct ggatcagaaa ggcggcggag      660
aagggacagc cggaaagcgt gcgggtgatg tcgaccttgt atgaggggtg ttatggcggt      720
gccc aaagtg agaagatgc gctcgtctgg ctggaaaaga gcgttgatgc gggcgacagg      780
gaagcgatga ggcggctggc agcgggtgat gaaaacggca cgttcggtgt gacgcccggac      840
aaggagaagg cccgtttgct gaaaaagaag gcagccgagg tgaaaaaggc tgtttttccg      900
gattatgccg tgatgtag      918
    
```

<210> SEQ ID NO 16
 <211> LENGTH: 305
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 16

Met Ser Lys Gln Gly Asp Arg Arg Arg Thr Gly Met Lys Met Ser Gly
 1 5 10 15

Phe Arg Phe Ala Leu Gly Val Ser Val Cys Leu Leu Thr Val Trp Val

-continued

20		25		30											
Pro	Ala	Thr	Ser	Met	Ala	Gly	Asn	Val	Ser	Ala	Ser	Ala	Tyr	Gln	Met
	35						40					45			
Ser	Gln	Gly	Glu	Asp	Glu	Ala	Trp	Thr	Phe	Tyr	Arg	Thr	Gly	Gln	Tyr
	50					55					60				
Gly	Lys	Ala	Leu	Ala	Ala	Phe	Arg	Gly	Leu	Glu	Thr	Lys	Gly	Asp	Val
	65				70					75					80
Ala	Ala	Leu	Tyr	Gly	Leu	Gly	Val	Met	Ala	Thr	Asn	Gly	Leu	Gly	Met
				85					90					95	
Pro	Arg	Asn	Asp	Glu	Lys	Ala	Leu	Val	Trp	Phe	Arg	Glu	Gly	Ala	Ala
			100					105						110	
Lys	Gly	Ser	Arg	Glu	Ala	Gln	Phe	Gly	Leu	Gly	Ala	Met	Tyr	Asp	Leu
		115					120						125		
Ser	Arg	Gly	Val	Arg	Gln	Asp	Met	Thr	Leu	Ala	Ile	Asp	Trp	Tyr	Glu
		130				135							140		
Lys	Ser	Ala	Arg	Ala	Gly	Tyr	Ala	Pro	Ala	Leu	Thr	Arg	Leu	Gly	Arg
					150					155					160
Met	Asn	Leu	Leu	Gly	Arg	Gly	Met	Ser	Arg	Asn	Tyr	Gly	Lys	Ala	Phe
				165					170						175
Arg	Phe	Phe	Lys	Gln	Ser	Ala	Gln	Arg	Gly	Asp	Arg	Asp	Gly	Glu	Phe
			180					185						190	
Tyr	Phe	Gly	Met	Met	Phe	Ile	Arg	Gly	Trp	Gly	Thr	Lys	Arg	Asp	Val
		195					200						205		
Glu	Glu	Ala	Ala	Gly	Trp	Ile	Arg	Lys	Ala	Ala	Glu	Lys	Gly	Gln	Pro
		210				215					220				
Glu	Ala	Met	Arg	Val	Met	Ser	Thr	Leu	Tyr	Glu	Gly	Gly	Tyr	Gly	Val
		225			230					235					240
Ala	Gln	Ser	Glu	Lys	Asp	Ala	Leu	Val	Trp	Leu	Glu	Lys	Ser	Val	Asp
			245						250						255
Ala	Gly	Asp	Arg	Glu	Ala	Met	Arg	Arg	Leu	Ala	Ala	Val	Tyr	Glu	Asn
			260					265						270	
Gly	Thr	Phe	Gly	Val	Thr	Pro	Asp	Lys	Glu	Lys	Ala	Arg	Leu	Leu	Lys
		275					280						285		
Lys	Lys	Ala	Ala	Glu	Val	Lys	Lys	Ala	Val	Phe	Pro	Asp	Tyr	Ala	Val
		290				295					300				
Met															
															305

<210> SEQ ID NO 17
 <211> LENGTH: 819
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 17

atgatgaaaa ccatcctcaa agccctcttc ttgcattca ttctgattgg ttccactctg	60
gcgttcgcg acaatgttga agaaggcaac catctctata atgctggaaa ataccaggaa	120
gccctgacct tttcatgaa accggatgcg gtcaataacc cagccaccat gaaccggatc	180
gggtatatgt acgacgaagg tcaggagtc aaaaagatc caaagaagc ctccaagtgg	240
tacaaaaaag cagctgatgc caattacca gttgccagc ttaatctggg gcttatgtat	300
caacatggca ccggcgtctc aaaagatc aatgaatcca ttaaatgggt tcgtaaagca	360

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gcagaacaaa atgatccoga cgctgaaatg aaaatgggct atttgaccgc aacaggaaca 420
ggggtcacaaa aagattatca agaagctata caatggatc aacgcgctgc tgaacatggc 480
gatagtgcag cttatgcaca aattggactt ttctatactc tgggcaatgg tgtcaaaaaa 540
gacgtcaacc gtgctgtcca gtattacatt atgggcgctc aaaaggggtga tgccagagca 600
caggcctttt tgggaaaagc atatgccttg ggcagaggta tccaaccgga tagtgaaaaa 660
gccctctact ggtacaaaac agccgccaga aacggcaacg tcaacgccat gaaagaactg 720
ggttccatct atgcaaaaagg ccgtctcggg gtcaagccag accagcagga agcacaacga 780
tggaacgaca tggccagaaa agctgaacag aaaaattaa 819

```

<210> SEQ ID NO 18

<211> LENGTH: 272

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 18

```

Met Met Lys Thr Ile Leu Lys Ala Leu Phe Phe Ala Phe Ile Leu Ile
 1          5          10          15
Gly Ser Thr Leu Ala Phe Ala Asp Asn Val Glu Glu Gly Asn His Leu
 20          25          30
Tyr Asn Ala Gly Lys Tyr Gln Glu Ala Leu Thr Phe Phe Met Lys Pro
 35          40          45
Asp Ala Val Asn Asn Pro Ala Thr Met Asn Arg Ile Gly Tyr Met Tyr
 50          55          60
Asp Glu Gly Gln Gly Val Lys Lys Asp Pro Lys Glu Ala Phe Lys Trp
 65          70          75          80
Tyr Lys Lys Ala Ala Asp Ala Asn Leu Pro Val Ala Gln Phe Asn Leu
 85          90          95
Gly Leu Met Tyr Gln His Gly Thr Gly Val Ser Lys Asp Ile Asn Glu
 100         105         110
Ser Ile Lys Trp Phe Arg Lys Ala Ala Glu Gln Asn Asp Pro Asp Ala
 115         120         125
Glu Met Lys Met Gly Tyr Leu Thr Ala Thr Gly Thr Gly Val Lys Lys
 130         135         140
Asp Tyr Gln Glu Ala Ile Gln Trp Tyr Gln Arg Ala Ala Glu His Gly
 145         150         155         160
Asp Ser Ala Ala Tyr Ala Gln Ile Gly Leu Phe Tyr Thr Leu Gly Asn
 165         170         175
Gly Val Lys Lys Asp Val Asn Arg Ala Val Gln Tyr Tyr Ile Met Gly
 180         185         190
Ala Gln Lys Gly Asp Ala Arg Ala Gln Ala Phe Leu Gly Lys Ala Tyr
 195         200         205
Ala Leu Gly Arg Gly Ile Gln Pro Asp Ser Glu Lys Ala Leu Tyr Trp
 210         215         220
Tyr Lys Thr Ala Ala Arg Asn Gly Asn Val Asn Ala Met Lys Glu Leu
 225         230         235         240
Gly Ser Ile Tyr Ala Lys Gly Arg Leu Gly Val Lys Pro Asp Gln Gln
 245         250         255
Glu Ala Gln Arg Trp Asn Asp Met Ala Arg Lys Ala Glu Gln Lys Asn
 260         265         270

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<210> SEQ ID NO 19

<211> LENGTH: 816

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 19

```

atgaaaattt cttacaagct atttctttct ttcattttta tgctgtgttc ttcgctgtt    60
tttgctgaca atgctcttac agggatcgag ttatacaaag caaaaaaata tgaacaggcc    120
atgacccacc tcctgacgcc tgatgcccag aaaaaccctg cagcacaaaa tcttattgga    180
tatctctatg ataagggctt aggtgtagaa aaaaacgctg aaatagccaa tcaatgggat    240
cttaaagcag ctgaacaggg atttgccaaa gctcaattca accttgact ctcttatgaa    300
aagggctactg gcatttcaaa aaatatgggt gaagctgtca aatggtatcg caaagcagct    360
gaacaaaatc acgccaaagc tgaatgaaa atgggggtatc tcacagtaga aggtatcggt    420
actcaaaaga attacaaaga agccttgcaa tggtatcggc gcgcagcaga acatgggtgat    480
aatagggctt atgcagacat tggcctcttc tatgatcagg gaaacgggtg caaaaaagac    540
cccaaccggg ctgtccagta ttacatcatg ggtgcagaaa agggcgatgg cgaagcacag    600
ctttttctcg cggattgcta cgcgaaagca agcgggattc cttatgatgc cgatcgcgcc    660
ttgtattggt acaaggaatc cgccaaaaac ggaaatatca ctgcatgaa ggtgtgtgctc    720
ggcatttaca aacttggcc aattgggtata gagaagaatc cggaaaaatc cgcgactcgg    780
cttgagatgg ccaaacaaaa agaagctcag ccatga                                816

```

<210> SEQ ID NO 20

<211> LENGTH: 271

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 20

```

Met Lys Ile Ser Tyr Lys Leu Phe Leu Ser Phe Ile Leu Met Leu Cys
 1           5           10          15
Ser Ser Ala Val Phe Ala Asp Asn Ala Leu Thr Gly Ile Glu Leu Tyr
          20          25          30
Lys Ala Lys Lys Tyr Glu Gln Ala Met Thr His Leu Met Thr Pro Asp
          35          40          45
Ala Gln Lys Asn Pro Ala Ala Gln Asn Leu Ile Gly Tyr Leu Tyr Asp
          50          55          60
Lys Gly Leu Gly Val Glu Lys Asn Ala Glu Ile Ala Asn Gln Trp Tyr
 65          70          75          80
Leu Lys Ala Ala Glu Gln Gly Phe Ala Lys Ala Gln Phe Asn Leu Gly
          85          90          95
Leu Ser Tyr Glu Lys Gly Thr Gly Ile Ser Lys Asn Met Val Glu Ala
          100         105         110
Val Lys Trp Tyr Arg Lys Ala Ala Glu Gln Asn His Ala Lys Ala Glu
          115         120         125
Met Lys Met Gly Tyr Leu Thr Val Glu Gly Ile Gly Thr Gln Lys Asn
          130         135         140
Tyr Lys Glu Ala Leu Gln Trp Tyr Arg Arg Ala Ala Glu His Gly Asp
          145         150         155         160
Asn Arg Ala Tyr Ala Asp Ile Gly Leu Phe Tyr Asp Gln Gly Asn Gly
          165         170         175

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Val Lys Lys Asp Pro Asn Arg Ala Val Gln Tyr Tyr Ile Met Gly Ala
 180 185 190

Glu Lys Gly Asp Gly Glu Ala Gln Leu Phe Leu Ala Asp Cys Tyr Ala
 195 200 205

Lys Ala Ser Gly Ile Pro Tyr Asp Ala Asp Arg Ala Leu Tyr Trp Tyr
 210 215 220

Lys Glu Ser Ala Lys Asn Gly Asn Ile Thr Ala Met Lys Val Leu Ser
 225 230 235 240

Gly Ile Tyr Lys Leu Gly Gln Leu Gly Ile Glu Lys Asn Pro Glu Lys
 245 250 255

Ser Arg His Trp Leu Glu Met Ala Lys Gln Lys Glu Ala Gln Pro
 260 265 270

<210> SEQ ID NO 21
 <211> LENGTH: 1614
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 21

atgtctgaaa caatgttggc tgttctggga tttcctgac tggatgagct tttgccgatg 60

gcacggctgt ttctgggtgc ttctttgctg gccgggtggt gtgtggggtc gatgagcttg 120

tttaggggcg gtttcttttc gggcagaggt gcccagggga acgtcgtcat gccgagggat 180

aaaatggatt ggaatttggg tcgtgttcgt caggggacgc tggctgtggt cggttgctctg 240

tttctggcga atgccgctg ggcgaacgat tattccgacg gcatgcagtt ctatcaggac 300

aaggattacg agaaagcttt ttcttctttc cagaaggcgg cgcacaaggg caatgcggcg 360

gcgcaatccg cactggctgc cctgtactat aacggggaag ggggtggagga agatgaggcc 420

gctgctgctg tgtgggtatc ggcgctgccc gaacatggcc gaacggatgc gcagtttgcg 480

cttgccgaga tgttcgaagc cggcgaaggg gtcaaacgcg attataaaaa agcggctttc 540

tggtataaaa aggcggcgga caagggccat ttgatggctg ccacgaagct gggcatcctc 600

tatatggaag gtcgctgagc caagcaagat gacgcgaaag cggctgcatt gctttctcat 660

gccgccaaac ggggcattgc gttggcgcag tcgaatctcg gtgttttcta tgcgagcggg 720

cgcggggttg aatccagtc gaaacgggcg ctggagtggt acaagaaggc ggctgttcag 780

ggaaattcac aagcccagtt ttcgctcggc aatatgtatg aggacggctc tgggtgttgaa 840

aagaatctgg cggtagcggc cgctctggtat cagaagtcgg ccgaacaggg gaatgccgag 900

gcccagaata atctgggacg ccttttatatg gaaggcggcg agtttgaggg gcgtgaagac 960

gaagcgttta tgtggttctc ccgtgcggcc gatcaggggt atgccgaggc gcaaacgaaat 1020

ctgggtgtct tgtattccta tgggcttgggt gtggacaagg atttgtccaa ggcgttttac 1080

tggtatcagc aggcggctga aaaggggcag gctgaagggg ctttttctcct ggctgaggcg 1140

tattacaagg gggaaaggtg tcaccgggac gacaacagcg cggttttctg gtatcagaaa 1200

gcgccgaagc tgggtgtccc agaaagtcag gacagggctg ggttgatgct gacgaacggg 1260

gtcggcgtca aacaggatta caagcaggca tatagctggt tcaggaaggg ggcctcag 1320

gggtatgccg aatcccagaa caatctgggg gtcttctgatg cccgtggact cggggtcgaa 1380

aaggattaca aacaggccgt ggctctggtat cgcaagggcg tgatgcagaa tctgcctcag 1440

gcgcagttca atctgggaac gatgtatttg caggggcatg gtgtcaaaca ggatgtcaaa 1500

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caggccagac actggtttac gaaggcggca gcgcaggggt tgccggaagc gcagagaagt 1560
 ctggacagga tgccgaaaaa cgggcagacc atcaatacgg atctctcgac ctga 1614

<210> SEQ ID NO 22

<211> LENGTH: 1074

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 22

Met Ser Glu Thr Met Leu Ala Val Leu Gly Phe Pro Asp Leu Asp Glu
 1 5 10 15
 Leu Leu Pro Met Ala Arg Leu Phe Leu Val Ser Ser Leu Leu Ala Gly
 20 25 30
 Cys Cys Val Gly Ser Met Ser Leu Phe Arg Gly Gly Phe Phe Ser Gly
 35 40 45
 Arg Gly Ala Gln Gly Asn Val Val Met Pro Arg Asp Lys Met Asp Trp
 50 55 60
 Asn Leu Gly Arg Val Arg Gln Gly Thr Leu Ala Val Val Gly Cys Leu
 65 70 75 80
 Phe Leu Ala Asn Ala Ala Trp Ala Asn Asp Tyr Ser Asp Gly Met Gln
 85 90 95
 Phe Tyr Gln Asp Lys Asp Tyr Glu Lys Ala Phe Ser Ser Phe Gln Lys
 100 105 110
 Ala Ala Asp Lys Gly Asn Ala Ala Ala Gln Ser Ala Leu Ala Ala Leu
 115 120 125
 Tyr Tyr Asn Gly Glu Gly Val Glu Glu Asp Glu Ala Ala Ala Ala Leu
 130 135 140
 Trp Tyr Ser Arg Ala Ala Glu His Gly Arg Thr Asp Ala Gln Phe Ala
 145 150 155 160
 Leu Gly Glu Met Phe Glu Ala Gly Glu Gly Val Lys Arg Asp Tyr Lys
 165 170 175
 Lys Ala Ala Phe Trp Tyr Lys Lys Ala Ala Asp Lys Gly His Leu Met
 180 185 190
 Ala Ala Thr Lys Leu Gly Ile Leu Tyr Met Glu Gly Arg Gly Val Lys
 195 200 205
 Gln Asp Asp Ala Lys Ala Ala Ala Leu Leu Ser His Ala Ala Lys Arg
 210 215 220
 Gly Ile Ala Leu Ala Gln Ser Asn Leu Gly Val Leu Tyr Ala Ser Gly
 225 230 235 240
 Arg Gly Val Glu Ser Ser Pro Lys Arg Ala Leu Glu Trp Tyr Lys Lys
 245 250 255
 Ala Ala Val Gln Gly Asn Ser Gln Ala Gln Phe Ser Leu Gly Asn Met
 260 265 270
 Tyr Glu Asp Gly Ser Gly Val Glu Lys Asn Leu Ala Val Ala Ala Ala
 275 280 285
 Trp Tyr Gln Lys Ser Ala Glu Gln Gly Asn Ala Glu Ala Gln Asn Asn
 290 295 300
 Leu Gly Arg Leu Tyr Met Glu Gly Gly Glu Phe Glu Gly Arg Glu Asp
 305 310 315 320
 Glu Ala Phe Met Trp Phe Ser Arg Ala Ala Asp Gln Gly Tyr Ala Glu
 325 330 335
 Ala Gln Thr Asn Leu Gly Val Leu Tyr Ser Tyr Gly Leu Gly Val Asp

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340					345					350					
Lys	Asp	Leu	Ser	Lys	Ala	Phe	Tyr	Trp	Tyr	Gln	Gln	Ala	Ala	Glu	Lys
	355						360					365			
Gly	Gln	Ala	Glu	Gly	Ala	Phe	Phe	Leu	Ala	Glu	Ala	Tyr	Tyr	Lys	Gly
	370					375					380				
Glu	Gly	Val	His	Arg	Asp	Asp	Lys	Gln	Ala	Val	Phe	Trp	Tyr	Gln	Lys
	385					390					395				400
Ala	Ala	Lys	Leu	Gly	Val	Pro	Glu	Ser	Gln	Asp	Arg	Leu	Gly	Leu	Met
			405						410					415	
Leu	Thr	Asn	Gly	Val	Gly	Val	Lys	Gln	Asp	Tyr	Lys	Gln	Ala	Tyr	Ser
		420						425						430	
Trp	Phe	Arg	Lys	Ala	Ala	Arg	Gln	Gly	Tyr	Ala	Glu	Ser	Gln	Asn	Asn
		435					440						445		
Leu	Gly	Val	Leu	Tyr	Ala	Arg	Gly	Leu	Gly	Val	Glu	Lys	Asp	Tyr	Lys
	450					455					460				
Gln	Ala	Val	Ala	Trp	Tyr	Arg	Lys	Ala	Val	Met	Gln	Asn	Leu	Pro	Gln
	465					470					475				480
Ala	Gln	Phe	Asn	Leu	Gly	Thr	Met	Tyr	Leu	Gln	Gly	His	Gly	Val	Lys
			485						490					495	
Gln	Asp	Val	Lys	Gln	Ala	Arg	His	Trp	Phe	Thr	Lys	Ala	Ala	Ala	Gln
			500					505						510	
Gly	Leu	Pro	Glu	Ala	Gln	Arg	Ser	Leu	Asp	Arg	Met	Pro	Lys	Asn	Gly
		515					520					525			
Gln	Thr	Ile	Asn	Thr	Asp	Leu	Ser	Thr	Met	Ser	Glu	Thr	Met	Leu	Ala
	530					535					540				
Val	Leu	Gly	Phe	Pro	Asp	Leu	Asp	Glu	Leu	Leu	Pro	Met	Ala	Arg	Leu
	545					550					555				560
Phe	Leu	Val	Ser	Ser	Leu	Leu	Ala	Gly	Cys	Cys	Val	Gly	Ser	Met	Ser
			565						570					575	
Leu	Phe	Arg	Gly	Gly	Phe	Phe	Ser	Gly	Arg	Gly	Ala	Gln	Gly	Asn	Val
		580						585						590	
Val	Met	Pro	Arg	Asp	Lys	Met	Asp	Trp	Asn	Leu	Gly	Arg	Val	Arg	Gln
		595					600					605			
Gly	Thr	Leu	Ala	Val	Val	Gly	Cys	Leu	Phe	Leu	Ala	Asn	Ala	Ala	Trp
	610					615					620				
Ala	Asn	Asp	Tyr	Ser	Asp	Gly	Met	Gln	Phe	Tyr	Gln	Asp	Lys	Asp	Tyr
	625					630					635				640
Glu	Lys	Ala	Phe	Ser	Ser	Phe	Gln	Lys	Ala	Ala	Asp	Lys	Gly	Asn	Ala
			645						650					655	
Ala	Ala	Gln	Ser	Ala	Leu	Ala	Ala	Leu	Tyr	Tyr	Asn	Gly	Glu	Gly	Val
		660						665						670	
Glu	Glu	Asp	Glu	Ala	Ala	Ala	Ala	Leu	Trp	Tyr	Ser	Arg	Ala	Ala	Glu
		675					680					685			
His	Gly	Arg	Thr	Asp	Ala	Gln	Phe	Ala	Leu	Gly	Glu	Met	Phe	Glu	Ala
	690					695					700				
Gly	Glu	Gly	Val	Lys	Arg	Asp	Tyr	Lys	Lys	Ala	Ala	Phe	Trp	Tyr	Lys
	705					710					715				720
Lys	Ala	Ala	Asp	Lys	Gly	His	Leu	Met	Ala	Ala	Thr	Lys	Leu	Gly	Ile
			725						730					735	
Leu	Tyr	Met	Glu	Gly	Arg	Gly	Val	Lys	Gln	Asp	Asp	Ala	Lys	Ala	Ala
		740							745					750	

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Ala Leu Leu Ser His Ala Ala Lys Arg Gly Ile Ala Leu Ala Gln Ser
 755 760 765

Asn Leu Gly Val Leu Tyr Ala Ser Gly Arg Gly Val Glu Ser Ser Pro
 770 775 780

Lys Arg Ala Leu Glu Trp Tyr Lys Lys Ala Ala Val Gln Gly Asn Ser
 785 790 795 800

Gln Ala Gln Phe Ser Leu Gly Asn Met Tyr Glu Asp Gly Ser Gly Val
 805 810 815

Glu Lys Asn Leu Ala Val Ala Ala Ala Trp Tyr Gln Lys Ser Ala Glu
 820 825 830

Gln Gly Asn Ala Glu Ala Gln Asn Asn Leu Gly Arg Leu Tyr Met Glu
 835 840 845

Gly Gly Glu Phe Glu Gly Arg Glu Asp Glu Ala Phe Met Trp Phe Ser
 850 855 860

Arg Ala Ala Asp Gln Gly Tyr Ala Glu Ala Gln Thr Asn Leu Gly Val
 865 870 875 880

Leu Tyr Ser Tyr Gly Leu Gly Val Asp Lys Asp Leu Ser Lys Ala Phe
 885 890 895

Tyr Trp Tyr Gln Gln Ala Ala Glu Lys Gly Gln Ala Glu Gly Ala Phe
 900 905 910

Phe Leu Ala Glu Ala Tyr Tyr Lys Gly Glu Gly Val His Arg Asp Asp
 915 920 925

Lys Gln Ala Val Phe Trp Tyr Gln Lys Ala Ala Lys Leu Gly Val Pro
 930 935 940

Glu Ser Gln Asp Arg Leu Gly Leu Met Leu Thr Asn Gly Val Gly Val
 945 950 955 960

Lys Gln Asp Tyr Lys Gln Ala Tyr Ser Trp Phe Arg Lys Ala Ala Arg
 965 970 975

Gln Gly Tyr Ala Glu Ser Gln Asn Asn Leu Gly Val Leu Tyr Ala Arg
 980 985 990

Gly Leu Gly Val Glu Lys Asp Tyr Lys Gln Ala Val Ala Trp Tyr Arg
 995 1000 1005

Lys Ala Val Met Gln Asn Leu Pro Gln Ala Gln Phe Asn Leu Gly
 1010 1015 1020

Thr Met Tyr Leu Gln Gly His Gly Val Lys Gln Asp Val Lys Gln
 1025 1030 1035

Ala Arg His Trp Phe Thr Lys Ala Ala Ala Gln Gly Leu Pro Glu
 1040 1045 1050

Ala Gln Arg Ser Leu Asp Arg Met Pro Lys Asn Gly Gln Thr Ile
 1055 1060 1065

Asn Thr Asp Leu Ser Thr
 1070

<210> SEQ ID NO 23
 <211> LENGTH: 588
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 23

atggttcata caggtttttc cgggcagtcc cgtccgtttt ccgatggat ggggactttt 60

ctgtttctgg tgttttgcc a gctggcattg gctgcccgtg c gatggggc ggaccggaa 120

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aagaaagcgg tggcttcaaa cgggcagctt tcgaaaaatg tggcatccgg ggatgaggat 180
gtgttgcgcg atctgatgga tttgaagagc aatgcggatt cgggtgatgt gtcgcccag 240
tttgagttga gccgcccgta cttgaacggg gatggccttg agcagaacga tgatgaggca 300
atccgctggc tccgatggc ggcggaaggt ggtttgccga gggcgcaggc gggctctggc 360
tggatgtatg cggcgggcag gggggggaat aaggatgaga cgctgtcttt ttcttggtat 420
gaacgggcgg cggttgccg ttttctgtg gcgcagtata tgctgggccc ttattatgaa 480
aagggtatcg gcgtcgccaa agaccgtgtg ctggctaaag agtggtatga aaaggcggca 540
gcgcagggta atgagaaagc gaagaagcgg ttgcaggact ggaaatga 588

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<210> SEQ ID NO 24
<211> LENGTH: 195
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

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<400> SEQUENCE: 24

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```

Met Val His Thr Gly Phe Ser Gly Gln Ser Arg Pro Phe Ser Gly Trp
1          5          10          15
Met Gly Thr Phe Leu Phe Leu Val Phe Cys Gln Leu Ala Leu Ala Ala
20          25          30
Gly Ala Met Ala Ala Asp Pro Glu Lys Lys Ala Val Ala Ser Asn Gly
35          40          45
Gln Leu Ser Lys Asn Val Ala Ser Gly Asp Glu Asp Val Leu Arg Asp
50          55          60
Leu Met Asp Leu Lys Ser Asn Ala Asp Ser Gly Asp Val Ser Ala Gln
65          70          75          80
Phe Glu Leu Ser Arg Arg Tyr Leu Asn Gly Asp Gly Leu Glu Gln Asn
85          90          95
Asp Asp Glu Ala Ile Arg Trp Leu Arg Met Ala Ala Glu Gly Gly Leu
100         105         110
Pro Arg Ala Gln Ala Gly Leu Gly Trp Met Tyr Ala Ala Gly Arg Gly
115         120         125
Val Asn Lys Asp Glu Thr Leu Ser Phe Ser Trp Tyr Glu Arg Ala Ala
130         135         140
Val Ala Gly Phe Pro Val Ala Gln Tyr Met Leu Gly Arg Tyr Tyr Glu
145         150         155         160
Lys Gly Ile Gly Val Ala Lys Asp Arg Val Leu Ala Lys Glu Trp Tyr
165         170         175
Glu Lys Ala Ala Ala Gln Gly Asn Glu Lys Ala Lys Lys Arg Leu Gln
180         185         190
Asp Trp Lys
195

```

```

<210> SEQ ID NO 25
<211> LENGTH: 1200
<212> TYPE: DNA
<213> ORGANISM: Oxalobacter formigenes

```

```

<400> SEQUENCE: 25

```

```

atgcgtagaa aactgttctt cttcatttgt ttattttgtt ctattacagc ttcagtctct 60
tttgctgaaa cgaaaatcaa tgaatcgttt ccggaattg aacggtcggg ggaagaggca 120
aagagagaaa cggctatcaa gctctataat ctgggcgtgg aatatgcaaa aggcagtcgt 180

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gtcgaaaaag atcgtaaaaa agccaattcc tatttcagac aagctgctga aataggactg 240
cctgaagcgc agtacaatct gggacgagcg tatttcgatg gcgatggtct ggaggtagac 300
aggaaagcag ctattgaatg gtacaaaaag gcagcagaac agggatttgc acaggctcaa 360
tacaacttgg gcgtaatcta ccaaaatggt ttgggcatca aacaggattt tgattcggtc 420
gtccagtggg atgagagagc ggcaaatcag ggatttgat tagcccaata caatcttgga 480
atggtatata taactggagc aggcgttggc aagaatccga aacgagggat tttgtggttg 540
cgcaaggctg ctgaaggggg ttacggtcag gcgcagcata atcttggaac cgtttattac 600
gagggtattg gcgtcagaaa aaattatccg gaagcgggtc aatggttcgc caaagctgct 660
aaacaagagc ttggcatggc gcaatacaat ctggggatgg cttactatca tggagagggc 720
gtcaaaaaaa atcctcagaa agcgggttca tgggtgaaaa aagcagcaaa acaaaatctt 780
cttatagctc aggctagtct tggctatata tatgttacgg acaggaattt caaaaataat 840
ctggcagaag gaattttctg gacaaaaaaa gcctccgcat atggtaatgc aagggtcag 900
gcgacacttg gtattgcata tcttattgga aagggtgtag aaaaaaatat tccagaaggc 960
gtttcgtgga taaaaaaagc agcgagacag ggtaattatc cggctcaaag catgcttgct 1020
tcctgttatg aaaatgggat tggcgtaaag caaaacaagg tattggctta tgcactttat 1080
ttacactcat ctccctatac ggaagttgct atggaagaac ggcaaaatct tgaaaaaaag 1140
ctaagcaaag atgaaattgt aaaagcccca tccatcaaca tcgaaaaact tttcgaatga 1200

```

<210> SEQ ID NO 26

<211> LENGTH: 399

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 26

```

Met Arg Arg Lys Leu Phe Phe Phe Ile Cys Leu Phe Cys Ser Ile Thr
 1           5           10           15
Ala Ser Val Ser Phe Ala Glu Thr Lys Ile Asn Glu Ser Phe Pro Glu
          20           25           30
Ile Glu Arg Ser Gly Glu Glu Ala Lys Arg Glu Thr Ala Ile Lys Leu
          35           40           45
Tyr Asn Leu Gly Val Glu Tyr Ala Lys Gly Ser Arg Val Glu Lys Asp
          50           55           60
Arg Lys Lys Ala Asn Ser Tyr Phe Arg Gln Ala Ala Glu Ile Gly Leu
          65           70           75           80
Pro Glu Ala Gln Tyr Asn Leu Gly Arg Ala Tyr Phe Asp Gly Asp Gly
          85           90           95
Leu Glu Val Asp Arg Lys Ala Ala Ile Glu Trp Tyr Lys Lys Ala Ala
          100          105          110
Glu Gln Gly Phe Ala Gln Ala Gln Tyr Asn Leu Gly Val Ile Tyr Gln
          115          120          125
Asn Gly Leu Gly Ile Lys Gln Asp Phe Asp Ser Ala Val Gln Trp Tyr
          130          135          140
Glu Arg Ala Ala Asn Gln Gly Phe Val Leu Ala Gln Tyr Asn Leu Gly
          145          150          155          160
Met Leu Tyr Ile Thr Gly Ala Gly Val Gly Lys Asn Pro Lys Arg Gly
          165          170          175

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Ile Leu Trp Leu Arg Lys Ala Ala Glu Gly Gly Tyr Gly Gln Ala Gln
 180 185 190

His Asn Leu Gly Thr Val Tyr Tyr Glu Gly Ile Gly Val Arg Lys Asn
 195 200 205

Tyr Pro Glu Ala Val Gln Trp Phe Ala Lys Ala Ala Lys Gln Glu Leu
 210 215 220

Gly Met Ala Gln Tyr Asn Leu Gly Met Ala Tyr Tyr His Gly Glu Gly
 225 230 235 240

Val Lys Lys Asn Pro Gln Lys Ala Val Ser Trp Leu Lys Lys Ala Ala
 245 250 255

Lys Gln Asn Leu Leu Ile Ala Gln Ala Ser Leu Gly Tyr Ile Tyr Val
 260 265 270

Thr Asp Arg Asn Phe Lys Asn Asn Leu Ala Glu Gly Ile Phe Trp Thr
 275 280 285

Lys Lys Ala Ser Ala Tyr Gly Asn Ala Arg Ala Gln Ala Thr Leu Gly
 290 295 300

Ile Ala Tyr Leu Ile Gly Lys Gly Val Glu Lys Asn Ile Pro Glu Gly
 305 310 315 320

Val Ser Trp Ile Lys Lys Ala Ala Arg Gln Gly Asn Tyr Pro Ala Gln
 325 330 335

Ser Met Leu Ala Ser Cys Tyr Glu Asn Gly Ile Gly Val Lys Gln Asn
 340 345 350

Lys Val Leu Ala Tyr Ala Leu Tyr Leu His Ser Ser Pro Tyr Thr Glu
 355 360 365

Val Ala Met Glu Glu Arg Gln Asn Leu Glu Lys Lys Leu Ser Lys Asp
 370 375 380

Glu Ile Val Lys Ala Arg Ser Ile Asn Ile Glu Lys Leu Phe Glu
 385 390 395

<210> SEQ ID NO 27
 <211> LENGTH: 909
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 27

```

atgaaaaaaaa atatttcact catctttttt atatttattt ctttattttc tttgactgga      60
tgtcaaaaatg aagataataa agagttaaca gatgaaaaaa caggaattga atactataaa      120
aaagtgact   acgaaaaatc attatccttt ctgaagaaag ccgctgattc cgggagtac      180
aaagcttccc ggtatttggg aaaaatgat  cagtatggca aagtggtcga taaagattat      240
ccgctttctt tcaagtggta tttaaacgca gccgaaaaag gagacaaaga atcatccggt      300
atggttggcg ccagttacta tttgggacaa ggcgtaaac  aagactacaa ggaatcgttt      360
agatggttgt taaaggcatc tgaaaaaatt gatgaaaaaa aaccaactga acgggacgga      420
aaactcatgc ttcttttggc taatttatat tttacgggaa aaggtacact tcaggatttc      480
agtgagttag caaaatgggc gagaagagct gccgaattgg gaaattccga gtcacaagcc      540
atgcttgcac tcttcctcta ttcggggcaa ggcattttac agaacagaac ggaagcgaaa      600
atctgggcag aaaagtctgc cgggcagggg gatagccttg gacaagtcat aatgggaatg      660
ctttatcagt atggcggcgg aacagatgaa ccggatatga agaaagcaat cgactggtac      720
gaaaaatcag ctgaaaaggg aaatccgatt gcacaatatc aactggcaac cctttacgaa      780
    
```

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aatgggaatg gcttgccgaa agatctggaa aaagcaaaa actattatga gcaatcggct 840
aaaagccaat cggaaatacc agtgaaagct ttggcggaat tcaaggccaa acagaaaaga 900
caaaattaa 909

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<210> SEQ ID NO 28
<211> LENGTH: 302
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

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<400> SEQUENCE: 28

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```

Met Lys Lys Asn Ile Ser Leu Ile Phe Phe Ile Phe Ile Ser Leu Phe
1           5           10           15
Ser Leu Thr Gly Cys Gln Asn Glu Asp Asn Lys Glu Leu Thr Asp Glu
20          25          30
Lys Thr Gly Ile Glu Tyr Tyr Lys Lys Gly Asp Tyr Glu Lys Ser Leu
35          40          45
Ser Phe Leu Lys Lys Ala Ala Asp Ser Gly Ser Ile Lys Ala Ser Arg
50          55          60
Tyr Leu Gly Lys Met Tyr Gln Tyr Gly Lys Gly Val Asp Lys Asp Tyr
65          70          75          80
Pro Leu Ser Phe Lys Trp Tyr Leu Asn Ala Ala Glu Lys Gly Asp Lys
85          90          95
Glu Ser Ser Gly Met Val Gly Ala Ser Tyr Tyr Leu Gly Gln Gly Val
100         105        110
Lys Gln Asp Tyr Lys Glu Ser Phe Arg Trp Leu Leu Lys Ala Ser Glu
115        120        125
Lys Ile Asp Glu Lys Lys Pro Thr Glu Arg Asp Gly Lys Leu Met Leu
130        135        140
Leu Leu Ala Asn Leu Tyr Phe Thr Gly Lys Gly Thr Leu Gln Asp Phe
145        150        155        160
Ser Glu Ser Ala Lys Trp Ala Arg Arg Ala Ala Glu Leu Gly Asn Ser
165        170        175
Glu Ser Gln Ala Met Leu Ala Phe Phe Leu Tyr Ser Gly Gln Gly Ile
180        185        190
Leu Gln Asn Arg Thr Glu Ala Lys Ile Trp Ala Glu Lys Ser Ala Gly
195        200        205
Gln Gly Asp Ser Leu Gly Gln Val Ile Met Gly Met Leu Tyr Gln Tyr
210        215        220
Gly Gly Gly Thr Asp Glu Pro Asp Met Lys Lys Ala Ile Asp Trp Tyr
225        230        235        240
Glu Lys Ser Ala Glu Lys Gly Asn Pro Ile Ala Gln Tyr Gln Leu Ala
245        250        255
Thr Leu Tyr Glu Asn Gly Asn Gly Leu Pro Lys Asp Leu Glu Lys Ala
260        265        270
Lys Tyr Tyr Tyr Glu Gln Ser Ala Lys Ser Gln Ser Glu Ile Pro Val
275        280        285
Lys Ala Leu Ala Glu Phe Lys Ala Lys Gln Lys Arg Gln Asn
290        295        300

```

```

<210> SEQ ID NO 29
<211> LENGTH: 885
<212> TYPE: DNA
<213> ORGANISM: Oxalobacter formigenes

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-continued

<400> SEQUENCE: 29

```

atgatgaata aaatatttca ggttttttat atcgtaatc tttctttttt tattagtcc      60
tgccacgaat cagacaacag aatcacagaa aataatgggt tcaatctcta caaggcaaa      120
caatatgaaa tggcattgcc tttacttgaa aaggctgcca atgcaggaga tcctcaggcg      180
ccattctatc tcggcataat gttcgatgaa ggatcaggag taatcaaaga tcagaaaaaa      240
tcatttgaat ggtttgaaaa agcggcaaaa aatggtaaca ctgacgcggt ttttgttatt      300
ggcagcaggt atttatatgg ttctggcggt gaaaaagatt ataaagaagc cctgaaatgg      360
tataaaagga gtgttgaaga gggcaaaaaa gacgacaaga caatctatct catgatcgga      420
tcgatgtact acaatgggtt ggttaccttg aaagatacca gcgaggcagc caaatgggat      480
gaaaaagcgg cagaaaaagg agatgctttt tcacaggcaa tgctcgccat gcaatattac      540
agtggtcagg gtattttgac gaatatgaa aaagccagat actgggccga aaaatccgcg      600
gaacaggatt acgatgccgg acaaatgatg atggggattc tgagccagta tggaacacct      660
gaaccagaca tgaagccgc aattgactgg tatgaaaaag ccgccagaca gggaaacccg      720
attgcacagt ttctattggc gagaagtat gaaaatggaa acggcgtgcc aaaagatctg      780
gaaaaagccc atgcttatta taaacaggct gccggtggca tgaagtcgga tgacctggcc      840
agggagttta tggaaattga agcaagacag aaaagacaga aataa                      885

```

<210> SEQ ID NO 30

<211> LENGTH: 294

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 30

```

Met Met Asn Lys Ile Phe Gln Val Phe Tyr Ile Val Ile Leu Ser Phe
 1          5          10          15
Phe Ile Ser Ser Cys His Glu Ser Asp Asn Arg Ile Ser Glu Asn Asn
 20          25          30
Gly Val Asn Leu Tyr Lys Ala Lys Gln Tyr Glu Met Ala Leu Pro Leu
 35          40          45
Leu Glu Lys Ala Ala Asn Ala Gly Asp Pro Gln Ala Pro Phe Tyr Leu
 50          55          60
Gly Ile Met Phe Asp Glu Gly Ser Gly Val Ile Lys Asp Gln Lys Lys
 65          70          75          80
Ser Phe Glu Trp Phe Glu Lys Ala Ala Lys Asn Gly Asn Thr Asp Ala
 85          90          95
Phe Phe Val Ile Gly Ser Arg Tyr Leu Tyr Gly Ser Gly Val Glu Lys
100          105          110
Asp Tyr Lys Glu Ala Leu Lys Trp Tyr Lys Arg Ser Val Glu Glu Gly
115          120          125
Lys Lys Asp Asp Lys Thr Ile Tyr Phe Met Ile Gly Ser Met Tyr Tyr
130          135          140
Asn Gly Leu Gly Thr Leu Lys Asp Thr Ser Glu Ala Ala Lys Trp Tyr
145          150          155          160
Glu Lys Ala Ala Glu Lys Gly Asp Ala Phe Ser Gln Ala Met Leu Ala
165          170          175
Met Gln Tyr Tyr Ser Gly Gln Gly Ile Leu Thr Asn Met Glu Lys Ala
180          185          190

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Arg Tyr Trp Ala Glu Lys Ser Ala Glu Gln Asp Tyr Asp Ala Gly Gln
 195 200 205

Met Met Met Gly Ile Leu Ser Gln Tyr Gly Thr Pro Glu Pro Asp Met
 210 215 220

Lys Ala Ala Ile Asp Trp Tyr Glu Lys Ala Ala Arg Gln Gly Asn Pro
 225 230 235 240

Ile Ala Gln Phe Leu Leu Ala Arg Ser Tyr Glu Asn Gly Asn Gly Val
 245 250 255

Pro Lys Asp Leu Glu Lys Ala His Ala Tyr Tyr Lys Gln Ala Ala Gly
 260 265 270

Gly Met Lys Ser Asp Asp Leu Ala Arg Glu Phe Met Glu Phe Glu Ala
 275 280 285

Arg Gln Lys Arg Gln Lys
 290

<210> SEQ ID NO 31
 <211> LENGTH: 813
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 31

```

atgctgggat atttgcacg tgaggggat ggagtcaagc aggattatca aaaggcgttt    60
tttctctatc ttgaaggagc aaaactgggc gacgcaaaaa gccagttcgg tttgggtttt    120
atgtatgagg ggggattatt cgtcaaacag gattatgcca aggcaaaaac atggtatgag    180
tattcatcca atcagggata tctgtctgcg atgaataatc ttggatcgct ttatgacgat    240
gaaaacactg gctttaaaaa cgagaaaaatc gcatttgaat ggatattgaa agctgcccac    300
aaagataacc caactgctca atttaatatc ggtttttttt acgaaaaagg cacaggaacc    360
aaaaaagact atgccgaagc ccgaaagtgg tatgaaaaag cagtcgatgca gggatatcct    420
cgggccaaa gcaatctggc aaatctctat ctgatggaa aaggtggccc caaagaccag    480
caaaagggcg ttgcctgat aaaagaagcg gcgaacgagg aatcgaaagc cgcacaatac    540
acactggcaa acctctacgc cgatggcgaa ggcgttccgc aaagcgatga acaggccggt    600
tactggttcc acaaggccgc tgaaaacgac agcgcgctgg ccatggacat gctcgccaaa    660
gcctacctga acgaaaaata cggactgccg aaaagcccga cgaaatggga ctactggcaa    720
aagagggcag ccgaaacgcg cgcccgtaca cacgaagtcg atccttcaag ctggacaata    780
ttcgactgga tcaaatatac gttaggcaat tga                                813
    
```

<210> SEQ ID NO 32
 <211> LENGTH: 270
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 32

Met Leu Gly Tyr Leu Tyr Arg Glu Gly Tyr Gly Val Lys Gln Asp Tyr
 1 5 10 15

Gln Lys Ala Phe Phe Leu Tyr Leu Glu Gly Ala Lys Leu Gly Asp Ala
 20 25 30

Lys Ser Gln Phe Gly Leu Gly Phe Met Tyr Glu Gly Gly Leu Phe Val
 35 40 45

Lys Gln Asp Tyr Ala Lys Ala Lys Thr Trp Tyr Glu Tyr Ser Ser Asn

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50		55				60									
Gln	Gly	Tyr	Leu	Ser	Ala	Met	Asn	Asn	Leu	Gly	Ser	Leu	Tyr	Asp	Asp
65					70					75					80
Glu	Asn	Thr	Gly	Phe	Lys	Asn	Glu	Lys	Ile	Ala	Phe	Glu	Trp	Ile	Leu
				85					90					95	
Lys	Ala	Ala	Gln	Lys	Asp	Asn	Pro	Thr	Ala	Gln	Phe	Asn	Ile	Gly	Phe
			100					105					110		
Phe	Tyr	Glu	Lys	Gly	Thr	Gly	Thr	Lys	Lys	Asp	Tyr	Ala	Glu	Ala	Arg
		115					120					125			
Lys	Trp	Tyr	Glu	Lys	Ala	Val	Met	Gln	Gly	Tyr	Leu	Pro	Ala	Lys	Ala
	130					135					140				
Asn	Leu	Ala	Asn	Leu	Tyr	Leu	Asp	Gly	Lys	Gly	Gly	Pro	Lys	Asp	Gln
145					150					155					160
Gln	Lys	Gly	Val	Ala	Leu	Ile	Lys	Glu	Ala	Ala	Asn	Glu	Glu	Ser	Lys
				165					170						175
Ala	Ala	Gln	Tyr	Thr	Leu	Ala	Asn	Leu	Tyr	Ala	Asp	Gly	Glu	Gly	Val
			180					185					190		
Pro	Gln	Ser	Asp	Glu	Gln	Ala	Val	Tyr	Trp	Phe	His	Lys	Ala	Ala	Glu
		195					200					205			
Asn	Asp	Ser	Ala	Leu	Ala	Met	Asp	Met	Leu	Ala	Lys	Ala	Tyr	Leu	Asn
	210					215					220				
Gly	Lys	Tyr	Gly	Leu	Pro	Lys	Ser	Pro	Thr	Lys	Trp	Asp	Tyr	Trp	Gln
225					230					235					240
Lys	Arg	Ala	Ala	Glu	Thr	Arg	Ala	Arg	Thr	His	Glu	Val	Asp	Pro	Ser
				245					250					255	
Ser	Trp	Thr	Ile	Phe	Asp	Trp	Ile	Lys	Tyr	Lys	Leu	Gly	Asn		
			260					265					270		

<210> SEQ ID NO 33
 <211> LENGTH: 726
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 33

```

ttgatatttt tcagaaaaaa aagcgacgct ggagaaagt gcctgagaga tactatgaaa      60
agccgttttt accgcttcaa acatcatttt tcttatatag aaaaatgggt gattttcaat    120
gttgtttttt tgaccatttc tgcttcgctc tttgcacaaa cgcatccgat aaccggaaca    180
tctgatcccg acagtcaaca tactcagggt catcaaaaac tctgtctctc cgcagaaaat    240
ggcgatcagg atgcgcagta caatctcgga cgtatatacc tgcaaggaaa aggaacacgt    300
caggattatc aggccgcccg taaatgggtc atgcgcgccg ccgaaaaaga ggatgccgga    360
gcccataaca atctgggaaa tatttatcaa aaaggacaag ggattcaaca ggattgcaaa    420
aaagcctttt tctggtacaa aaaggcagct gcaaaattct atgcgccggc tcagtaagct    480
cttgcaagc tttactcaag tggatgtggt gtcaatcaaa attcatataa atcgacagaa    540
tggattctta aggcagccta taatggcatg ccggaagccc aatttcagat aggatatacg    600
tatctcacag gctatggaat tcaagtcgat aaaaacaagg cctatgaatg gctgttgaag    660
gcgcaaaaac aggataaacc agcggcaata aaggttctga aaacaaatth tccacagaa    720
atatga                                           726
    
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<210> SEQ ID NO 34
 <211> LENGTH: 241
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 34

Met Ile Phe Phe Arg Lys Lys Ser Asp Ala Gly Glu Ser Cys Leu Arg
 1 5 10 15
 Asp Thr Met Lys Ser Arg Phe Tyr Arg Phe Lys His His Phe Ser Tyr
 20 25 30
 Ile Glu Lys Trp Val Ile Phe Asn Val Val Phe Leu Thr Ile Ser Ala
 35 40 45
 Ser Ser Phe Ala Gln Thr His Pro Ile Thr Gly Thr Ser Asp Pro Asp
 50 55 60
 Ser Gln His Thr Gln Val His Gln Lys Leu Leu Ser Leu Ala Glu Asn
 65 70 75 80
 Gly Asp Gln Asp Ala Gln Tyr Asn Leu Gly Arg Ile Tyr Leu Gln Gly
 85 90 95
 Lys Gly Thr Arg Gln Asp Tyr Gln Ala Ala Arg Lys Trp Phe Met Arg
 100 105 110
 Ala Ala Glu Lys Glu Asp Ala Gly Ala Gln Tyr Asn Leu Gly Asn Ile
 115 120 125
 Tyr Gln Lys Gly Gln Gly Ile Gln Gln Asp Cys Lys Lys Ala Phe Phe
 130 135 140
 Trp Tyr Lys Lys Ala Ala Ala Lys Phe Tyr Ala Pro Ala Gln Tyr Ala
 145 150 155 160
 Leu Gly Lys Leu Tyr Ser Ser Gly Cys Gly Val Asn Gln Asn Ser Tyr
 165 170 175
 Lys Ser Thr Glu Trp Ile Leu Lys Ala Ala Tyr Asn Gly Met Pro Glu
 180 185 190
 Ala Gln Phe Gln Ile Gly Tyr Arg Tyr Leu Thr Gly Tyr Gly Ile Gln
 195 200 205
 Val Asp Lys Asn Lys Ala Tyr Glu Trp Leu Leu Lys Ala Ala Lys Gln
 210 215 220
 Asp Asn Pro Ala Ala Ile Lys Val Leu Lys Thr Asn Phe Ser Thr Glu
 225 230 235 240
 Ile

<210> SEQ ID NO 35
 <211> LENGTH: 453
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 35

atgttttctt ttctgaaaaa agcggttccg gtcagtctcg ttctggtttc tgcgggggtt 60
 ttgtctgcct gtactacagg acaggaaatg acggccaagt tgacggcaca ggcggaaaag 120
 ggggatgtgg aagcgatggt cgagctggcg gaggtttatt gcggtggcaa aaacatcgaa 180
 caggacgacc agatttgcgg catgtggatg aaacgggcag ctgaaaaagg ccatgtccgt 240
 gcccagtata tgctcggcag aatgtatgaa ctgggtctgg gtatgagagc cgatccggta 300
 caggcataca agtggatag cttgtctgcc cccattatc atatgtccca gactggtgca 360
 gagactgtat atgccgtcat gacacctgtc cagcagtcag aagccagaaa agtggctgat 420

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gaggcgaaga aaaccattcc ttcaaaacaa taa

453

<210> SEQ ID NO 36

<211> LENGTH: 150

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 36

Met Phe Ser Phe Leu Lys Lys Ala Val Pro Val Ser Leu Val Leu Val
 1 5 10 15
 Ser Val Gly Val Leu Ser Ala Cys Thr Thr Gly Gln Glu Met Thr Ala
 20 25 30
 Lys Leu Thr Ala Gln Ala Glu Lys Gly Asp Val Glu Ala Met Val Glu
 35 40 45
 Leu Ala Glu Val Tyr Cys Gly Gly Lys Asn Ile Glu Gln Asp Asp Gln
 50 55 60
 Ile Cys Gly Met Trp Met Lys Arg Ala Ala Glu Lys Gly His Val Arg
 65 70 75 80
 Ala Gln Tyr Met Leu Gly Arg Met Tyr Glu Leu Gly Leu Gly Met Arg
 85 90 95
 Ala Asp Pro Val Gln Ala Tyr Lys Trp Tyr Ser Leu Ser Ala Pro His
 100 105 110
 Tyr His Met Ser Gln Thr Gly Ala Glu Thr Val Tyr Ala Val Met Thr
 115 120 125
 Pro Val Gln Gln Ser Glu Ala Arg Lys Val Ala Asp Glu Ala Lys Lys
 130 135 140
 Thr Ile Pro Ser Lys Gln
 145 150

<210> SEQ ID NO 37

<211> LENGTH: 1536

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 37

atggctaadc gagaagaatt acaactggtg cgtgaagcac gatccaacca tcacgaagcc 60
 cagctccagt tgggcaaact gtatttggtc ggaacaaaaa gcctgcccga aagtctgacg 120
 accgcccctgc actgggtaag ccgtgcccgc acacaatcca ataaggaagc gtgccttctg 180
 attggcgacc atattcctta tgaattgcc agaaattttc cggatcaggt ggcgatccag 240
 tcctgggtaca gaacagccct gacagaaggg cattatgagg ctggtctggt attggcccgg 300
 ctgattcttt cagatcccaa tcagatcgat gacaaaaagt atgccgaagc gattcgcatt 360
 ctcgaaacaa ttgcagatca tgatattgcc gaagcacaat gggtactggc tgaattggcc 420
 aaagatccga atgcccgccc atccattatt aacaatgccc ttaaatggac agcgagagcc 480
 gccgatgccg gcattgtoga tgcacagatc gccctgattg aacatgcctg ggaaaacagg 540
 gactaccocg tttttctgca gcattgocctg ccgattgccc gatcgattac tcaggcagtc 600
 cagcaggggtg atgtcattaa tattgacgaa caatcttccc gtttactggt tcgggtgccc 660
 caactgctcc tgaaacagga cggtcatgca tcagaagaga ttcagtccat gtgggaattg 720
 gccgccagac agaaaaatgc cgaagccgct ttttcattag gcctgtggta cgcgcgcatg 780
 aatgaagacg gtatccgctg cagtattggc gcggccgcga ccagtttcaa aaaagcgatt 840

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cgatggctga ctcaggccgg tgaacagggt ctggccaaag catggtacgc actttcactt 900
atttaccaga aggcggaatt ctcacagaga aatatgaatg atgcccaacg ttatctggaa 960
ttggcagcag atcttggaca tgcaacagcc cagtatgaaa gaggcacgca tgcgtggcgg 1020
gcacgccggg atgatgaatc aaatgacatt caggctgttt actggctgca gaaagcaaac 1080
ggaaacggaa aagcggacgc ggcagctgtc cttgataaaa ttgctttcaa ggcgaatcct 1140
gcatcatggg cggaagcgc acgggaatgc ctgactcagc aaattttcag cagccatcca 1200
tttctggcgg cccgtatoga actggctgcc gtattcggac tgacacgtcc ggaagcgtc 1260
ttgctcgata tccacaatgc cgaaaaggc cattgccttc tggctogatat tcgtgaattt 1320
tacagacgca gcaagcgsaa gctcatcctg attcaaactg gccaggaacg ccagattctc 1380
tcccgaatcg gacgtctttt tgaaaaagtg gattggcgcc tgaatggtcc cgagggtaat 1440
tacagacaac gtcaatatcg cctcaaaaac atgctcccg gcgcttatca ggaaaattca 1500
gatgatgaag aatatcgga aacagccgaa gcttag 1536

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<210> SEQ ID NO 38

<211> LENGTH: 511

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 38

```

Met Ala Asn Arg Glu Glu Leu Gln Leu Leu Arg Glu Ala Arg Ser Asn
1           5           10           15
His His Glu Ala Gln Leu Gln Leu Gly Lys Leu Tyr Leu Phe Gly Thr
20           25           30
Lys Ser Leu Pro Gln Ser Leu Thr Thr Ala Leu His Trp Leu Ser Arg
35           40           45
Ala Ala Thr Gln Ser Asn Lys Glu Ala Cys Leu Leu Ile Gly Asp His
50           55           60
Ile Pro Tyr Glu Ile Ala Arg Asn Phe Pro Asp Gln Val Ala Ile Gln
65           70           75           80
Ser Trp Tyr Arg Thr Ala Leu Thr Glu Gly His Tyr Glu Ala Gly Leu
85           90           95
Val Leu Ala Arg Leu Ile Leu Ser Asp Pro Asn Gln Ile Asp Asp Lys
100          105          110
Lys Tyr Ala Glu Ala Ile Arg Ile Leu Glu Thr Ile Ala Asp His Asp
115          120          125
Ile Ala Glu Ala Gln Trp Leu Leu Ala Glu Leu Ala Lys Asp Pro Asn
130          135          140
Ala Arg Pro Ser Ile Ile Asn Asn Ala Leu Lys Trp Thr Ala Arg Ala
145          150          155          160
Ala Asp Ala Gly Ile Val Asp Ala Gln Ile Ala Leu Ile Glu His Ala
165          170          175
Trp Glu Asn Arg Asp Tyr Pro Val Phe Leu Gln His Ala Leu Pro Ile
180          185          190
Ala Arg Ser Ile Thr Gln Ala Val Gln Gln Gly Asp Val Ile Asn Ile
195          200          205
Asp Glu Gln Ser Ser Arg Leu Leu Phe Arg Cys Gly Gln Leu Leu Leu
210          215          220
Lys Gln Asp Gly His Ala Ser Glu Glu Ile Gln Ser Met Trp Glu Leu
225          230          235          240

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Ala Ala Arg Gln Lys Asn Ala Glu Ala Ala Phe Ser Leu Gly Leu Trp
 245 250 255

Tyr Ala Arg Met Asn Glu Asp Gly Ile Arg Val Ser Ile Gly Ala Ala
 260 265 270

Ala Thr Ser Phe Lys Lys Ala Ile Arg Trp Leu Thr Gln Ala Gly Glu
 275 280 285

Gln Gly Leu Ala Lys Ala Trp Tyr Ala Leu Ser Leu Ile Tyr Gln Lys
 290 295 300

Ala Glu Phe Ser Gln Arg Asn Met Asn Asp Ala Gln Arg Tyr Leu Glu
 305 310 315 320

Leu Ala Ala Asp Leu Gly His Ala Thr Ala Gln Tyr Glu Arg Gly Met
 325 330 335

His Ala Trp Arg Ala Arg Arg Asp Asp Glu Ser Asn Asp Ile Gln Ala
 340 345 350

Val Tyr Trp Leu Gln Lys Ala Asn Gly Asn Gly Lys Ala Asp Ala Ala
 355 360 365

Ala Val Leu Asp Lys Ile Ala Phe Lys Ala Asn Pro Ala Ser Trp Ala
 370 375 380

Val Ser Ala Arg Glu Cys Leu Thr His Glu Ile Phe Ser Ser His Pro
 385 390 395 400

Phe Leu Ala Ala Arg Ile Glu Leu Ala Ala Val Phe Gly Leu Thr Arg
 405 410 415

Pro Glu Ala Leu Leu Leu Asp Ile His Asn Ala Asp Lys Gly His Cys
 420 425 430

Leu Leu Val Asp Ile Arg Glu Phe Tyr Arg Arg Ser Lys Arg Lys Leu
 435 440 445

Ile Leu Ile Gln Thr Gly Gln Glu Arg Gln Ile Leu Ser Arg Ile Gly
 450 455 460

Arg Leu Phe Glu Lys Val Asp Cys Gly Leu Asn Gly Pro Glu Gly Asn
 465 470 475 480

Tyr Arg Gln Arg Gln Tyr Arg Leu Lys Thr Met Leu Pro Ala Ala Tyr
 485 490 495

Gln Glu Asn Ser Asp Asp Glu Glu Tyr Arg Glu Thr Ala Glu Ala
 500 505 510

<210> SEQ ID NO 39

<211> LENGTH: 798

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 39

ttgaaaaaac ttaatttctt taagaagcaa cctgaactct ctatcaatga attagctttc 60
 aaagcaaggc aaggcgatct cgatgcttta aaacgattgc aagaagcagc cgagcaaat 120
 gatgcgaatg cccagaatag acttggggtt atatatgccc atggcaaagg catcccgaga 180
 aacgagaact tggctgctga caggttccaa aaagcagctg aattagaaaa cgctgaagca 240
 caggctaacc tagccgcctt atatagaaat tcattagtgt tcccacgtga taatgcgaag 300
 gttatttact gggctcagaa ggctgctgaa catggaaatc ctaggaggaca gaatattcta 360
 gggtttatgt atatgatcgg agaagtgta cagcaagatg acgccaagc agcttcttgg 420
 tatcaaaaag ccgctgagca gggattcgca ggagggcaga ggaatttagc gtttatgtat 480

-continued

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ctcaatggaa aggggtgttcc gcaagatgac gctacagcaa cttattggta tcaaaaagcg 540
gcaaatcaag gtgacataca agcgcaaaaa agtttaaaga tgattcaaga aaaaagtaaa 600
gaaatgagta actataattht aaatacggta caacaatcac ctccccaggc atcatctcat 660
cagccacttc ccaaagcatc accttcaaat gaaagaaatt taccaccttt aaaaaatatt 720
tttgacaac tgaagcagaa ttctgcaaag atgaaatctg ctctctccgg caacgaagaa 780
caggattacc tgaatatag 798

```

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<210> SEQ ID NO 40
<211> LENGTH: 265
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

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```

<400> SEQUENCE: 40

```

```

Met Glu Lys Leu Asn Phe Phe Lys Lys Gln Pro Glu Leu Ser Ile Asn
1          5          10          15
Glu Leu Ala Phe Lys Ala Arg Gln Gly Asp Leu Asp Ala Leu Lys Arg
20          25          30
Leu Gln Glu Ala Ala Glu Gln Asn Asp Ala Asn Ala Gln Asn Arg Leu
35          40          45
Gly Val Ile Tyr Ala Asp Gly Lys Gly Ile Pro Arg Asn Glu Asn Leu
50          55          60
Ala Ala Asp Arg Phe Gln Lys Ala Ala Glu Leu Glu Asn Ala Glu Ala
65          70          75          80
Gln Ala Asn Leu Ala Ala Leu Tyr Arg Asn Ser Leu Val Val Pro Arg
85          90          95
Asp Asn Ala Lys Val Ile Tyr Trp Ala Gln Lys Ala Ala Glu His Gly
100         105         110
Asn Pro Arg Gly Gln Asn Ile Leu Gly Phe Met Tyr Met Ile Gly Glu
115        120        125
Gly Val Gln Gln Asp Asp Ala Lys Ala Ala Ser Trp Tyr Gln Lys Ala
130        135        140
Ala Glu Gln Gly Phe Ala Gly Gly Gln Arg Asn Leu Ala Phe Met Tyr
145        150        155        160
Leu Asn Gly Lys Gly Val Pro Gln Asp Asp Ala Thr Ala Thr Tyr Trp
165        170        175
Tyr Gln Lys Ala Ala Asn Gln Gly Asp Ile Gln Ala Gln Lys Ser Leu
180        185        190
Lys Met Ile Gln Glu Lys Ser Lys Glu Met Ser Asn Tyr Asn Leu Asn
195        200        205
Thr Val Gln Gln Ser Pro Pro Gln Ala Ser Ser His Gln Pro Leu Pro
210        215        220
Lys Ala Ser Pro Ser Asn Glu Arg Asn Leu Ser Pro Leu Lys Asn Ile
225        230        235        240
Phe Asp Lys Leu Lys Gln Asn Ser Ala Lys Met Lys Ser Ala Pro Pro
245        250        255
Gly Asn Glu Glu Gln Asp Tyr Leu Lys
260          265

```

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<210> SEQ ID NO 41
<211> LENGTH: 888
<212> TYPE: DNA
<213> ORGANISM: Oxalobacter formigenes

```

-continued

<400> SEQUENCE: 41

```

atgaacagcc agactgccag aaaaatgttt tctttcgata tcagggcaaa ccattttttt    60
cgagcttttt tgctgcttgc catgtcactt cttgccggtg ttgcctttgc tgacgatttc    120
aacgatggtg tcagcgccta caaaagcggg aactaccagc aggcactgtc gctatttgag    180
gctggcgcga aaaaggacga cccgaaatcg acgatgccc tcggccttct gtataaaaac    240
ggcgtgatcg tcagaaaaga taccggacgt ggcctgaatc tgatcatgaa atctgccaat    300
cagggattcg cgagagccca gaattatctc ggagtcacct actacgatgg caatgaggtc    360
gaacaggact acaaggaagc gttcgactgg tatggcaagg cagccgttca gggttatcct    420
gatgcggaat acaatctggc cgtcatgtat ggtcttggga agggaacccg gcaggatttt    480
tccgaaacca tcaaatggct gcgcaaggcc gccatgcacc agcttctga agcccagtac    540
ggtcttggcg taatgtattc ccgaggactt ggtgtcgtga aaaacgatga acagtccgct    600
tactggtttt caaaggcagc gctggccggt tacctgaaag cgcaaaaaca actgggtatt    660
ttatattccg aaggaaaagg tctggaaaag gacgagaaaa aagcgttcca ctggttcgag    720
gccgctgcgc aaaaaggcta tgcaaaagcc caattcaatc tggcggtcac gtacgacaag    780
ggaattggcg ttgcaaaaga tgtgtccaaa gccattatgt ggtaccgcaa agcggctaca    840
caaggcaatg tggtgcaca gaaacggttg aaacaactgc attcctaa    888

```

<210> SEQ ID NO 42

<211> LENGTH: 295

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 42

```

Met Asn Ser Gln Thr Ala Arg Lys Met Phe Ser Phe Asp Ile Arg Ala
1          5          10          15
Asn His Phe Phe Arg Ala Phe Leu Leu Leu Ala Met Ser Leu Leu Ala
20        25        30
Gly Val Ala Phe Ala Asp Asp Phe Asn Asp Gly Val Ser Ala Tyr Lys
35        40        45
Ser Gly Asn Tyr Gln Gln Ala Leu Ser Leu Phe Glu Ala Gly Ala Lys
50        55        60
Lys Asp Asp Pro Lys Ser Thr Tyr Ala Leu Gly Leu Leu Tyr Lys Asn
65        70        75        80
Gly Val Ile Val Arg Lys Asp Ile Gly Arg Gly Leu Asn Leu Ile Met
85        90        95
Lys Ser Ala Asn Gln Gly Phe Ala Arg Ala Gln Asn Tyr Leu Gly Val
100       105       110
Thr Tyr Tyr Asp Gly Asn Glu Val Glu Gln Asp Tyr Lys Glu Ala Phe
115      120      125
Asp Trp Tyr Gly Lys Ala Ala Val Gln Gly Tyr Pro Asp Ala Glu Tyr
130      135      140
Asn Leu Ala Val Met Tyr Gly Leu Gly Lys Gly Thr Arg Gln Asp Phe
145      150      155      160
Ser Glu Thr Ile Lys Trp Leu Arg Lys Ala Ala Met His Gln Leu Pro
165      170      175
Glu Ala Gln Tyr Gly Leu Gly Val Met Tyr Ser Arg Gly Leu Gly Val
180      185      190

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-continued

Val Lys Asn Asp Glu Gln Ser Ala Tyr Trp Phe Ser Lys Ala Ala Arg
 195 200 205

Ala Gly Tyr Leu Lys Ala Gln Asn Lys Leu Gly Ile Leu Tyr Ser Glu
 210 215 220

Gly Lys Gly Leu Glu Lys Asp Glu Lys Lys Ala Phe His Trp Phe Glu
 225 230 235 240

Ala Ala Ala Glu Lys Gly Tyr Ala Lys Ala Gln Phe Asn Leu Ala Val
 245 250 255

Met Tyr Asp Lys Gly Ile Gly Val Ala Lys Asp Val Ser Lys Ala Ile
 260 265 270

Met Trp Tyr Arg Lys Ala Ala Thr Gln Gly Asn Val Ala Ala Gln Lys
 275 280 285

Arg Leu Lys Gln Leu His Ser
 290 295

<210> SEQ ID NO 43
 <211> LENGTH: 459
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 43

```

ttgaaagagt tcaatatacct gcaaggagcg attatgagca aaccagaggt tcaaaagacg      60
gatgccgaat ggagagcgca gctttcccct gtggcgatg ccgtgaccgc acaggcagcg      120
acggaaccgc cctttaccgg cgagtactgg aaccatgacg agaccggtgt ttatacgtgc      180
gtcaattgtg gaacgcccct gttcatttcc gatacgaat tcgacgccgg ttgtggtggtg      240
ccaagttttt ttgaccgat cgaccctgaa aatgtaaggg aaaaggtcga tgtctcgctg      300
ggcatggtgc gtaccgaaat tatctgtgcc atttgtgatg cccatctggg ccatgtgttt      360
gacgacggcc caccccaac ggggctgctg tattgtatta attctgccgc tttgcggttc      420
gaccgctac cgaagaaggc agaattctct gacaagtga      459

```

<210> SEQ ID NO 44
 <211> LENGTH: 152
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 44

Met Lys Glu Phe Asn Ile Leu Gln Gly Ala Ile Met Ser Lys Pro Arg
 1 5 10 15

Val Gln Lys Thr Asp Ala Glu Trp Arg Ala Gln Leu Ser Pro Val Ala
 20 25 30

Tyr Ala Val Thr Arg Gln Ala Ala Thr Glu Pro Pro Phe Thr Gly Glu
 35 40 45

Tyr Trp Asn His Asp Glu Thr Gly Val Tyr Thr Cys Val Asn Cys Gly
 50 55 60

Thr Pro Leu Phe Ile Ser Asp Thr Lys Phe Asp Ala Gly Cys Gly Trp
 65 70 75 80

Pro Ser Phe Phe Ala Pro Ile Asp Pro Glu Asn Val Arg Glu Lys Val
 85 90 95

Asp Val Ser Leu Gly Met Val Arg Thr Glu Ile Ile Cys Ala Ile Cys
 100 105 110

Asp Ala His Leu Gly His Val Phe Asp Asp Gly Pro Pro Pro Thr Gly

-continued

115	120	125
Glu Lys Ala Ala Asn Gln Asn Tyr Gly Pro Ala Gln Thr Leu Val Gly 130 135 140		
Val Met Tyr Tyr Lys Gly Met Gly Val Glu Gln Asn Phe Gly Thr Ala 145 150 155 160		
Lys Met Trp Leu Glu Lys Ala Ser Ala Gln Gly Glu Lys Asp Ala Gln 165 170 175		
Ser Phe Leu Gly Leu Met Tyr Leu Glu Gly Asp Asp Asn Asn Lys Asn 180 185 190		
Pro Lys Lys Ala Val Glu Leu Leu Thr Lys Ala Ala Asp Gln Asn Glu 195 200 205		
Pro Leu Ala Gln Thr Val Leu Gly Ile Met Tyr Ile Gln Gly Lys Phe 210 215 220		
Val Lys Gln Asp Tyr Lys Lys Ala Glu Glu Leu Leu Thr Lys Gly Ala 225 230 235 240		
Glu Ala Gly Asn Thr Asp Ala Ala Thr Phe Leu Gly Asn Met Tyr Tyr 245 250 255		
Arg Gly Gln Gly Val Asp Lys Asp Lys Ala Lys Ala Val Lys Trp Leu 260 265 270		
Glu Lys Ala Ala Ile Arg Gly Asp Val Asp Ala Gln Glu Leu Leu His 275 280 285		
Arg Ile His Tyr Gly Thr Pro Gly Ala Pro Lys 290 295		

<210> SEQ ID NO 47
 <211> LENGTH: 669
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 47

```

atggcaaaca aaacttttca gaaaaccctg ctgggttgct ccatgcacct gatgtccgct    60
ttcgtcgccg gcaatgccag cgcggccaaa aaagtgcagc acacctactc cagactgaac    120
cagtgtatcg gctactacaa ccagagccag tacaccagg caatcccctg ctttactcca    180
ctggcaaaa cgggtaacga tcaggcacag acctatctgg gcatcatgta ccagcacggt    240
ttcgttacc aaaagacat ggcaaccgct gccatgtggt ataacagggc cgcccgtcag    300
ggagacaaat gggcctacga caacctcaaa gccatgggta atccgaaact ggcaaccggc    360
aaaacctggg cggattacag aaacaccgtc gaacagaaag ccgctgcggg tgaagcctcg    420
gccagaccg ctctgggcag cctgtactac ttcggtgtcg gtggcgtcaa acaggactac    480
aacaccgcca aaaactggta tgccaaagcc gccgtcaatg gcgatgcagc cgccatgaac    540
tacctcggtc gcatgtatta ctacgcactg ggcgtogaac agaattccat gatggccaaa    600
caatactga acgcagccgc caaagccggt aacgttcagg caaaaaacct gctgaaaaaa    660
atcaataa                                         669
    
```

<210> SEQ ID NO 48
 <211> LENGTH: 222
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 48

Met Ala Asn Lys Thr Phe Gln Lys Thr Leu Leu Gly Cys Ser Ile Ala

-continued

1	5	10	15
Leu Met Ser Ala Phe Val Ala Gly Asn Ala Ser Ala Ala Lys Lys Val	20	25	30
Asp Asp Thr Tyr Ser Arg Leu Asn Gln Cys Ile Gly Tyr Tyr Asn Gln	35	40	45
Ser Gln Tyr Thr Gln Ala Ile Pro Cys Phe Thr Pro Leu Ala Lys Ala	50	55	60
Gly Asn Asp Gln Ala Gln Thr Tyr Leu Gly Ile Met Tyr Gln His Gly	65	70	80
Phe Gly Thr Gln Lys Asp Met Ala Thr Ala Ala Met Trp Tyr Asn Arg	85	90	95
Ala Ala Arg Gln Gly Asp Lys Trp Ala Tyr Asp Asn Leu Lys Ala Met	100	105	110
Gly Asn Pro Lys Leu Ala Thr Gly Lys Thr Trp Ala Asp Tyr Arg Asn	115	120	125
Thr Val Glu Gln Lys Ala Ala Ala Gly Glu Ala Ser Ala Gln Thr Ala	130	135	140
Leu Gly Ser Leu Tyr Tyr Phe Gly Val Gly Gly Val Lys Gln Asp Tyr	145	150	155
Asn Thr Ala Lys Asn Trp Tyr Ala Lys Ala Ala Val Asn Gly Asp Ala	165	170	175
Ala Ala Met Asn Tyr Leu Gly Arg Met Tyr Tyr Tyr Ala Leu Gly Val	180	185	190
Glu Gln Asn Ser Met Met Ala Lys Gln Tyr Leu Asn Ala Ala Ala Lys	195	200	205
Ala Gly Asn Val Gln Ala Lys Asn Leu Leu Lys Lys Ile Lys	210	215	220

<210> SEQ ID NO 49
 <211> LENGTH: 942
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes
 <400> SEQUENCE: 49

```

atgttgctga tggatattcg aatagaaaaa ctgaagataa aaaacgatct caaagataat      60
aatttcttga ccgaacttag caatgctgat gataatgaat cagcattgct tctttattat      120
gaaatgatga aaactatcctt aaaagcttta tttgtttgtc ttttttctt cctcggcaat      180
catgcaattg ccgataatct catggaagga aataggctat tcgatgctgg acaatataag      240
gaagccatga cctttcttat gcaaccgat gtacatgatg atccagaagc attaaatctt      300
gtcgcttata tgtataacca tggccttggc gtcagcaaaa atgctgaaaa agcatttatg      360
tgctatatga aatcggcaga atccggatta gctattgctc aatttaacgt gggactggcg      420
tatgaacagg gtaacggtat cctgaaaaat cttccagaag cagtcaagtg gtatagaaaa      480
gctgctgaac aagaagatgc tgatgctgaa gcaaaaatgg gttatttgac tgtaaacggc      540
ataggtatcg gaaagaatta taaagaagcc atgaaatggt atcaaagggc ggccgaacat      600
ggagattatg attcgtatgc cgatatcggg atgatgtatt ccaggggaga tgggtgcaaaa      660
aggaatttga accatgcagt tcagtattac atttttgggg ctcaaaaagg cagtacatat      720
tcacaggctt tattgggaaa tgcatatgca tacggaaaag gtatccaaaa ggacatagaa      780
caagcactct actggtacaa gcaggctgcc agaaaaggaa acgtcaatgc catgaaagaa      840
    
```

-continued

ctcggctaca tatacgagac cggctgactt ggtgtgaaaa aagacccaaa agaagcacia 900

tactggaaaag acatggctga aagagcgggc aaaaaacagt aa 942

<210> SEQ ID NO 50

<211> LENGTH: 313

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 50

Met Leu Leu Met Asp Ile Arg Ile Glu Lys Leu Lys Ile Lys Asn Asp
1 5 10 15Leu Lys Asp Asn Asn Phe Leu Thr Glu Leu Ser Asn Ala Asp Asp Asn
20 25 30Glu Ser Ala Leu Leu Leu Tyr Tyr Glu Met Met Lys Thr Ile Leu Lys
35 40 45Ala Leu Phe Val Cys Leu Phe Phe Phe Leu Gly Asn His Ala Ile Ala
50 55 60Asp Asn Leu Met Glu Gly Asn Arg Leu Phe Asp Ala Gly Gln Tyr Lys
65 70 75 80Glu Ala Met Thr Phe Leu Met Gln Pro Asp Val His Asp Asp Pro Glu
85 90 95Ala Leu Asn Leu Val Ala Tyr Met Tyr Asn His Gly Leu Gly Val Ser
100 105 110Lys Asn Ala Glu Lys Ala Phe Met Cys Tyr Met Lys Ser Ala Glu Ser
115 120 125Gly Leu Ala Ile Ala Gln Phe Asn Val Gly Leu Ala Tyr Glu Gln Gly
130 135 140Asn Gly Ile Leu Lys Asn Leu Pro Glu Ala Val Lys Trp Tyr Arg Lys
145 150 155 160Ala Ala Glu Gln Glu Asp Ala Asp Ala Glu Ala Lys Met Gly Tyr Leu
165 170 175Thr Val Asn Gly Ile Gly Ile Gly Lys Asn Tyr Lys Glu Ala Met Lys
180 185 190Trp Tyr Gln Arg Ala Ala Glu His Gly Asp Tyr Asp Ser Tyr Ala Asp
195 200 205Ile Gly Met Met Tyr Ser Arg Gly Asp Gly Val Lys Arg Asn Leu Asn
210 215 220His Ala Val Gln Tyr Tyr Ile Phe Gly Ala Gln Lys Gly Ser Thr Tyr
225 230 235 240Ser Gln Ala Leu Leu Gly Asn Ala Tyr Ala Tyr Gly Lys Gly Ile Gln
245 250 255Lys Asp Ile Glu Gln Ala Leu Tyr Trp Tyr Lys Gln Ala Ala Arg Asn
260 265 270Gly Asn Val Asn Ala Met Lys Glu Leu Gly Tyr Ile Tyr Glu Thr Gly
275 280 285Arg Leu Gly Val Lys Lys Asp Pro Lys Glu Ala Gln Tyr Trp Lys Asp
290 295 300Met Ala Glu Arg Ala Gly Lys Lys Gln
305 310

<210> SEQ ID NO 51

<211> LENGTH: 195

-continued

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 51

```

atgcctatgg aaaaggaatt tccaaagata ccaaagccc tctactggta caaaaccgca    60
gccaaaaacg gaaacgtcaa tgccatgaaa gaactggggtt ccatttatgc agaaggtgat    120
ctcggagtcc aaaaggacat acaagaagcg aaacgatgga acgacatggc cagaaaagcg    180
gaacaaaaga aataa                                           195

```

<210> SEQ ID NO 52

<211> LENGTH: 64

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 52

```

Met Pro Met Glu Lys Glu Phe Pro Lys Ile Pro Lys Ala Leu Tyr Trp
 1          5          10         15
Tyr Lys Thr Ala Ala Lys Asn Gly Asn Val Asn Ala Met Lys Glu Leu
 20         25         30
Gly Ser Ile Tyr Ala Glu Gly Asp Leu Gly Val Gln Lys Asp Ile Gln
 35         40         45
Glu Ala Lys Arg Trp Asn Asp Met Ala Arg Lys Ala Glu Gln Lys Lys
 50         55         60

```

<210> SEQ ID NO 53

<211> LENGTH: 948

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 53

```

atggttacgc ccgaatcgga taatgaaatc gacgacatca tctggtcaga aattgaaatcc    60
agtgattcgc acggtgattt cgtctgctat gccgttcacg ctccatataa ggcaaaatat    120
ctggagaatg ccaaagctcg tatcgaatcc ggaaaatc tggacaacct cgctcctgtc    180
cgctatctga aggctgttga acgtattgaa aaactggccg gaacgggcga cccggcggca    240
acattccaca tgggaaaaat atatgccatt ggcattgccg ttccgcaaaa tgtacccaaa    300
gccgttgaat ggtatgaaaa ggccatcgca ctcggagaac ccagagctta cgccaatctc    360
ggctggtttt atcaatcggg ttacgggtgt ccgactgaca aatcgaaagc tttcgaattg    420
ctgtcgttcg gtgctgaaaa cggagttttg tctgcaaagg ctgccatcgg catgatgctt    480
ttgaacgggg aaggatgtac cctgaatccg gaactcgggt ttcagaaact tgaggaatcg    540
ttcaacagcg gttatctgaa tgcgggcaac cacatttctg atgtgtatct tgaaggcaaa    600
ctggttccag gggatatoga gcttgcccat gaatggctac agaaagtggc ggatagcggc    660
gatgaaagat caatggcgat ccttgccac tatcttgtga caggaagtca tggaaagaca    720
gacacggcaa aagggttga tttgcttgag caggcaacct gactgcaata tctgcctgcc    780
tatttatggc tgggtacctt gtacaaaaaa gggcttggcg ttgagctgga tgcgaaaaa    840
gccattgaat ggttcaaaaa gggcatcaag gcggtctgtc gcgactgcca gatagcactt    900
acgatgatgt ccatgcctga aacggacaat ccgaagtcgt atcattaa                       948

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<210> SEQ ID NO 54

<211> LENGTH: 315

-continued

<212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes
 <400> SEQUENCE: 54

Met Val Thr Pro Glu Ser Asp Asn Glu Ile Asp Asp Ile Ile Trp Ser
 1 5 10 15
 Glu Ile Glu Ser Ser Asp Ser His Gly Asp Phe Val Cys Tyr Ala Val
 20 25 30
 His Ala Pro Tyr Lys Ala Lys Tyr Leu Glu Asn Ala Lys Ala Arg Ile
 35 40 45
 Glu Ser Gly Lys Tyr Leu Asp Asn Leu Ala Pro Val Arg Tyr Leu Lys
 50 55 60
 Ala Val Glu Arg Ile Glu Lys Leu Ala Gly Thr Gly Asp Pro Ala Ala
 65 70 75 80
 Thr Phe His Met Gly Lys Ile Tyr Ala Ile Gly Ile Ala Val Pro Gln
 85 90 95
 Asn Val Pro Lys Ala Val Glu Trp Tyr Glu Lys Ala Ile Ala Leu Gly
 100 105 110
 Glu Pro Arg Ala Tyr Ala Asn Leu Gly Trp Phe Tyr Gln Ser Gly Tyr
 115 120 125
 Gly Val Pro Thr Asp Lys Ser Lys Ala Phe Glu Leu Leu Ser Phe Gly
 130 135 140
 Ala Glu Asn Gly Val Leu Ser Ala Lys Ala Ala Ile Gly Met Met Leu
 145 150 155 160
 Leu Asn Gly Glu Gly Cys Thr Leu Asn Pro Glu Leu Gly Phe Gln Lys
 165 170 175
 Leu Glu Glu Ser Phe Asn Ser Gly Tyr Leu Asn Ala Gly Asn His Ile
 180 185 190
 Ser Asp Val Tyr Phe Glu Gly Lys Leu Val Pro Gly Asp Ile Glu Leu
 195 200 205
 Ala His Glu Trp Leu Gln Lys Val Ala Asp Ser Gly Asp Glu Arg Ser
 210 215 220
 Met Ala Ile Leu Gly His Tyr Leu Val Thr Gly Ser His Gly Lys Thr
 225 230 235 240
 Asp Thr Ala Lys Gly Leu Asp Leu Leu Glu Gln Ala Thr Arg Leu Gln
 245 250 255
 Tyr Leu Pro Ala Tyr Leu Trp Leu Gly Thr Leu Tyr Lys Lys Gly Leu
 260 265 270
 Gly Val Glu Leu Asp Ala Glu Lys Ala Ile Glu Trp Phe Lys Lys Gly
 275 280 285
 Ile Lys Ala Gly Cys Arg Asp Cys Gln Ile Ala Leu Thr Met Met Ser
 290 295 300
 Met Pro Glu Thr Asp Asn Pro Lys Ser Tyr His
 305 310 315

<210> SEQ ID NO 55
 <211> LENGTH: 912
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes
 <400> SEQUENCE: 55

atgaacaagc cttatccaaa cctgtcattt caaatgaaat cttttttcaa gatcttctgt 60
 atagtctgtc tgtttctttt cagtgttcct tcttttgccg acaacgcca ggaaggcatg 120

-continued

```

cgtctgttcc aggctggaaa atatcagcag gccatgacat atttcatgaa gccggatgcc 180
caaaagaatc cggatgtatt aaaccgcata gcttatatgt atgacaaagg ctttgggtgtt 240
gaaaagaatc tgcaacaacg tgcaagtgg tataaaaaag cagctgaaat gggattcaaa 300
gtagcacagt tcaatctcgg attaagttat caaaaaggct tgggagtcc aaaagatata 360
aatgaagcca tcaaatggta ccgtaaatca gcagaacagg gatatcccag cgccgaatcc 420
aaaatggggtt atttcaactgt caagggaaaa ggaataaac aggattttgc acaagccttg 480
aatgggtatc gtctcgtgc tgaacatggc gatgaccgtt cttatgcoga tatcggcatt 540
ttttatgccc aaggttatgg cgtcaaaaaa gacaggaacc gtgccgtgca gtattacatc 600
atgggtgctg aaaaaggcga cgcttatgcc cagtatcttt taggccgcgc ctatgagcag 660
gggcggggca ttcaatactc tcccgaacgt tcaactttact ggctcaaaaa agctgccgac 720
aatggaagtt tccttgccat gaagaactc gggatcgttt atgccaatgg acttctggac 780
cagaaaatgg ataccgatgc tgccgcaaaa tggggtgaaa aagcctggga aaccgcgaag 840
aagaacgggg aatccgatcc ggaggtcgac cgcagactgc gctttttcgg catagacccg 900
gatgacttat ag 912

```

<210> SEQ ID NO 56

<211> LENGTH: 303

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 56

```

Met Asn Lys Pro Tyr Pro Asn Leu Ser Phe Gln Met Lys Ser Phe Phe
 1          5          10          15
Lys Ile Phe Cys Ile Val Cys Leu Phe Leu Phe Ser Val Pro Ser Phe
 20          25          30
Ala Asp Asn Ala Lys Glu Gly Met Arg Leu Phe Gln Ala Gly Lys Tyr
 35          40          45
Gln Gln Ala Met Thr Tyr Phe Met Lys Pro Asp Ala Gln Lys Asn Pro
 50          55          60
Asp Val Leu Asn Arg Ile Ala Tyr Met Tyr Asp Lys Gly Phe Gly Val
 65          70          75          80
Glu Lys Asn Leu Gln Thr Ser Val Lys Trp Tyr Lys Lys Ala Ala Glu
 85          90          95
Met Gly Phe Lys Val Ala Gln Phe Asn Leu Gly Leu Ser Tyr Gln Lys
100          105          110
Gly Leu Gly Val Pro Lys Asp Ile Asn Glu Ala Ile Lys Trp Tyr Arg
115          120          125
Lys Ser Ala Glu Gln Gly Tyr Pro Ser Ala Glu Ser Lys Met Gly Tyr
130          135          140
Phe Thr Val Lys Gly Lys Gly Ile Lys Gln Asp Phe Ala Gln Ala Leu
145          150          155          160
Lys Trp Tyr Arg Leu Ala Ala Glu His Gly Asp Asp Arg Ser Tyr Ala
165          170          175
Asp Ile Gly Ile Phe Tyr Ala Glu Gly Tyr Gly Val Lys Lys Asp Arg
180          185          190
Asn Arg Ala Val Gln Tyr Tyr Ile Met Gly Ala Glu Lys Gly Asp Ala
195          200          205

```

-continued

Tyr Ala Gln Tyr Leu Leu Gly Arg Ala Tyr Glu Gln Gly Arg Gly Ile
 210 215 220

Gln Tyr Ser Pro Glu Arg Ser Leu Tyr Trp Leu Lys Lys Ala Ala Asp
 225 230 235 240

Asn Gly Ser Phe Leu Ala Met Lys Glu Leu Gly Ile Val Tyr Ala Asn
 245 250 255

Gly Leu Leu Asp Gln Lys Met Asp Thr Asp Ala Ala Ala Lys Trp Gly
 260 265 270

Glu Lys Ala Trp Glu Thr Arg Lys Lys Asn Gly Glu Ser Asp Pro Glu
 275 280 285

Val Asp Arg Arg Leu Arg Phe Phe Gly Ile Asp Pro Asp Asp Leu
 290 295 300

<210> SEQ ID NO 57
 <211> LENGTH: 936
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 57

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atggcactcc cttcttccc gtcacccca ttcacaaagt ggctctccc tctggcgctg    60
gtctgtttca tcctgccgc gatgccgatc gaagccgttg caaaagacc gagcgaaggt    120
gtgaagcttt actataacca gcaatacaaa aaggccgccc ctttctgat gaaagaagcc    180
aaaaaaggca atgccaaggc acaggctctgc ctcgggatga tgtatcagga gggctcttggc    240
ctgaaacaga actacatgct ggccagacgc tggatgaaa aatcggcgaa gaaaaaccgg    300
gctgacgcac aaactttcct cgccatgctc tacagtcagg ggcttgggtg ggcgaaagat    360
ttcgaaaaag ccaaatattg gttcgacaag gccgcccggc agggctttgc ccctgccag    420
acactcgtcg gctgatgta cgccaaaggc gtcggcaccg caaaaagcat gtcgcaagcg    480
gaaaaatggc tgcggctggc cgccaaacag ggtgaaccgg acgcccagac ctatctgggc    540
cttttgatcc ttgacggcac tgaactgccg caggacgtag gtgaagccgc gcgacttttg    600
aaagaagcgg ccgtaaaagg cgaccgaac gccagctctg cactgggcat gatgtatttt    660
tccggcaagg gggctgatca ggacatgaac gagtcggaaa aatggcttga aaaggcggcc    720
attgccgata acgtcgatgc acagactttc ctcggcaacc tctattacaa gggatcggc    780
gtcgcgaaag acgatacccg cgcacgctac tggcttcaga aagcggcgat tgcgggtgat    840
ccggacgcac aggccacatt aatgatatg ttgaaagacg atgcagcggc cgttttgccc    900
ttcgaggaca agaagcccct ttcgggttcc ctgtaa                                936

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<210> SEQ ID NO 58
 <211> LENGTH: 311
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 58

Met Ala Leu Pro Ser Phe Pro Ser Ser Pro Phe Ile Lys Trp Leu Ser
 1 5 10 15

Arg Leu Ala Leu Val Cys Phe Ile Leu Pro Ala Met Pro Ile Glu Ala
 20 25 30

Val Ala Lys Asp Arg Ser Glu Gly Val Lys Leu Tyr Tyr Asn Gln Gln
 35 40 45

Tyr Lys Lys Ala Ala Pro Phe Leu Met Lys Glu Ala Lys Lys Gly Asn

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50		55				60									
Ala	Lys	Ala	Gln	Val	Cys	Leu	Gly	Met	Met	Tyr	Gln	Glu	Gly	Leu	Gly
65					70					75					80
Leu	Lys	Gln	Asn	Tyr	Met	Leu	Ala	Arg	Arg	Trp	Tyr	Glu	Lys	Ser	Ala
				85					90						95
Lys	Lys	Asn	Arg	Ala	Asp	Ala	Gln	Thr	Phe	Leu	Gly	Met	Leu	Tyr	Ser
			100					105					110		
Gln	Gly	Leu	Gly	Val	Ala	Lys	Asp	Phe	Glu	Lys	Ala	Lys	Tyr	Trp	Phe
		115					120					125			
Asp	Lys	Ala	Ala	Gly	Gln	Gly	Phe	Ala	Pro	Ala	Gln	Thr	Leu	Val	Gly
	130					135					140				
Leu	Met	Tyr	Ala	Lys	Gly	Val	Gly	Thr	Ala	Lys	Ser	Met	Ser	Gln	Ala
145					150					155					160
Glu	Lys	Trp	Leu	Arg	Leu	Ala	Ala	Lys	Gln	Gly	Glu	Pro	Asp	Ala	Gln
			165						170						175
Thr	Tyr	Leu	Gly	Leu	Leu	Tyr	Leu	Asp	Gly	Thr	Glu	Leu	Pro	Gln	Asp
		180						185						190	
Val	Gly	Glu	Ala	Ala	Arg	Leu	Leu	Lys	Glu	Ala	Ala	Val	Lys	Gly	Asp
		195					200						205		
Pro	Asn	Ala	Gln	Ser	Ala	Leu	Gly	Met	Met	Tyr	Phe	Ser	Gly	Lys	Gly
	210					215					220				
Val	Asp	Gln	Asp	Met	Asn	Glu	Ser	Glu	Lys	Trp	Leu	Glu	Lys	Ala	Ala
225					230					235					240
Ile	Ala	Gly	Asn	Val	Asp	Ala	Gln	Thr	Phe	Leu	Gly	Asn	Leu	Tyr	Tyr
			245						250						255
Lys	Gly	Ile	Gly	Val	Ala	Lys	Asp	Asp	Thr	Arg	Ala	Arg	Tyr	Trp	Leu
		260						265						270	
Gln	Lys	Ala	Ala	Ile	Ala	Gly	Asp	Pro	Asp	Ala	Gln	Ala	Thr	Leu	Asn
		275					280						285		
Asp	Met	Leu	Lys	Asp	Asp	Ala	Ala	Pro	Val	Leu	Pro	Phe	Glu	Asp	Lys
	290					295					300				
Lys	Pro	Leu	Ser	Gly	Ser	Leu									
305					310										

<210> SEQ ID NO 59
 <211> LENGTH: 660
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 59

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atgtccaggg gttccaaaa caaaaagctt tttttatggc tcattctcgc cgccacgac 60
atcctgatca tcatcagcca gactgctttc gcagtcggtg agcgcgacct tgaacaggcg 120
atgcagtcct attccgcggg acaatacaaa gaggctgttc cccaattcga aaaagcggca 180
acgctgggca acaacaaggc acaaacctatg ctgggtgtcc tgtattttca gggcaaggga 240
tgcgaaaagg actacgtgaa ggcgcggcaa tggttcgacc gtgcggccaa tggcggcaat 300
atcgaagcgc aaaccttcat gggcatcadc aatcttgaag gcctcggtac tcccaaaaac 360
gaaaaaacgg cctattactg gttcgaaaaa gccgcccgtg gcggtgaaac cagcgcacag 420
aattatctcg gcaccctgct catgaatgga cagggcacga aaagagactc tgcaaaagcg 480
gctgaatggt ttaccaaagc cgccgaaaag ggtgacctga acgcccggaa aatccttggc 540
    
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 gcgatgtact ttcagggcac aggcgttgcc aaagacatgg taaaagcccg gtactggctg 600

caaaaagcgg ctgatgacgg tgacatggac gcaaaaaaca tgctctccga actcaataa 660

<210> SEQ ID NO 60

<211> LENGTH: 219

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 60

Met Ser Arg Gly Ser Lys Asn Lys Lys Leu Phe Leu Trp Leu Ile Leu
1 5 10 15Ala Ala Thr Ile Ile Leu Ile Ile Ile Ser Gln Thr Ala Phe Ala Val
20 25 30Gly Glu Arg Asp Leu Glu Gln Ala Met Gln Ser Tyr Ser Ala Gly Gln
35 40 45Tyr Lys Glu Ala Val Pro Gln Phe Glu Lys Ala Ala Thr Leu Gly Asn
50 55 60Asn Lys Ala Gln Thr Met Leu Gly Val Leu Tyr Phe Gln Gly Lys Gly
65 70 75 80Cys Glu Lys Asp Tyr Val Lys Ala Ala Glu Trp Phe Asp Arg Ala Ala
85 90 95Asn Gly Gly Asn Ile Glu Ala Gln Thr Phe Met Gly Ile Ile Asn Leu
100 105 110Glu Gly Leu Gly Thr Pro Lys Asn Glu Lys Thr Ala Tyr Tyr Trp Phe
115 120 125Glu Lys Ala Ala Arg Gly Gly Glu Thr Ser Ala Gln Asn Tyr Leu Gly
130 135 140Thr Leu Leu Met Asn Gly Gln Gly Thr Lys Arg Asp Ser Ala Lys Ala
145 150 155 160Ala Glu Trp Phe Thr Lys Ala Ala Glu Lys Gly Asp Leu Asn Ala Arg
165 170 175Lys Ile Leu Gly Ala Met Tyr Phe Gln Gly Thr Gly Val Ala Lys Asp
180 185 190Met Val Lys Ala Arg Tyr Trp Leu Gln Lys Ala Ala Asp Asp Gly Asp
195 200 205Met Asp Ala Lys Asn Met Leu Ser Glu Leu Lys
210 215

<210> SEQ ID NO 61

<211> LENGTH: 3546

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 61

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atcgtcgcgg cggttttcat tcccttttca tegacaacag cagggacagc aaaccgccc 120

accattcaga aaactccoga acttgcccgt caggtcaatg aaacgcttaa cttcgatcac 180

aaaacagaag acatcctgaa aacactgacg gaatcaaaaa acccgtccga taaaaccctg 240

atgggagcgt tttacggttc cggtgcaggt ggaaaacagg actggggaaa agcccggatg 300

tggtttgaaa aagcggccag tgaaggcgat gcgcacgctg aatattttct gggctcttct 360

tacatggggg gtctgggcac ccccaaagat tatgacaagg cgtttcattg gctattgtt 420

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gccgcccgta	aggacattcc	cgacgcacag	taccagttga	gttggttcta	tgccaacgga	480
aaaggtacga	gccaaagtct	gcgtgaaacg	gtttactgga	tacaaaaggc	agcccacaag	540
gggcatgtag	tagccatgcg	aagcatggga	atgctcagtt	actcgggatt	aggcatgccc	600
gaaaacaagg	tcgacgcttt	caaatggttt	gaaaaagcgg	cgagtgcagg	cgatgccgaa	660
gcccaatatac	accttggecat	gtcctatatg	gcaggcaagg	gaacggaaaa	agacggaaaa	720
aagggagaag	aatggcttta	tcgtgccgcg	ctgcaaaatc	agacgaaagc	ccaggattat	780
ctttccgtcc	tttatgtgca	gcgattgctt	gataaaaaaa	acaggccggg	cgaaatcgaa	840
caggcaagac	agtggctgga	aaacgcccgc	cgacgaaacg	ataaaaatgc	gatatacatg	900
ctgaacgtca	ttgagcagta	ttctcgcaaa	acatcgaaaa	atcaggccga	aaccctcgac	960
acccttcgtc	gtcgcgcoga	acagggcaat	gccgattccc	aattcatgct	gggagaggcc	1020
ctgcttgccg	gaacgggaat	gaagaaaaat	ccggaagaag	ctgtccgctg	gtttgaaaaa	1080
gcggcataac	aggccaacat	tgatgcacag	tccgcccctg	gatatatgca	ttatttcggc	1140
gtccatgtcc	ctgtcgatta	cgcaaaagcc	attcccctgt	taaaacaggg	tgccgataag	1200
gggaacagcc	aggcacaac	cgctatggga	ttcgcctatg	ccagtggaac	aggaattgcc	1260
aaaaacgagc	aaaaagcatt	tgaactgttt	gaaaaagcgg	cccggaaaca	tgtccgaagt	1320
gcccagtttt	acctcgggtga	aatgctttaa	aatggcattg	gaaccaacg	aaacgtccca	1380
gagggattgg	catggatcga	aaaatcggcg	aaggcaggat	acgaccaggc	ccagtttacg	1440
atgggaataa	acgctctccg	tggcaaggac	aagatgcaaa	atattgacga	agcgcgcaaa	1500
tggatgcgtc	tggccgcca	acagaatcat	gcagaagccc	agtacatgct	cggcgatgca	1560
tattttctgg	gagaactgac	accggaaaat	caaaaggaag	gtattttctg	gtgggacaag	1620
gccgcccgcc	aaaatcaogt	cgaagcccgg	ctcatgctgt	tcaagtatca	ctgcgatccc	1680
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cgtgaccctg	acgcgcttta	ttcccttgg	gaattatttt	ttttcgtaa	tgacaaccat	1800
aaaaaaaaacg	tcccgaagc	tgtcgatttt	ttcagcaagg	cggtgatct	gaaccataca	1860
gaaagccagt	acatgctggg	gctgattctc	tattccaaaa	cggatgtcgg	acaaaacaaa	1920
aaacaagcct	gccagtgggt	tgaaaaagcg	gcaagtcaca	atcaccggga	aagccaatat	1980
atgctcggaa	tctgtgttct	cgaggggaac	cacacttccg	ccgacaaaaa	aaaggcgctt	2040
gaactgatca	gactggcagc	cgacaaaaac	gtcagcatcg	cccagaacaa	aatgggggat	2100
ctgtatgaaa	caggccatat	cgttccgaaa	gacatgaaaa	aagccatcga	atggtacaca	2160
cttgccgaac	agaatggatt	tacggatgcg	gcttaccatc	tggcccctgt	gtaccttgcc	2220
agttccccac	ccctgcaaaa	cgatcccgtg	gctctgcgtt	atcttgaaaa	agcggctagc	2280
gcaacaata	caaatgtctc	ttataaactg	ggaacgtttt	atctocacgg	tcaatactcg	2340
gccacaaaag	atcggaaaaa	agcggcggag	tatttccgcc	gtgcgcgcaa	actcggccat	2400
aaaaacagcc	aaatagccta	tgccgacatc	ctccaaaaag	gtaagggagt	tgaaaaaac	2460
gaaaaactgg	cttgcaaat	ctatgaaaaa	acagccaaag	aaggtagccc	ttacggacaa	2520
ttccgctcag	gactgtgcta	tcagacaggt	cttggaata	gacccaaaaa	tcctgcaaaa	2580
gcggtcagtc	tttttgaa	ggcagccaga	caaacctgc	cggatggcca	gattgcccctg	2640
gcatactgtt	atgaaaccgg	acagggcggt	gcccaaaatc	ttgcgctggc	gttcaaatgg	2700

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tacaaaatgg cagctgaaaa gggagatgtc ggcagcatga tcacaaccgg aaagatgctg 2760
gacaagggag aaggaacagc acgcgacagc aaacaggctt tttactggtt ttcaaaagcg 2820
gctgagaaaag gatcaccoga agcagaagtt cagttgggcc aactgtatta tgccggacgt 2880
ggcatttccg ctgacatgaa aaaagcggtt tccctcttcg accattcggc cagacaaggc 2940
aacgctttgg ctcaactactg gatgggttat ctctgtcttc acggaaaagg ggtcgaaaaa 3000
aatgagccac tggtctgta ctggctcgaa aaagccgccc tccagaatca gacaggcgcc 3060
gcttttgaac tggcaaaaaca gtattggaat ggcaacggca tcccgtccga tccggaacag 3120
gctattgtct ggttcaccaa agccgcacaa aataacgatg ttcaggcaca aagagcactg 3180
gcggttcattt atagcgtcca tggcgcgaaa aaagggataa agccggatga ccaaaaagcc 3240
ttttactggg caaaciaaagc cgccagatat aacgatgatc cctcgaccag actggtgctg 3300
ggaacgtttt atatctcggg aaaagggacc gctgtagatg aaaaaaagg ccttctcctt 3360
ctgaaagaag ccgagagaaa aaactatccc caggcaatgg cgatgctggg tgagttctat 3420
cttgagaaaa aagacaggaa agaagcacia atgtggttca agcgtgctgc cgcctccggt 3480
gatcagaagg tcatcgaat aatgaagaaa aaagcgcttt atccgcaaaa ccccgccat 3540
ccctga

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<210> SEQ ID NO 62

<211> LENGTH: 1181

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 62

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Met Asn Ser Phe Gln Trp His Ile Val Lys Leu Lys His Arg Ala Leu
1          5          10          15
Leu Ser Ser Leu Ile Val Ala Ala Ala Phe Ile Pro Phe Ser Ser Thr
20        25        30
Thr Ala Gly Thr Ala Asn Pro Pro Thr Ile Gln Lys Thr Pro Glu Leu
35        40        45
Ala Arg Gln Val Asn Glu Thr Leu Asn Phe Asp His Lys Thr Glu Asp
50        55        60
Ile Leu Lys Thr Leu Thr Glu Ser Lys Asn Pro Ser Asp Lys Thr Leu
65        70        75        80
Met Gly Ala Phe Tyr Gly Ser Gly Ala Gly Gly Lys Gln Asp Trp Gly
85        90        95
Lys Ala Arg Met Trp Phe Glu Lys Ala Ala Ser Glu Gly Asp Ala His
100       105       110
Ala Glu Tyr Phe Leu Gly Leu Leu Tyr Met Gly Gly Leu Gly Thr Pro
115      120      125
Lys Asp Tyr Asp Lys Ala Phe His Trp Leu Leu Leu Ala Ala Arg Lys
130      135      140
Asp Ile Pro Asp Ala Gln Tyr Gln Leu Ser Trp Phe Tyr Ala Asn Gly
145      150      155      160
Lys Gly Thr Ser Gln Ser Leu Arg Glu Thr Val Tyr Trp Ile Gln Lys
165      170      175
Ala Ala His Lys Gly His Val Val Ala Met Arg Ser Met Gly Met Leu
180      185      190
Ser Tyr Ser Gly Leu Gly Met Pro Glu Asn Lys Val Asp Ala Phe Lys
195      200      205

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Trp Phe Glu Lys Ala Ala Ser Ala Gly Asp Ala Glu Ala Gln Tyr His
 210 215 220
 Leu Gly Met Ser Tyr Met Ala Gly Lys Gly Thr Glu Lys Asp Gly Lys
 225 230 235 240
 Lys Gly Glu Glu Trp Leu Tyr Arg Ala Ala Leu Gln Asn Gln Thr Lys
 245 250 255
 Ala Gln Asp Tyr Leu Ser Val Leu Tyr Val Gln Arg Leu Leu Asp Lys
 260 265 270
 Lys Asn Arg Pro Gly Glu Ile Glu Gln Ala Arg Gln Trp Leu Glu Asn
 275 280 285
 Ala Ala Arg Arg Asn Asp Lys Asn Ala Ile Tyr Met Leu Asn Val Ile
 290 295 300
 Glu Gln Tyr Ser Arg Lys Thr Ser Lys Asn Gln Ala Glu Thr Leu Asp
 305 310 315 320
 Thr Leu Arg Arg Arg Ala Glu Gln Gly Asn Ala Asp Ser Gln Phe Met
 325 330 335
 Leu Gly Glu Ala Leu Leu Ala Gly Thr Gly Met Lys Lys Asn Pro Glu
 340 345 350
 Glu Ala Val Arg Trp Phe Glu Lys Ala Ala Lys Gln Gly Asn Ile Asp
 355 360 365
 Ala Gln Ser Ala Leu Gly Tyr Met His Tyr Phe Gly Val His Val Pro
 370 375 380
 Val Asp Tyr Ala Lys Ala Ile Pro Leu Leu Lys Gln Gly Ala Asp Lys
 385 390 395 400
 Gly Asn Ser Gln Ala Gln Thr Ala Met Gly Phe Ala Tyr Ala Ser Gly
 405 410 415
 Thr Gly Ile Ala Lys Asn Glu Gln Lys Ala Phe Glu Leu Phe Glu Lys
 420 425 430
 Ala Ala Arg Asn Asn Val Arg Ser Ala Gln Phe Tyr Leu Gly Glu Met
 435 440 445
 Leu Glu Asn Gly Ile Gly Thr Gln Arg Asn Val Pro Glu Gly Leu Ala
 450 455 460
 Trp Ile Glu Lys Ser Ala Lys Ala Gly Tyr Asp Gln Ala Gln Phe Thr
 465 470 475 480
 Met Gly Ile Asn Ala Leu Arg Gly Lys Asp Lys Met Gln Asn Ile Asp
 485 490 495
 Glu Ala Arg Lys Trp Met Arg Leu Ala Ala Lys Gln Asn His Ala Glu
 500 505 510
 Ala Gln Tyr Met Leu Gly Met Ser Tyr Phe Leu Gly Glu Leu Thr Pro
 515 520 525
 Glu Asn Gln Lys Glu Gly Ile Phe Trp Trp Asp Lys Ala Ala Ala Gln
 530 535 540
 Asn His Val Glu Ala Arg Leu Met Leu Phe Lys Tyr His Cys Asp Pro
 545 550 555 560
 Lys Met His Tyr Ala Asp Arg Lys Arg Cys Ser Ser Ile Ser Glu Gln
 565 570 575
 Met His Asp Ile Arg Asp Pro Asp Ala Leu Tyr Ser Leu Gly Glu Leu
 580 585 590
 Phe Phe Phe Gly Asn Asp Asn His Lys Lys Asn Val Pro Lys Ala Val
 595 600 605

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Asp Phe Phe Ser Lys Ala Ala Asp Leu Asn His Thr Glu Ser Gln Tyr
 610 615 620
 Met Leu Gly Leu Ile Leu Tyr Ser Lys Thr Asp Val Gly Gln Asn Lys
 625 630 635 640
 Lys Gln Ala Cys Gln Trp Phe Glu Lys Ala Ala Ser His Asn His Pro
 645 650 655
 Glu Ser Gln Tyr Met Leu Gly Ile Cys Val Leu Glu Gly Asn His Thr
 660 665 670
 Ser Ala Asp Lys Lys Lys Ala Leu Glu Leu Ile Arg Leu Ala Ala Asp
 675 680 685
 Lys Asn Val Ser Ile Ala Gln Asn Lys Met Gly Tyr Leu Tyr Glu Thr
 690 695 700
 Gly His Ile Val Pro Lys Asp Met Lys Lys Ala Ile Glu Trp Tyr Thr
 705 710 715 720
 Leu Ala Glu Gln Asn Gly Phe Thr Asp Ala Ala Tyr His Leu Ala Leu
 725 730 735
 Leu Tyr Leu Ala Ser Ser Pro Pro Leu Gln Asn Asp Pro Leu Ala Leu
 740 745 750
 Arg Tyr Leu Glu Lys Ala Ala Ser Ala Asn Asn Thr Asn Ala Leu Tyr
 755 760 765
 Lys Leu Gly Thr Phe Tyr Phe His Gly Gln Tyr Ser Ala Thr Lys Asp
 770 775 780
 Arg Lys Lys Ala Ala Glu Tyr Phe Arg Arg Ala Ala Lys Leu Gly His
 785 790 795 800
 Lys Asn Ser Gln Ile Ala Tyr Ala Asp Ile Leu Gln Lys Gly Lys Gly
 805 810 815
 Val Glu Lys Asn Glu Lys Leu Ala Cys Glu Ile Tyr Glu Lys Thr Ala
 820 825 830
 Lys Glu Gly Ser Pro Tyr Gly Gln Phe Arg Ser Gly Leu Cys Tyr Gln
 835 840 845
 Thr Gly Leu Gly Asn Arg Pro Lys Asn Pro Ala Lys Ala Val Ser Leu
 850 855 860
 Phe Glu Gln Ala Ala Arg Gln Asn Leu Pro Asp Gly Gln Ile Ala Leu
 865 870 875 880
 Ala Tyr Cys Tyr Glu Thr Gly Gln Gly Val Ala Gln Asn Leu Ala Leu
 885 890 895
 Ala Phe Lys Trp Tyr Lys Met Ala Ala Glu Lys Gly Asp Val Gly Ser
 900 905 910
 Met Ile Thr Thr Gly Lys Met Leu Asp Lys Gly Glu Gly Thr Ala Arg
 915 920 925
 Asp Ser Lys Gln Ala Phe Tyr Trp Phe Ser Lys Ala Ala Glu Lys Gly
 930 935 940
 Ser Pro Glu Ala Glu Val Gln Leu Gly Gln Leu Tyr Tyr Ala Gly Arg
 945 950 955 960
 Gly Ile Ser Ala Asp Met Lys Lys Ala Val Ser Leu Phe Asp His Ser
 965 970 975
 Ala Arg Gln Gly Asn Ala Leu Ala Gln Tyr Trp Met Gly Tyr Leu Cys
 980 985 990
 Leu His Gly Lys Gly Val Glu Lys Asn Glu Pro Leu Ala Arg Asp Trp
 995 1000 1005
 Leu Glu Lys Ala Ala Val Gln Asn Gln Thr Gly Ala Ala Phe Glu

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1010	1015	1020
Leu Ala Lys Gln Tyr Trp Asn Gly Asn Gly Ile Pro Ser Asp Pro		
1025	1030	1035
Glu Gln Ala Ile Val Trp Phe Thr Lys Ala Ala Gln Asn Asn Asp		
1040	1045	1050
Val Gln Ala Gln Arg Ala Leu Ala Phe Ile Tyr Ser Val His Gly		
1055	1060	1065
Ala Lys Lys Gly Ile Lys Pro Asp Asp Gln Lys Ala Phe Tyr Trp		
1070	1075	1080
Ala Asn Lys Ala Ala Arg Tyr Asn Asp Asp Pro Ser Thr Arg Leu		
1085	1090	1095
Leu Leu Gly Thr Phe Tyr Ile Ser Gly Lys Gly Thr Ala Val Asp		
1100	1105	1110
Glu Lys Lys Gly Leu Leu Leu Leu Lys Glu Ala Ala Glu Lys Asn		
1115	1120	1125
Tyr Pro Gln Ala Met Ala Met Leu Gly Glu Phe Tyr Leu Glu Lys		
1130	1135	1140
Lys Asp Arg Lys Glu Ala Gln Met Trp Phe Lys Arg Ala Ala Ala		
1145	1150	1155
Ser Gly Asp Gln Lys Val Ile Glu Ile Met Lys Lys Lys Gly Val		
1160	1165	1170
Tyr Pro Gln Asn Pro Val His Pro		
1175	1180	

<210> SEQ ID NO 63

<211> LENGTH: 3852

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 63

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atgaaccctt tgccatcaga cagccatttt ttcaaacggt ttgtacgggg atatatoctg    60
ccttgccctt tcttccccat cagcgtagcc acagcctctc cggctcagca ccttgtttat    120
cgggagatca tggaaatgcc acaggagata gaccaaacc tgctggtcga tcatgacagt    180
gaaaaaatac tgaacaaact ggcggactca aacgcccccg ccatgaaaac actcatggga    240
gcctggtagc ctatcggagc cggaggcaaa agggactgga tcaaagcccc catctggttt    300
gaaaaagccg cactgaagg cgacacaaga gccgcatac ccctcggtct tttatactcg    360
gcccgtctgg gcacccccat cgattacgac aaggcctttt actggctgtc aatcgcggca    420
cgccagaata ttccggatgc ccagtaccgg ctggcaggac tttatcagga aggaaaagga    480
accgccaaaa gcgaacgtga attcgcttac tgggtaaaaa aagccgctgg aaacggacac    540
attgacgccc agagagccat gggaatgatc cttcattatg gtctcggagt tcacaaaaac    600
cttctgaat cgggtcaaatg gtttgaaaaa gcagccaatg ccgaaaacgc aaccgcacaa    660
tattatctcg gcattgatta tatgaatggt aatggcctcg ccaaaaatga aagggaagga    720
gaaaagtggc tttatcgcgc cgccatgcaa gaccatcttg aagctcaaac atatctcggc    780
accatttata tgaaacgcaa gcttcaaaat gaaggtcagc caccggaaac cgcccttgcc    840
atccagtggg tggaaaatgc cgctacgcca aatgacccgt acgctatccg gcttctatcc    900
atggtgtaca ggcatttcc cgaactgcaa aacaacgcca aaggcatgct tcatcttcgc    960
cgcagtgcgg agatggggaa tgcacagcc caattcgatc tcggaagaac cctgtttcag   1020

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gataaaaaact cacctgcaaa acgcaaggaa gctgtcgtct ggcttgaaaa agcgcgcaaa 1080
caggacgaac gtagggcaca ggcttttctt ggtaacatgt attactatgg cgagtctgtt 1140
cctgtcgatt atgtcaaggc gctccctctt ttgatgaggg ctgccgacaa gggagacagc 1200
tttgacaat acacgcttg tttggcttat atcgatggga atggcattgc caaagatgaa 1260
cgaaaggctt tttcatgggt gaaaaaatca gccagtcaaa acagggcaag tgcccagtat 1320
tttttaggac ttatgtacct tgatggaaca gggacacctg tcaatgagga aaaaggcatc 1380
cgtttctca aagaattagc aaagacaggg tatgtctatg cccaataata gcttgggtacc 1440
tacgcccata gcgggcttca tatggccaaa gatctggcag aagcagagaaa atggtaccag 1500
ttggcggcca gtcaggatca tatcaaggcg aaatactggc tcggcatgtc cctctttcag 1560
ggcccggaca gtgaacagga caggaaaaaa ggcgtttact ggttcacoga agcagcaaaa 1620
caggatgacc cggatgcgca gcttgaactg gaaaaatccc tgctctacgg cgatggtatc 1680
gacaaaaacg aaaagcaggc gtcacagtgg ttcaaaaaag cgcacaataa ccaacagcac 1740
accggccagt attacgcagg aatgtgcctt atgcggggca tcaatgggccc ggtcgatatt 1800
ccgaaaggaa tgtcactcat tgaatgtct cgaataata aggtctcaat ggcccagttc 1860
cagcttggga aactgtatga atatggtttg aaagagttgc cgaaagacat ctgaaagcc 1920
attggctggt acacacgtgc cgcagaaaat ggacatgcaa ctgcccaata cgggctggga 1980
aagctttatc tcaaagctga tactccactc aaaaatattc cgctcgccct cgaattcctt 2040
gaaaaatccg cttcacaanaa tataaccagt gccatttttg atctgggtaa catttactat 2100
gacggaaaaa ttgtcaaaaca ggatatggca aaagcgttga attactttca gaagggaaacc 2160
ggactgggcc atctcccag ccagaathtt gtcggattca tgatcgagaa tggaagcggga 2220
gttaaaaagg ataagaaaa agcatgcaaa atctatgatg aaaccggtcg acgcggaagt 2280
gcataatggac tttaccgcta tggcttgtgc cagctttccg atcccgatcc atcccggaa 2340
aaccagaaaa aagcgtttat tcttttoga caggctgccc gaaaaatctt ggcagacgcc 2400
caatactttt tagccctgtg ttacgaatat ggcaaaagga ctcccaaaaa ccctggcgaa 2460
gctattgaat ggtacaggcg tgctcggaa aacgacaaac cgggaagcact gtaccagttg 2520
ggtatgcttt atatcaccag cccgtctcct cgccaaaaa taccgctggg tctggattat 2580
ctggagaaa gacccgccc ggatctgtcg tccgcattca acgaactggg ccgaatttat 2640
tatgacggga aaatcgtcag gcaggatctc aaaaaatccg tcttctggta tcgcaagggga 2700
gcacaatcgg gagatacaag aagccagaat gatctggctt atatgatgga atacggtaaa 2760
ggtcttga aagacgaaaa ggcggcctgc acgatgtatg aaaaaacggc aggggaaaaa 2820
aacgcttatg gccagttccg tctggggctc tgctatctga atggaaaagg cggtaaagcg 2880
aaggatcaac gggaaagcagt ccggctcttc gaatctgccg ccggacaaaa tctggcgtct 2940
gcccataatt ttctgggaat ttaccataaa gaaggtaaag gtgtcgtcaa aaacatgaat 3000
gaggctttca aatggtacct gacggcagcc gataacggtc acgtcagttc catgtttgaa 3060
gtcggtaaaa tgttcgctaa tggcagaggt acggaacgcg acgacaaaaa ggcatttcac 3120
tggtttgaga aagcagctga aaacgggtct gattccgctt tgaccagct gggcattatg 3180
tattataaag gcttgggcat ttcagccgac aaaagcaaa cgcctcctt tttttgaaa 3240
gctgccgaaa aaaacaattc atacgcccac cactggctcg gctatatgta cctgtaacgtt 3300

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aaggggctgg aaaaaaacgg ggagctggca aatcagtggt tatcaaaagc ggctgaccag 3360
aatgaaacgg gcgcgatttt tgaactggga aaacaatact ggtatggcat gggcgtcccg 3420
gtcaatcctg aaaaggcaat cgctcctgctt caaaaagcag ggaacgatgg ccatgttatt 3480
gcacaaagga tactgggcta tatttatgct gatggcggcc cggaaaaagg gattcccctt 3540
gatttcgaaa aggccgtcca atggtttgaa aaagcggccc gacaagacga cgccgcccgg 3600
aaaatggggg tggcattgct cactctgacc ggaaaaggaa ctccaaaaaa tggaggagaaa 3660
ggtatccggc ttttgacgca gtctgcaaac atgaattatc catccgcgat ggaactgctc 3720
ggggattttt atcgtgaaga aaaagtgat aaaaaagaag cggaaaaatg gtatcgtcgt 3780
gccgccgaaa ccgcgacaaa aaccgttatc gaatccatga agaaaaaagg cgtttatccg 3840
gcaaaccctg aa 3852

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<210> SEQ ID NO 64

<211> LENGTH: 1283

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 64

```

Met Asn Pro Leu Pro Ser Asp Ser His Phe Phe Lys Arg Phe Val Arg
1           5           10           15
Gly Tyr Ile Leu Pro Cys Leu Phe Phe Pro Ile Thr Val Ala Thr Ala
20           25           30
Ser Pro Ala Gln His Leu Val Tyr Pro Glu Ile Met Glu Met Pro Gln
35           40           45
Glu Ile Asp Gln Thr Leu Leu Val Asp His Asp Ser Glu Lys Ile Leu
50           55           60
Asn Lys Leu Ala Asp Ser Asn Ala Pro Ala Met Lys Thr Leu Met Gly
65           70           75           80
Ala Trp Tyr Ala Ile Gly Ala Gly Gly Lys Arg Asp Trp Ile Lys Ala
85           90           95
Arg Ile Trp Phe Glu Lys Ala Ala Thr Glu Gly Asp Thr Arg Ala Ala
100          105          110
Tyr Pro Leu Gly Leu Leu Tyr Ser Ala Gly Leu Gly Thr Pro Ile Asp
115          120          125
Tyr Asp Lys Ala Phe Tyr Trp Leu Ser Ile Ala Ala Arg Gln Asn Ile
130          135          140
Pro Asp Ala Gln Tyr Arg Leu Ala Gly Leu Tyr Gln Glu Gly Lys Gly
145          150          155          160
Thr Ala Lys Ser Glu Arg Glu Phe Ala Tyr Trp Val Lys Lys Ala Ala
165          170          175
Gly Asn Gly His Ile Asp Ala Gln Arg Ala Met Gly Met Ile Leu His
180          185          190
Tyr Gly Leu Gly Val His Lys Asn Leu Pro Glu Ser Val Lys Trp Phe
195          200          205
Glu Lys Ala Ala Asn Ala Gly Asn Ala Thr Ala Gln Tyr Tyr Leu Gly
210          215          220
Met Asp Tyr Met Asn Gly Asn Gly Leu Ala Lys Asn Glu Arg Glu Gly
225          230          235          240
Glu Lys Trp Leu Tyr Arg Ala Ala Met Gln Asp His Leu Glu Ala Gln
245          250          255

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Thr Tyr Leu Gly Thr Ile Tyr Leu Lys Arg Lys Leu Gln Asn Glu Gly
 260 265 270
 Gln Pro Pro Glu Thr Ala Leu Ala Ile Gln Trp Met Glu Asn Ala Ala
 275 280 285
 Thr Arg Asn Asp Pro Tyr Ala Ile Arg Leu Leu Ser Met Val Tyr Arg
 290 295 300
 His Ile Pro Glu Leu Gln Asn Asn Ala Lys Gly Met Leu His Leu Arg
 305 310 315 320
 Arg Ser Ala Glu Met Gly Asn Ala Ser Ala Gln Phe Asp Leu Gly Arg
 325 330 335
 Thr Leu Phe Gln Asp Lys Asn Ser Pro Ala Lys Arg Lys Glu Ala Val
 340 345 350
 Val Trp Leu Glu Lys Ala Ala Gln Gln Asp Glu Arg Arg Ala Gln Ala
 355 360 365
 Phe Leu Gly Asn Met Tyr Tyr Tyr Gly Glu Phe Val Pro Val Asp Tyr
 370 375 380
 Val Lys Ala Leu Pro Leu Leu Met Arg Ala Ala Asp Lys Gly Asp Ser
 385 390 395 400
 Phe Ala Gln Tyr Thr Leu Gly Leu Ala Tyr Ile Asp Gly Asn Gly Ile
 405 410 415
 Ala Lys Asp Glu Arg Lys Ala Phe Ser Trp Leu Glu Lys Ser Ala Ser
 420 425 430
 Gln Asn Arg Ala Ser Ala Gln Tyr Phe Leu Gly Leu Met Tyr Leu Asp
 435 440 445
 Gly Thr Gly Thr Pro Val Asn Glu Glu Lys Gly Ile Arg Leu Leu Lys
 450 455 460
 Glu Leu Ala Lys Thr Gly Tyr Val Tyr Ala Gln Tyr Lys Leu Gly Thr
 465 470 475 480
 Tyr Ala His Ser Gly Leu His Met Ala Lys Asp Leu Ala Glu Ala Arg
 485 490 495
 Lys Trp Tyr Gln Leu Ala Ala Ser Gln Asp His Ile Lys Ala Lys Tyr
 500 505 510
 Trp Leu Gly Met Ser Leu Phe Gln Gly Pro Asp Ser Glu Gln Asp Arg
 515 520 525
 Lys Lys Gly Val Tyr Trp Phe Thr Glu Ala Ala Lys Gln Asp Asp Pro
 530 535 540
 Asp Ala Gln Leu Glu Leu Gly Lys Ser Leu Leu Tyr Gly Asp Gly Ile
 545 550 555 560
 Asp Lys Asn Glu Lys Gln Ala Cys Thr Trp Phe Lys Lys Ala Ala Asn
 565 570 575
 Asn Gln Gln His Thr Gly Gln Tyr Tyr Ala Gly Met Cys Leu Met Arg
 580 585 590
 Gly Ile Asn Gly Pro Val Asp Ile Pro Lys Gly Met Ser Leu Ile Glu
 595 600 605
 Met Ser Ala Asn Asn Lys Val Ser Met Ala Gln Phe Gln Leu Gly Lys
 610 615 620
 Leu Tyr Glu Tyr Gly Leu Lys Glu Leu Pro Lys Asp Ile Ser Lys Ala
 625 630 635 640
 Ile Gly Trp Tyr Thr Arg Ala Ala Glu Asn Gly His Ala Thr Ala Gln
 645 650 655

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Tyr Arg Leu Gly Lys Leu Tyr Leu Lys Ala Asp Thr Pro Leu Lys Asn
 660 665 670
 Ile Pro Leu Gly Leu Glu Phe Leu Glu Lys Ser Ala Ser Gln Asn Ile
 675 680 685
 Thr Ser Ala Ile Phe Asp Leu Gly Asn Ile Tyr Tyr Asp Gly Lys Ile
 690 695 700
 Val Lys Gln Asp Met Ala Lys Ala Leu Asn Tyr Phe Gln Lys Gly Thr
 705 710 715 720
 Gly Leu Gly His Leu Pro Ser Gln Asn Phe Val Gly Phe Met Ile Glu
 725 730 735
 Asn Gly Ser Gly Val Lys Lys Asp Lys Glu Lys Ala Cys Lys Ile Tyr
 740 745 750
 Asp Glu Thr Gly Arg Arg Gly Ser Ala Tyr Gly Leu Tyr Arg Tyr Gly
 755 760 765
 Leu Cys Gln Leu Ser Asp Pro Asp Pro Ser Pro Glu Asn Gln Lys Lys
 770 775 780
 Ala Phe Ile Leu Phe Glu Gln Ala Ala Arg Lys Asn Leu Ala Asp Ala
 785 790 795 800
 Gln Tyr Phe Leu Ala Leu Cys Tyr Glu Tyr Gly Lys Gly Thr Pro Lys
 805 810 815
 Asn Pro Gly Glu Ala Ile Glu Trp Tyr Arg Arg Ala Ser Glu Asn Asp
 820 825 830
 Lys Pro Glu Ala Leu Tyr Gln Leu Gly Met Leu Tyr Ile Thr Ser Pro
 835 840 845
 Ser Pro Arg Gln Asn Ile Pro Leu Gly Leu Asp Tyr Leu Glu Lys Ala
 850 855 860
 Ala Ala Arg Asp Leu Ser Ser Ala Phe Asn Glu Leu Gly Arg Ile Tyr
 865 870 875 880
 Tyr Asp Gly Lys Ile Val Arg Gln Asp Leu Lys Lys Ser Val Phe Trp
 885 890 895
 Tyr Arg Lys Gly Ala Gln Ser Gly Asp Thr Arg Ser Gln Asn Asp Leu
 900 905 910
 Ala Tyr Met Met Glu Tyr Gly Lys Gly Leu Glu Lys Asp Glu Lys Ala
 915 920 925
 Ala Cys Thr Met Tyr Glu Lys Thr Ala Gly Glu Lys Asn Ala Tyr Gly
 930 935 940
 Gln Phe Arg Leu Gly Leu Cys Tyr Leu Asn Gly Lys Gly Gly Lys Ala
 945 950 955 960
 Lys Asp Gln Arg Glu Ala Val Arg Leu Phe Glu Ser Ala Ala Gly Gln
 965 970 975
 Asn Leu Ala Ser Ala Gln Tyr Phe Leu Gly Ile Tyr His Lys Glu Gly
 980 985 990
 Lys Gly Val Val Lys Asn Met Asn Glu Ala Phe Lys Trp Tyr Leu Thr
 995 1000 1005
 Ala Ala Asp Asn Gly His Val Ser Ser Met Phe Glu Val Gly Lys
 1010 1015 1020
 Met Phe Ala Asn Gly Arg Gly Thr Glu Arg Asp Asp Lys Lys Ala
 1025 1030 1035
 Phe His Trp Phe Glu Lys Ala Ala Glu Asn Gly Ser Asp Ser Ala
 1040 1045 1050
 Leu Thr Gln Leu Gly Ile Met Tyr Tyr Lys Gly Leu Gly Ile Ser

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1055	1060	1065
Ala Asp Lys Ser Lys Ala Ala Ser Phe Phe Leu Lys Ala Ala Glu 1070 1075 1080		
Lys Asn Asn Ser Tyr Ala Gln His Trp Leu Gly Tyr Met Tyr Leu 1085 1090 1095		
Tyr Gly Lys Gly Leu Glu Lys Asn Gly Glu Leu Ala Asn Gln Trp 1100 1105 1110		
Leu Ser Lys Ala Ala Asp Gln Asn Glu Thr Gly Ala Ile Phe Glu 1115 1120 1125		
Leu Gly Lys Gln Tyr Trp Tyr Gly Met Gly Val Pro Val Asn Pro 1130 1135 1140		
Glu Lys Ala Ile Val Leu Leu Gln Lys Ala Gly Asn Asp Gly His 1145 1150 1155		
Val Ile Ala Gln Arg Ile Leu Gly Tyr Ile Tyr Ala Asp Gly Gly 1160 1165 1170		
Pro Glu Lys Gly Ile Pro Leu Asp Phe Glu Lys Ala Val Gln Trp 1175 1180 1185		
Phe Glu Lys Ala Ala Arg Gln Asp Asp Ala Ala Gly Lys Met Gly 1190 1195 1200		
Leu Ala Leu Leu Thr Leu Thr Gly Lys Gly Thr Pro Lys Asn Glu 1205 1210 1215		
Glu Lys Gly Ile Arg Leu Leu Thr Gln Ser Ala Asn Met Asn Tyr 1220 1225 1230		
Pro Ser Ala Met Glu Leu Leu Gly Asp Phe Tyr Arg Glu Glu Lys 1235 1240 1245		
Gly Asp Lys Lys Glu Ala Glu Lys Trp Tyr Arg Arg Ala Ala Glu 1250 1255 1260		
Thr Gly Asp Lys Thr Val Ile Glu Ser Met Lys Lys Lys Gly Val 1265 1270 1275		
Tyr Pro Ala Asn Pro 1280		

<210> SEQ ID NO 65
 <211> LENGTH: 1245
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 65

```

atgccgtatt ggtccggcgc tcatctgcgc tacaatgaca gcaaaataaa atttcccgtc    60
atgactctat ccctgttccg aaaaattcat atgaaagctg cctttgcctg ctgcgctttc    120
ctgatagcct tgcctttttc cgcaagccat gcaggaaata tcgtccattc cggggtgccc    180
ggagcgcgcc ttattgtcgg gacggaaaac cggaaacctc cggaaaccgt ctctgaaaac    240
aatgttcccg aagcagcccg ttcaccagtc gatcagaacg acgcaaaaac acagtttcat    300
cttgggctga tgtcgaaaaa cggtatgga gtcctgtcgc acccggtaaa agcccgcgaa    360
tggtttcgca aagcggcccg acaaaattac cagcccgcgc aataccagct tgcctcatg    420
caattttcag gcacagggcg cacagaaaac aaaagtgcgc cgatcgaaca attcaaaaaa    480
ctggcttctg aaggatatgc tcccgcccaa tacacgctcg gatatctgaa cctgaaaggc    540
gatggcatcc cgcaaaatc cggagaagcc cgcttctggt tcgaaaaggc cgccgcaaaa    600
aacgatgtgc gcgcaaccgc tgcactggca tggctttatc tgaaggcgt cggcgctccc    660
    
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atcgatgaga aaaaagcgc cgtcctgttt gaaaaagctg ccaatatggg tgacgcatac   720
agccaggatc aattcggcat gatgctcggc cagggaaactg gcatgaatgc cgaaccggaa   780
aaagcgtttt tatggattga aaaagcggca aaccagcaat atccgggtgc cgaataccat   840
atggccatga tgtacctgac aggttcggga acggaaaaaa atccggagct agccgtcaaa   900
tggttgaaa aagccgcttt ccacgggaac gtcgatgccc agaatttcta tgccagcctg   960
ctttatctcg gttatggaat caaacaggac attcccaggc ccatcgggta ttttaccgaa  1020
gcgccgagg gcgacacgc cgaatcccag ttcttgctgg gaaccatcta cgtcaaggga  1080
aatggcgttt taacgaacct gaaaactgcc cgaaactggt ttgaaaaagc ggaaaagaac  1140
ggacaccccg atgcaaaagc cgcactggaa aaactggatg aaatggaagg caaaggccgt  1200
gcaacacctg aaaagaccga taccggcaat cactatccat cttga                    1245

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<210> SEQ ID NO 66
<211> LENGTH: 414
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

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<400> SEQUENCE: 66

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```

Met Pro Tyr Trp Ser Gly Ala His Leu Arg Tyr Asn Asp Ser Lys Ile
1          5          10          15

Lys Phe Pro Val Met Thr Leu Ser Leu Phe Arg Lys Ile His Met Lys
20        25        30

Ala Ala Phe Ala Cys Cys Ala Phe Leu Ile Ala Leu Pro Phe Ser Ala
35        40        45

Ser His Ala Gly Asn Ile Val His Ser Gly Leu Pro Gly Ala Pro Val
50        55        60

Ile Val Gly Thr Glu Asn Arg Lys Pro Pro Glu Thr Val Ser Glu Asn
65        70        75        80

Asn Val Pro Glu Ala Ala Arg Ser Pro Val Asp Gln Asn Asp Ala Lys
85        90        95

Thr Gln Phe His Leu Gly Leu Met Ser Lys Asn Gly Tyr Gly Val Pro
100       105       110

Val Asp Pro Val Lys Ala Arg Glu Trp Phe Ala Lys Ala Ala Gly Gln
115       120       125

Asn Tyr Gln Pro Ala Gln Tyr Gln Leu Ala Leu Met Gln Phe Ser Gly
130       135       140

Thr Gly Gly Thr Glu Asn Lys Ser Ala Ala Ile Glu Gln Phe Lys Lys
145       150       155       160

Leu Ala Ser Glu Gly Tyr Ala Pro Ala Gln Tyr Thr Leu Gly Tyr Leu
165       170       175

Asn Leu Lys Gly Asp Gly Ile Pro Gln Asn Ser Gly Glu Ala Arg Phe
180       185       190

Trp Phe Glu Lys Ala Ala Ala Lys Asn Asp Val Arg Ala Thr Ala Ala
195       200       205

Leu Ala Trp Leu Tyr Leu Lys Gly Val Gly Ala Pro Ile Asp Glu Lys
210       215       220

Lys Ala Ala Val Leu Phe Glu Lys Ala Ala Asn Met Gly Asp Ala Tyr
225       230       235       240

Ser Gln Asp Gln Phe Gly Met Met Leu Gly Gln Gly Thr Gly Met Asn
245       250       255

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Ala Glu Pro Glu Lys Ala Phe Leu Trp Ile Glu Lys Ala Ala Asn Gln
 260 265 270

Gln Tyr Pro Val Ala Glu Tyr His Met Ala Met Met Tyr Leu Thr Gly
 275 280 285

Ser Gly Thr Glu Lys Asn Pro Glu Leu Ala Val Lys Trp Leu Glu Lys
 290 295 300

Ala Ala Phe His Gly Asn Val Asp Ala Gln Asn Phe Tyr Ala Ser Leu
 305 310 315 320

Leu Tyr Leu Gly Tyr Gly Ile Lys Gln Asp Ile Pro Arg Ala Ile Gly
 325 330 335

Tyr Phe Thr Glu Ala Ala Glu Gly Gly His Ala Glu Ser Gln Phe Leu
 340 345 350

Leu Gly Thr Ile Tyr Val Lys Gly Asn Gly Val Leu Thr Asn Leu Lys
 355 360 365

Thr Ala Arg Asn Trp Phe Glu Lys Ala Glu Lys Asn Gly His Pro Asp
 370 375 380

Ala Lys Ala Ala Leu Glu Lys Leu Asp Glu Met Glu Gly Lys Gly Arg
 385 390 395 400

Ala Thr Pro Glu Lys Thr Asp Thr Gly Asn His Tyr Pro Ser
 405 410

<210> SEQ ID NO 67
 <211> LENGTH: 630
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 67

```

atggagtgct tgagcctgaa aaaaacaggc tgttccctga tggcaatgac cctttgtttc    60
attttccttt cggcgtgcaa accaaaaaac attttgaca ccaactccgct gaaagcggaa    120
atcgcgcagg aatttgccaa agccgataaa ggcgacgcgg ctgcccgggt caatatcggc    180
ctcatgtact tcaggggaga aggtgtccct caggattacc ggcagggcgc cgtctggttt    240
gaaaaagcgg ccatgcaagg gctcggcgaa gcccaataca atatgggagc catgctggag    300
cagggactcg gcacagaaaa gaatgccgcc acagccagaa cctggtatga aaaagcggca    360
gcacagggca acgctcacgc ccagtacaac cttgccaggc tttacctgac aggcaaaggc    420
acgtcacaaa acatcggcaa ggcaggtgaa tggatgcaaa aagcggcagc acagggaatg    480
acagaagcaa aagaacgctt ttccattcta ttcgacagtc aatccggctc atttaaaccg    540
gcaaaaatcc agtcatggat tgaacaatcg gtccagaccg gtggggaaaa agcgaaaatc    600
ctgcaggaaa aaataagaag ccatagctga    630
    
```

<210> SEQ ID NO 68
 <211> LENGTH: 209
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 68

Met Glu Cys Leu Ser Leu Lys Lys Thr Gly Cys Ser Leu Met Ala Met
 1 5 10 15

Thr Leu Cys Phe Ile Phe Leu Ser Ala Cys Lys Pro Lys Thr Ile Leu
 20 25 30

Asp Thr Thr Pro Leu Lys Ala Glu Ile Ala Gln Glu Phe Ala Lys Ala

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35	40	45
Asp Lys Gly Asp Ala Ala Ala Arg Phe Asn Ile Gly Leu Met Tyr Phe		
50	55	60
Arg Gly Glu Gly Val Pro Gln Asp Tyr Arg Gln Ala Arg Val Trp Phe		
65	70	75 80
Glu Lys Ala Ala Met Gln Gly Leu Gly Glu Ala Gln Tyr Asn Met Gly		
	85	90 95
Ala Met Leu Glu Gln Gly Leu Gly Thr Glu Lys Asn Ala Ala Thr Ala		
	100	105 110
Arg Thr Trp Tyr Glu Lys Ala Ala Ala Gln Gly Asn Ala His Ala Gln		
	115	120 125
Tyr Asn Leu Ala Arg Leu Tyr Leu Thr Gly Lys Gly Thr Ser Gln Asn		
	130	135 140
Ile Gly Lys Ala Gly Glu Trp Met Gln Lys Ala Ala Ala Gln Gly Met		
	145	150 155 160
Thr Glu Ala Lys Glu Arg Phe Ser Ile Leu Phe Asp Ser Gln Ser Gly		
	165	170 175
Ser Phe Lys Pro Ala Lys Ile Gln Ser Trp Ile Glu Gln Ser Val Gln		
	180	185 190
Thr Gly Gly Glu Lys Ala Lys Ile Leu Gln Glu Lys Ile Arg Ser His		
	195	200 205

Ser

<210> SEQ ID NO 69
 <211> LENGTH: 1461
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 69

```

atgaatgaat cgacgtcttt tcaaacagtg aaatcccatt tcatgaaaac cggctttatt    60
tcaatctgca aacgggggat tcccgctctg ctcttctcat tcatggcttc cctgtcgaat    120
ccggcagcgg gacaaaccga cgccgatgcg aaaaccgaag gcatcaggct gtatcaggaa    180
aaaaaataca tcgaagccct tcctacctc aatgcgcccg atgcccmeta agaccgctt    240
gtccagagcg cgctcgcaa tatgttttca atggggctgg gcgtcgacgt caatcaggaa    300
aaagccttcg actggtacct gaaagccgcc aaacagaata acgcatggc ccagctctac    360
gtcgcctata tgcttgmeta aggacttggc gtccgmeta acgacaggga agccttcaac    420
tggtacmeta aagcagccga acagaacgta ccgaacgcac aatacaaaact cggcactctt    480
tacgmetaaag gcatcggcac ccgaatcaat ctgmetaaag ccctgaaactg gtaccggmeta    540
gccgcccgaag gagcctgtc gggagcggaa gtmetaactgg gccgctgtg cagcgaaggc    600
atcggcgctca aacgcgacta taccgaagcc gcccgctggt tctaccctgc agccgmetaa    660
ggggacgtca tggcacagac tgccttggc ttctctttg metaacgggct tggcgtccag    720
caggatgaag ccttcgctat cagctggtac tccaaagcgg ccgmetaagg tttcgtctct    780
gcccagaaca atctgggtta cctgtacgac aacggcatcg gcgtcctgcg tgactacacc    840
accgccagaa aatggtacga agccgcccga aaacagggga atgtggaagc ccagtccaac    900
ctcggccagc tctacacctc tggccacggc accgttcagg actacggcmeta ggccgcccga    960
tggctcmeta aagccgctgc caaaggccac ccgmetaagccc tgaacaaactc cggcatggcc    1020
    
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agtcttgaagc gcatgggctg tccgatggat cgcgtcaagg ccggagaata cttccggaaa 1080
gccgcctcgc tcggcaacgc ccatgcccag tacaatctgg ctaccctgta cgtccagcat 1140
ccagacgcac tgacaaaaga caagcgaagc aaaaccgacg ctttggccct gcaatggttc 1200
aaaaaaagcg cggctgcccgg tcaccctgcc gcgatggcct atctggctga cgtctatacc 1260
tacggcaaac tgggacagag acccaaccgg aaactggccg ccagctggaa acagaaagcc 1320
gatgccgccc aagaaaaacg caaaagagca cagaccatca ctccgctgcc cagaaccgaa 1380
gcggccagac ccgcaaaaac cgaaccggtc aagcccgta caaaggattt caatcccaac 1440
gaccgcgact ggctggaata g 1461

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<210> SEQ ID NO 70

<211> LENGTH: 486

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 70

```

Met Asn Glu Ser Thr Ser Phe Gln Thr Val Lys Ser His Phe Met Lys
1          5          10          15
Thr Gly Phe Ile Ser Ile Cys Lys Arg Ala Ile Pro Val Leu Leu Phe
20          25          30
Ser Phe Met Ala Ser Leu Ser Asn Pro Ala Ala Gly Gln Thr Asp Ala
35          40          45
Asp Ala Lys Thr Glu Gly Ile Arg Leu Tyr Gln Glu Lys Lys Tyr Ile
50          55          60
Glu Ala Leu Pro Tyr Leu Asn Ala Pro Asp Ala Gln Lys Asp Pro Leu
65          70          75          80
Val Gln Ser Ala Leu Gly Asn Met Phe Ser Met Gly Leu Gly Val Asp
85          90          95
Val Asn Gln Glu Lys Ala Phe Asp Trp Tyr Leu Lys Ala Ala Lys Gln
100         105         110
Asn Asn Ala Met Ala Gln Leu Tyr Val Ala Tyr Met Leu Glu Lys Gly
115         120         125
Leu Gly Val Arg Lys Asn Asp Arg Glu Ala Phe Asn Trp Tyr Lys Lys
130         135         140
Ala Ala Glu Gln Asn Val Pro Asn Ala Gln Tyr Lys Leu Gly Thr Leu
145         150         155         160
Tyr Glu Lys Gly Ile Gly Thr Arg Ile Asn Leu Lys Glu Ala Leu Asn
165         170         175
Trp Tyr Arg Lys Ala Ala Glu Gly Gly Leu Ser Gly Ala Gln Val Lys
180         185         190
Leu Gly Arg Leu Tyr Ser Glu Gly Ile Gly Val Lys Arg Asp Tyr Thr
195         200         205
Glu Ala Ala Arg Trp Phe Tyr Pro Ala Ala Glu Lys Gly Asp Val Met
210         215         220
Ala Gln Thr Ala Leu Ala Phe Leu Phe Glu Asn Gly Leu Gly Val Gln
225         230         235         240
Gln Asp Asp Ala Phe Ala Ile Ser Trp Tyr Ser Lys Ala Ala Glu Lys
245         250         255
Gly Phe Ala Pro Ala Gln Asn Asn Leu Gly Tyr Leu Tyr Asp Asn Gly
260         265         270
Ile Gly Val Leu Arg Asp Tyr Thr Thr Ala Arg Lys Trp Tyr Glu Ala

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275		280		285	
Ala	Ala Lys Gln Gly Asn Val	Glu Ala Gln Phe	Asn Leu Gly Gln Leu		
	290	295	300		
Tyr	Thr Leu Gly His Gly Thr Val Gln Asp Tyr	Gly Lys Ala Ala Glu			
	305	310	315	320	
Trp	Leu Glu Lys Ala Ala Ala Lys Gly His Pro	Lys Ala Leu Asn Asn			
		325	330	335	
Leu	Gly Met Ala Ser Leu Asp Gly Met Gly Val	Pro Met Asp Arg Val			
		340	345	350	
Lys	Ala Gly Glu Tyr Phe Arg Lys Ala Ala Leu	Leu Gly Asn Ala His			
		355	360	365	
Ala	Gln Tyr Asn Leu Ala Thr Leu Tyr Val Gln	His Pro Asp Ala Leu			
		370	375	380	
Thr	Lys Asp Lys Arg Ser Lys Thr Asp Ala Leu	Ala Leu Gln Trp Phe			
		385	390	395	400
Lys	Lys Ser Ala Ala Ala Gly His Pro Ala Ala	Met Ala Tyr Leu Ala			
		405	410	415	
Asp	Val Tyr Thr Tyr Gly Lys Leu Gly Gln Arg	Pro Asn Arg Lys Leu			
		420	425	430	
Ala	Ala Ser Trp Lys Gln Lys Ala Asp Ala Ala	Glu Glu Lys Arg Lys			
		435	440	445	
Arg	Ala Gln Thr Ile Thr Pro Leu Pro Arg Thr	Glu Ala Ala Arg Pro			
		450	455	460	
Ala	Lys Thr Glu Pro Val Lys Pro Val Thr Lys	Asp Phe Asn Pro Asn			
		465	470	475	480
Asp	Arg Asp Trp Leu Glu				
		485			

<210> SEQ ID NO 71
 <211> LENGTH: 885
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 71

```

atgatcaatc gttctttcaa gaatgcaaaa attatatatta tcctttttct tttatcattt    60
tgtataaatg tgtttgccgg tgatatagaa aaaggattaa agctgttcaa aaaacaggaa    120
tatgaaaagg cattgcctta ttttcagaaa ccggttgccg aaaaaaatcc ggacgttcag    180
gctgcattgg gatacatgta tcgtgaaggt ttagccgttc cgaaagacat tcaaaaggca    240
tttgatttat ttctggagtc agccagacaa aacaatccga gaggacagta tggaatggga    300
accatgtatg accttgact gattgtcaaa caggataaag aaaaagcgtt caaatggat    360
atgtatgcag ctgaaaatgg atacaagaat gcccaatata atattgggat tatgtatgcc    420
agaggaagag gaacaaaacg ggattacaaa aaagccaggg aatggtatga aaaagccgtt    480
ttgcagggac acaagggtgc tatgacgaat ttgggacttc tttattatcg gggatggggc    540
ggcccgaag attatgctaa atcagcagaa ctgaatacac gggcggctaa attaggtgat    600
gatatagccc aatataatct ggcacgtgat tatgaaaatg gtaccggcgt tccaaaagat    660
tacaagcagg ctgtttactg gtatttcaag ggggctgaaa atggaaatgc aatggcgtatg    720
gaacgattgt atgaagccta tcacctcaac cgattgggtt tgccgagaga cgatgaaaaa    780
gcgcttact gggctgaaaa agcccgggaa acccgcgta aacaggaga actggctccc    840
    
```

-continued

gagtcaatga gcattatoga aaaaatagag cgaatctggt tctga

885

<210> SEQ ID NO 72

<211> LENGTH: 294

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 72

```

Met Ile Asn Arg Ser Phe Lys Asn Ala Lys Ile Ile Phe Ile Leu Phe
1           5           10           15
Leu Leu Ser Phe Cys Ile Asn Val Phe Ala Gly Asp Ile Glu Lys Gly
20           25           30
Leu Lys Leu Phe Lys Lys Gln Glu Tyr Glu Lys Ala Leu Pro Tyr Phe
35           40           45
Gln Lys Pro Val Ala Gln Lys Asn Pro Asp Val Gln Ala Ala Leu Gly
50           55           60
Tyr Met Tyr Arg Glu Gly Leu Ala Val Pro Lys Asp Ile Gln Lys Ala
65           70           75           80
Phe Asp Leu Phe Leu Glu Ser Ala Arg Gln Asn Asn Pro Arg Gly Gln
85           90           95
Tyr Gly Met Gly Thr Met Tyr Asp Leu Gly Leu Ile Val Lys Gln Asp
100          105          110
Lys Glu Lys Ala Phe Lys Trp Tyr Met Tyr Ala Ala Glu Asn Gly Tyr
115          120          125
Lys Asn Ala Gln Tyr Asn Ile Gly Ile Met Tyr Ala Arg Gly Arg Gly
130          135          140
Thr Lys Arg Asp Tyr Lys Lys Ala Arg Glu Trp Tyr Glu Lys Ala Val
145          150          155          160
Leu Gln Gly His Lys Gly Ala Met Thr Asn Leu Gly Leu Leu Tyr Tyr
165          170          175
Arg Gly Trp Gly Gly Pro Lys Asp Tyr Ala Lys Ser Ala Glu Leu Asn
180          185          190
Thr Arg Ala Ala Lys Leu Gly Asp Asp Ile Ala Gln Tyr Asn Leu Ala
195          200          205
Arg Asp Tyr Glu Asn Gly Thr Gly Val Pro Lys Asp Tyr Lys Gln Ala
210          215          220
Val Tyr Trp Tyr Phe Lys Gly Ala Glu Asn Gly Asn Ala Met Ala Met
225          230          235          240
Glu Arg Leu Tyr Glu Ala Tyr His Leu Asn Arg Leu Gly Leu Pro Arg
245          250          255
Asp Asp Glu Lys Ala His Tyr Trp Ala Glu Lys Ala Arg Glu Thr Arg
260          265          270
Arg Lys Thr Gly Glu Leu Ala Pro Glu Ser Met Ser Ile Ile Glu Lys
275          280          285
Ile Glu Arg Ile Trp Phe
290

```

<210> SEQ ID NO 73

<211> LENGTH: 888

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 73

-continued

```

atggttgatt ttttatttaa aaatattcaga atttatattta tctctctttt ggtttcattc   60
agcctgaacc ttgaagccgc cgataccata aaccaaggac tggatttatt tcaaaaaaaaa   120
caattcgaaa aagcgttacc ctatctggaa gcatcacatt ccaaaaaacga tctctgagta   180
caagcggcat taggatatat gtatcgggat gggctgggag tcgaaaaaga ttatcagaaa   240
gcgtttatgt tatttctgga gtctgccaaa caatccaatc ccaagggaca gttcggcgta   300
ggaagtatgt atgatcgtgg attttttgtt aaacgtaata aggaaaaagc gtttaaatgg   360
tatctatacg cagctgagaa tggttatagt tctgctcaaa ataatgtagc ctggtcttat   420
gtacatggag agggaattcc taaagattac aaaaaagcca gagaatggta tgaaaaagca   480
atgattcaag gacattcga tgcctatggt ggtttatcct ttatgtatta ttggggaaaa   540
ggcgtaaaaa aagatcgtaa taaagcagcg gaactggaca aacgggcggc agcgttggga   600
aacagaaaag gacagtcaa tctgggccgg gattacgaag atggaacagg tgttcccaaa   660
gactacaagc aagccgttta ctgggtacttt aaagccgctg aaaacggtga tccccggct   720
atggaacgtc tttatgaagc ctatcaoctc aaccgattgg gtctgaagag agacgatgaa   780
aaagcccatt actgggctga gaaagcccgg gaaacccgtc gaaaaacggg agaagtcgat   840
ccgcgttcat tgtctgtcat tgaaaaagtc agaatgatct ggttctga   888

```

<210> SEQ ID NO 74

<211> LENGTH: 295

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 74

```

Met Val Asp Phe Leu Phe Lys Asn Ile Arg Ile Ile Phe Ile Leu Ser
 1           5           10          15
Leu Val Ser Phe Ser Leu Asn Leu Glu Ala Ala Asp Thr Ile Asn Gln
 20          25          30
Gly Leu Asp Leu Phe Gln Lys Lys Gln Phe Glu Lys Ala Leu Pro Tyr
 35          40          45
Leu Glu Ala Ser His Ser Lys Asn Asp Pro Arg Val Gln Ala Ala Leu
 50          55          60
Gly Tyr Met Tyr Arg Asp Gly Leu Gly Val Glu Lys Asp Tyr Gln Lys
 65          70          75          80
Ala Phe Met Leu Phe Leu Glu Ser Ala Lys Gln Ser Asn Pro Lys Gly
 85          90          95
Gln Phe Gly Val Gly Ser Met Tyr Asp Arg Gly Phe Phe Val Lys Arg
100         105         110
Asn Lys Glu Lys Ala Phe Lys Trp Tyr Leu Tyr Ala Ala Glu Asn Gly
115         120         125
Tyr Ser Ser Ala Gln Asn Asn Val Ala Trp Ser Tyr Val His Gly Glu
130         135         140
Gly Ile Pro Lys Asp Tyr Lys Lys Ala Arg Glu Trp Tyr Glu Lys Ala
145         150         155         160
Met Ile Gln Gly His Ser Asn Ala Met Val Gly Leu Ser Phe Met Tyr
165         170         175
Tyr Trp Gly Lys Gly Val Lys Lys Asp Arg Asn Lys Ala Ala Glu Leu
180         185         190
Asp Lys Arg Ala Ala Ala Leu Gly Asn Arg Lys Gly Gln Phe Asn Leu
195         200         205

```

-continued

Gly Arg Asp Tyr Glu Asp Gly Thr Gly Val Pro Lys Asp Tyr Lys Gln
 210 215 220

Ala Val Tyr Trp Tyr Phe Lys Ala Ala Glu Asn Gly Asp Pro Met Ala
 225 230 235 240

Met Glu Arg Leu Tyr Glu Ala Tyr His Leu Asn Arg Leu Gly Leu Lys
 245 250 255

Arg Asp Asp Glu Lys Ala His Tyr Trp Ala Glu Lys Ala Arg Glu Thr
 260 265 270

Arg Arg Lys Thr Gly Glu Val Asp Pro Arg Ser Leu Ser Val Ile Glu
 275 280 285

Lys Val Arg Met Ile Trp Phe
 290 295

<210> SEQ ID NO 75
 <211> LENGTH: 954
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 75

```

atggaagcc ataaagaat caagtcaatt ttaaaagaac ttagccagat gaaaaaacia 60
tctttatttc tcatcataat tgcaatgatt ctgacaatat cgccttgtag agaaaaagag 120
tataaatacc tcgcagatga aaaagcagga atcgaatact atcaaaacgg atcatatgat 180
aaagcgttgg catcactaaa aaaaacatat ggttcogga gtatggaagc cgcttattat 240
cttgcgaaa tgtatcgta cggtaatggg gttgaaaaag atagaatcgt ttcttgcaat 300
tactatcaaa aatcagcaga agggggaaac aggaaagcct ttttgagagc aggaacctgt 360
catataccgg ataccagaga tggagaagga ttcaaagaaa cattcaaatg gtttaaaaa 420
gcgtctgaag aattaaaga aaccgatttg aatgaagcag aaaaaaaga catgtatata 480
aggcttgga ttatgtatta cgtggaaaa ggtacactac aggattggag tgaagccgcg 540
aagtggtttg aaaaagctgc cgaatgggg gatgcctact cacagggagg tctggcgta 600
ttatattatt cggcgcgatg tgttttaact gacaggaaaa aagcgcgta ttgggcagaa 660
aaagcagcgg cacaaggtaa tagtacagga gaaggtattc ttggaatggt ataccaatat 720
agtcttccgc cggataaaga tatgagaaaa gctgtctatt ggtacaagaa atcagccgat 780
cagggaaatc caatatccca gtatcaattg gctgttatct atgaaaatgg tgatggtggt 840
ccaaaaaacc tcgaaaaagc gcgctattac tatgaacagg catcgaaaag caaatatgat 900
ctgacaaaaa acagtttaaa ggaatttgaa tcgcgaaaca aaagtttgca gtga 954

```

<210> SEQ ID NO 76
 <211> LENGTH: 317
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 76

Met Glu Ser His Lys Glu Ile Lys Ser Ile Leu Lys Glu Leu Ser Gln
 1 5 10 15

Met Lys Lys Gln Ser Leu Phe Leu Ile Ile Ile Ala Met Ile Leu Thr
 20 25 30

Ile Ser Ala Cys Thr Glu Lys Glu Tyr Lys Tyr Leu Ala Asp Glu Lys
 35 40 45

-continued

Ala Gly Ile Glu Tyr Tyr Gln Asn Gly Ser Tyr Asp Lys Ala Leu Ala
 50 55 60

Ser Leu Lys Lys Thr Tyr Gly Ser Gly Ser Met Glu Ala Ala Tyr Tyr
 65 70 75 80

Leu Gly Glu Met Tyr Arg Gln Gly Asn Gly Val Glu Lys Asp Arg Ile
 85 90 95

Val Ser Cys Asn Tyr Tyr Gln Lys Ser Ala Glu Gly Gly Asn Arg Lys
 100 105 110

Ala Phe Leu Arg Ala Gly Thr Cys His Ile Pro Asp Thr Arg Asp Gly
 115 120 125

Glu Gly Phe Lys Glu Thr Phe Lys Trp Phe Lys Lys Ala Ser Glu Glu
 130 135 140

Leu Lys Glu Thr Asp Leu Asn Glu Ala Glu Lys Lys Asp Met Tyr Ile
 145 150 155 160

Arg Leu Gly Ile Met Tyr Tyr Ala Gly Lys Gly Thr Leu Gln Asp Trp
 165 170 175

Ser Glu Ala Ala Lys Trp Phe Glu Lys Ala Ala Glu Met Gly Asp Ala
 180 185 190

Tyr Ser Gln Gly Gly Leu Ala Leu Leu Tyr Tyr Ser Gly Asp Gly Val
 195 200 205

Leu Thr Asp Arg Lys Lys Ala Arg Tyr Trp Ala Glu Lys Ala Ala Ala
 210 215 220

Gln Gly Asn Ser Thr Gly Glu Gly Ile Leu Gly Met Leu Tyr Gln Tyr
 225 230 235 240

Ser Leu Pro Pro Asp Lys Asp Met Arg Lys Ala Val Tyr Trp Tyr Lys
 245 250 255

Lys Ser Ala Asp Gln Gly Asn Pro Ile Ser Gln Tyr Gln Leu Ala Val
 260 265 270

Ile Tyr Glu Asn Gly Asp Gly Val Pro Lys Asn Leu Glu Lys Ala Arg
 275 280 285

Tyr Tyr Tyr Glu Gln Ala Ser Lys Ser Lys Tyr Asp Leu Thr Lys Asn
 290 295 300

Ser Leu Lys Glu Phe Glu Ser Arg Asn Lys Ser Leu Gln
 305 310 315

<210> SEQ ID NO 77
 <211> LENGTH: 1083
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 77

```

atgctgaaaa aatcattatc tctttcgatt atcctttttc tcttgtttc cttatcgggc    60
tgtgagaaga acggtgagga aaacaagagc aagggggatg ttttttattc cagaggggat    120
tatgaaaatg ccctgtctgt ttacaaacag gaggcaaac ggggcgattc gcatgccttg    180
ctcggtttgt gcaagatgta tgaggatgga aaggggggtca aggaaaacga tcggctggct    240
ttccagtatt gcatgcaggc ggccaacgg ggtgaggtga aggcacagaa gcgtctgggg    300
atgatgtatt acaagggtag cgggggtggcg aggaatgtgc atgaagcgcg tttctggttc    360
aatcagggcg ccttgtcgga cgatccggag gcgctttatt atctcgggat cgcttatttg    420
aaagggatcg gcggggagaa agactttcat caggcgcgatg acctttttga aagggcggcc    480
gatgagggac atgtgaatgc catgtggaag ctgtatgaga tgttcaatga aggaaccggt    540
    
```

-continued

```

gtcaggcagg acaggcagga ggcggtcaaa tggctgatga agctggcggc cgatggcgac   600
atacgcgcgc agttccttgc cggttcgtcg tatctgaccg gcaatggggg gaaggctgat   660
cctgcccagg cggtcagggt gtttgaaaag gcggcgcagc agggcagtgt tgatgccag   720
acttcaactgg ttttatatta tctgggcaac gagccagtcc gtctggatga agcgcgaagc   780
tgggcggcga aactggaaaa ccagacgggt tcgcgcgggt tcggggcggg gttcaggcgt   840
tttttcaactg agatgggaga gtggcctgag ccggaacgca cgcatgaacg gataaaagac   900
tggttcgatt ctcttttttc gacgaaagac ccggatctgc tttatgggtc cggtttttg   960
tacgagctgg gttatcaggg gcagcgcgat atggacaagg cgctgcacct ctataaagg   1020
gcggcggctc tggggaacgt gaatgcgatt aaaaaactga aagaactcca gtcggttttt   1080
taa                                                                                   1083

```

<210> SEQ ID NO 78

<211> LENGTH: 360

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 78

```

Met Leu Lys Lys Ser Leu Ser Leu Ser Ile Ile Leu Phe Leu Leu Phe
 1          5          10          15
Ser Leu Ser Gly Cys Glu Lys Asn Gly Glu Glu Asn Lys Ser Lys Gly
 20          25          30
Asp Val Phe Tyr Ser Arg Gly Asp Tyr Glu Asn Ala Leu Ser Val Tyr
 35          40          45
Lys Gln Glu Ala Lys Arg Gly Asp Ser His Ala Leu Leu Gly Leu Cys
 50          55          60
Lys Met Tyr Glu Asp Gly Lys Gly Val Lys Glu Asn Asp Arg Leu Ala
 65          70          75          80
Phe Gln Tyr Cys Met Gln Ala Ala Lys Arg Gly Glu Val Lys Ala Gln
 85          90          95
Lys Arg Leu Gly Met Met Tyr Tyr Lys Gly Thr Gly Val Ala Arg Asn
100          105          110
Val His Glu Ala Arg Phe Trp Phe Asn Gln Ala Ala Leu Ser Asp Asp
115          120          125
Pro Glu Ala Leu Tyr Tyr Leu Gly Ile Ala Tyr Leu Lys Gly Ile Gly
130          135          140
Gly Glu Lys Asp Phe His Gln Ala His Asp Leu Phe Glu Arg Ala Ala
145          150          155          160
Asp Glu Gly His Val Asn Ala Met Trp Lys Leu Tyr Glu Met Phe Asn
165          170          175
Glu Gly Thr Gly Val Arg Gln Asp Arg Gln Glu Ala Phe Lys Trp Leu
180          185          190
Met Lys Leu Ala Ala Asp Gly Asp Ile Arg Ala Gln Phe Leu Ala Gly
195          200          205
Ser Ser Tyr Leu Thr Gly Asn Gly Val Lys Ala Asp Pro Ala Glu Ala
210          215          220
Val Arg Trp Phe Glu Lys Ala Ala Gln Gln Gly Ser Val Asp Ala Gln
225          230          235          240
Thr Ser Leu Val Leu Tyr Tyr Leu Gly Asn Glu Pro Val Arg Leu Asp
245          250          255

```


-continued

Ala Gln Lys Ser Pro Glu Ile Met Asn Arg Ile Gly Phe Met Tyr Asp
50 55 60

Ala Gly Arg Gly Val Glu Arg Asn Gly Asn Ile Ala Phe Gln Trp Tyr
65 70 75 80

Arg Lys Ala Ala Glu Thr Gly Leu Ala Lys Ala Gln Tyr Asn Leu Gly
85 90 95

Leu Cys Phe Gln Asn Gly Ile Gly Val Lys Lys Asp Ile Asn Glu Ala
100 105 110

Ile Lys Trp Tyr Leu Lys Ala Ala Glu Gln Gly Tyr Pro Asp Ala Glu
115 120 125

Ser Lys Met Gly Tyr Leu Thr Val Thr Gly Lys Gly Val Lys Gln Asp
130 135 140

Phe Lys Gln Ala Met Gln Trp Tyr Arg Arg Ala Val Glu His Gly Asn
145 150 155 160

Ile His Ala Ile Pro Glu Leu Gly Ile Met Tyr Glu Glu Gly Leu Gly
165 170 175

Val Lys Lys Asp Lys Thr His Ala Val Gln Tyr Tyr Ile Met Gly Ala
180 185 190

Glu Lys Gly Asn Val Arg Ala Gln Phe Leu Leu Ala Glu Ala Tyr Arg
195 200 205

Tyr Gly Arg Gly Ile Lys Asn Asp Asp Glu Arg Ser Leu Tyr Trp Tyr
210 215 220

Lys Lys Ala Ala Glu Asn Gly Ser Ala Asp Ala Tyr Asp Ala Leu Gly
225 230 235 240

Ser Val Tyr Ala Asn Glu Gln Leu Gly Gln Lys Lys Asp Arg Lys Lys
245 250 255

Ala Gly Glu Met Val Glu Lys Ala Ile Glu Ile Arg Lys Gln Asn Gly
260 265 270

Glu Gly Asp Pro Asp Ala Arg Arg Arg Leu Lys Phe Phe Gly Val Lys
275 280 285

Leu Asp Asp
290

<210> SEQ ID NO 81

<211> LENGTH: 852

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 81

```

atgtgtaaaa actattcaat tgattacaga atatattttc gcgttttaat tcttttttta      60
ttggcttggt ttcttttaac gctgcatac gccaatgaca ttactcaggg attgaaatac      120
tataaaaaag aagaatatca aaaagcatat cgttattttt ccacaccggc tgctcaaaaag      180
aatccccggg ttcaacgatat attgggatat atgtatttaa agggctcttcg cgtaacgagc      240
gattatcaaa aggcgatggt ctgggatggc aaatcagctg atcagggtaa tcctcaagcc      300
atgtacgata ttggcggttat gtacgatttt ggacaggggt tgaaacagga tcatgaaaaa      360
gcgattcaat ggtatcagcg ctccgcttta aaaggatatg ctgacgcaca atataacctt      420
ggtatcgctt atgaaaaggg agaagggtact cagcaaaatt acgctaaagc gcgtgaatgg      480
tatcaaaaag cagtcacaca gggtaatgtc agtgctatgg tcaatttagg caatctttat      540
ggggaaggtc tcggcggtga aaaaaacgac tcaaaagcct ttgatcttta caaaaaagct      600

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```

gccgaaaaag gtgattcggc cgcccaatat aatctcggcg aatactaccg tgcgggotta 660
gtacccccac gagacctoga caaagccatc tactggtagc aaaagagcgc tgcagagggc 720
accatcaagg caatggacaa actggccaga atctatcggg tcggctacaa acacatccct 780
gccaatcagg ccttgtccga tgaatgggcc gacaaggctg caaaggcagc tgcaagggaa 840
gccgtcaggt aa 852

```

```

<210> SEQ ID NO 82
<211> LENGTH: 283
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

```

```

<400> SEQUENCE: 82

```

```

Met Cys Lys Asn Tyr Ser Ile Asp Tyr Arg Ile Tyr Phe Arg Val Leu
1           5           10          15
Ile Leu Phe Leu Leu Ala Cys Phe Leu Leu Thr Pro Ala Tyr Ala Asn
20          25          30
Asp Ile Thr Gln Gly Leu Lys Tyr Tyr Lys Lys Glu Glu Tyr Gln Lys
35          40          45
Ala Tyr Arg Tyr Phe Ser Thr Pro Ala Ala Gln Lys Asn Pro Arg Val
50          55          60
Gln Arg Ile Leu Gly Tyr Met Tyr Leu Lys Gly Leu Ala Val Lys Gln
65          70          75          80
Asp Tyr Gln Lys Ala Met Phe Trp Tyr Gly Lys Ser Ala Asp Gln Gly
85          90          95
Asn Pro Gln Ala Met Tyr Asp Ile Gly Val Met Tyr Asp Phe Gly Gln
100         105        110
Gly Val Lys Gln Asp His Glu Lys Ala Ile Gln Trp Tyr Gln Arg Ser
115        120        125
Ala Leu Lys Gly Tyr Ala Asp Ala Gln Tyr Asn Leu Gly Ile Ala Tyr
130        135        140
Glu Lys Gly Glu Gly Thr Gln Gln Asn Tyr Ala Lys Ala Arg Glu Trp
145        150        155        160
Tyr Gln Lys Ala Val Thr Gln Gly Asn Val Ser Ala Met Val Asn Leu
165        170        175
Gly Asn Leu Tyr Gly Glu Gly Leu Gly Gly Glu Lys Asn Asp Ser Lys
180        185        190
Ala Phe Asp Leu Tyr Lys Lys Ala Ala Glu Lys Gly Asp Ser Ala Ala
195        200        205
Gln Tyr Asn Leu Ala Glu Tyr Tyr Arg Ala Gly Leu Ala Thr Pro Arg
210        215        220
Asp Leu Asp Lys Ala Ile Tyr Trp Tyr Glu Lys Ser Ala Ala Glu Gly
225        230        235        240
Thr Ile Lys Ala Met Asp Lys Leu Ala Arg Ile Tyr Arg Val Gly Tyr
245        250        255
Lys His Ile Pro Ala Asn Gln Ala Leu Ser Asp Glu Trp Ala Asp Lys
260        265        270
Ala Ala Lys Ala Arg Ala Arg Glu Ala Val Arg
275        280

```

```

<210> SEQ ID NO 83
<211> LENGTH: 909

```

-continued

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 83

```

atgaaacagc cttatccata tttgttcttc aaaatgaaat ttctttcaaa aatattcttt    60
ttttatTTTT ttttaggtat aagcagcagt gtttttgtag ataatgcaaa aacagggatg    120
agtttggtcg agtcagcga atagagcaa gcgatgactt atttctgaa accggatgct    180
cagagaaatg cagaagtact taatcatatt ggttatatgt atgacaatgg attaggtggt    240
aggcaaaatc caaaactggc taatcaatgg tatagaaaag catcagaaaa aggatttcct    300
gctgctgatt tcaatattgg attaagtttt gaatcaggat ctggtgttaa aaaagatata    360
aacgaagcca taaaatggta tctcaaggct gcagaacagg gatatcccga tgcagaatcc    420
aaaatggggtt atctgaccgt aaccggaaaa ggcgtcaaac aggatttcaa acaggccatg    480
caatggtaacc ggcgtgacct tgaacatggc agtggtcatg ccatttcaga gctcggatc    540
ctgtatgagg aaggtcttgg cgtcaaaaag gacaagacct atgccgtcca gtattacatc    600
atgggtgccc aaaagggaaa cgtcagggca cagttctgtg tggcggaagc ctaccgttac    660
ggcctgggca tcaaaaacga cgatgagcgt tcgctctact ggtacaacaa ggcagcggaa    720
aacgggagtg cccgatgccta cgatgcgtta ggcagcgtgt atgccaacga acagctgggg    780
cagaaaaaag acaggaaaaa agcagggcga atgggtgaaa aagccattga aatccgcaag    840
cagaatggag agggcgatcc cgatgccaga cgccgcctga ggttttctgg ggtgaagctg    900
gatgactga                                         909

```

<210> SEQ ID NO 84

<211> LENGTH: 302

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 84

```

Met Lys Gln Pro Tyr Pro Tyr Leu Phe Phe Lys Met Lys Phe Leu Ser
1          5          10          15
Lys Ile Phe Phe Phe Tyr Phe Phe Leu Gly Ile Ser Ser Ser Val Phe
20          25          30
Ala Asp Asn Ala Lys Thr Gly Met Ser Leu Phe Glu Ser Ala Lys Tyr
35          40          45
Glu Gln Ala Met Thr Tyr Phe Leu Lys Pro Asp Ala Gln Arg Asn Ala
50          55          60
Glu Val Leu Asn His Ile Gly Tyr Met Tyr Asp Asn Gly Leu Gly Val
65          70          75          80
Arg Gln Asn Pro Lys Leu Ala Asn Gln Trp Tyr Arg Lys Ala Ser Glu
85          90          95
Lys Gly Phe Pro Ala Ala Asp Phe Asn Ile Gly Leu Ser Phe Glu Ser
100         105         110
Gly Ser Gly Val Lys Lys Asp Ile Asn Glu Ala Ile Lys Trp Tyr Leu
115         120         125
Lys Ala Ala Glu Gln Gly Tyr Pro Asp Ala Glu Ser Lys Met Gly Tyr
130         135         140
Leu Thr Val Thr Gly Lys Gly Val Lys Gln Asp Phe Lys Gln Ala Met
145         150         155         160
Gln Trp Tyr Arg Arg Ala Val Glu His Gly Ser Val His Ala Ile Ser

```

-continued

	165		170		175										
Glu	Leu	Gly	Ile	Leu	Tyr	Glu	Glu	Gly	Leu	Gly	Val	Lys	Lys	Asp	Lys
	180							185						190	
Thr	His	Ala	Val	Gln	Tyr	Tyr	Ile	Met	Gly	Ala	Glu	Lys	Gly	Asn	Val
	195						200					205			
Arg	Ala	Gln	Phe	Leu	Leu	Ala	Glu	Ala	Tyr	Arg	Tyr	Gly	Leu	Gly	Ile
	210					215					220				
Lys	Asn	Asp	Asp	Glu	Arg	Ser	Leu	Tyr	Trp	Tyr	Asn	Lys	Ala	Ala	Glu
	225				230					235					240
Asn	Gly	Ser	Ala	Asp	Ala	Tyr	Asp	Ala	Leu	Gly	Ser	Val	Tyr	Ala	Asn
			245						250					255	
Glu	Gln	Leu	Gly	Gln	Lys	Lys	Asp	Arg	Lys	Lys	Ala	Gly	Glu	Met	Val
			260					265						270	
Glu	Lys	Ala	Ile	Glu	Ile	Arg	Lys	Gln	Asn	Gly	Glu	Gly	Asp	Pro	Asp
		275					280						285		
Ala	Arg	Arg	Arg	Leu	Arg	Phe	Phe	Gly	Val	Lys	Leu	Asp	Asp		
	290					295					300				

<210> SEQ ID NO 85
 <211> LENGTH: 789
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 85

```

atgcggtgtg ttgaagctgg aaaataaccag caggccatga cgtatttcat gaaggaggat      60
gccagaaga atccggccgt gatcaatcgt attggctata tgtatgatta cgggctcggc      120
gttgaaaaaa acaggcaaat cagttttcag tggataaaa aagcaggaga aatgggggag      180
gccgtgcgc agttaaagt cggactatc tatgagaaag gttatggcgt tcctcaagat      240
ataaatatgg ctatcgaatg gtttcgtaa tctgcaaac aacaatatcc gaatgctgaa      300
gcgaaaatgg gttatctgac ggcgacagga aaaggaacca acaaaagttt tgtggaagct      360
atgaagtggg atcggtcagc tgcagagcat ggcaatatta atgttttctc agaaatagga      420
attatgtatg aggaggggta tggcgtcaaa aaaaacaaga accgtgctgt ccagtattac      480
attatgggag ctgataaagg aaatgccaaa gcgcagatc ttttaggtca tgccatccaa      540
tatggtcgtg gcataaaga tgatcctgaa cgggcgctgc actggtatcg taaagcggcg      600
gaacagggaa atgccgatgc cttgcaggca ctggggggta tttatgtaca tggcctgttg      660
aaccagaagg aagacaggga aaaaggtgaa aaatatattg aagaggcgat tagaatccgc      720
aagcaaacgg gtcagctaga tccggctgcg atgaggcggt tgccgctttt gggcatcgat      780
atagagtga                                     789
    
```

<210> SEQ ID NO 86
 <211> LENGTH: 262
 <212> TYPE: PRT
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 86

Met	Arg	Leu	Phe	Glu	Ala	Gly	Lys	Tyr	Gln	Gln	Ala	Met	Thr	Tyr	Phe
1				5					10					15	
Met	Lys	Glu	Asp	Ala	Gln	Lys	Asn	Pro	Ala	Val	Ile	Asn	Arg	Ile	Gly
		20					25					30			

-continued

Tyr Met Tyr Asp Tyr Gly Leu Gly Val Glu Lys Asn Arg Gln Ile Ser
 35 40 45

Phe Gln Trp Tyr Lys Lys Ala Gly Glu Met Gly Glu Ala Ala Ala Gln
 50 55 60

Phe Asn Val Gly Leu Phe Tyr Glu Lys Gly Tyr Gly Val Pro Gln Asp
 65 70 75 80

Ile Asn Met Ala Ile Glu Trp Phe Arg Lys Ser Ala Lys Gln Gln Tyr
 85 90 95

Pro Asn Ala Glu Ala Lys Met Gly Tyr Leu Thr Ala Thr Gly Lys Gly
 100 105 110

Thr Lys Gln Ser Phe Val Glu Ala Met Lys Trp Tyr Arg Ser Ala Ala
 115 120 125

Glu His Gly Asn Ile Asn Val Phe Ser Glu Ile Gly Ile Met Tyr Glu
 130 135 140

Glu Gly Tyr Gly Val Lys Lys Asn Lys Asn Arg Ala Val Gln Tyr Tyr
 145 150 155 160

Ile Met Gly Ala Asp Lys Gly Asn Ala Lys Ala Gln Tyr Leu Leu Gly
 165 170 175

His Ala Tyr Gln Tyr Gly Arg Gly Ile Lys Asp Asp Pro Glu Arg Ala
 180 185 190

Leu His Trp Tyr Arg Lys Ala Ala Glu Gln Gly Asn Ala Asp Ala Leu
 195 200 205

Gln Ala Leu Gly Gly Ile Tyr Val His Gly Leu Leu Asn Gln Lys Glu
 210 215 220

Asp Arg Glu Lys Gly Glu Lys Tyr Ile Glu Glu Ala Ile Arg Ile Arg
 225 230 235 240

Lys Gln Thr Gly Gln Leu Asp Pro Ala Ala Met Arg Arg Leu Arg Leu
 245 250 255

Leu Gly Ile Asp Ile Glu
 260

<210> SEQ ID NO 87
 <211> LENGTH: 939
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 87

```

atgttaaaat acatcctgtt aattcagata gtatcaatta tgaaaaaatt tttattttat    60
tttttgtttt tatcaatttc attactgata atgtcaggat gtgacgataa aaatatactt    120
gatgaaaag aagggtgtga atattataaa cagaacaat atgaaaaagc gttacctcta    180
ttgaaaaagt cggcttcatc aggtaattcc atcgcattctt attatttagg agagatgttt    240
tataaagggtg aaggagtga aaagaatgaa aataccgggtt gtcaacgata tctggaatct    300
tcaaaagggg gcaataaaaa cgcttatttg gtgacggcca tctgttttta tacaggtagg    360
ggattggaag aagaatgatgc cgaagcattc aaatgggcga aaaaagttga aaatgaaata    420
aatgaaacag gtcttggtga gtatgaccgg aaacattttt attttcta atgcagaatc    480
tatttgctcag gaagaggtagc tttgcaggat ttgagcggagg cagccaaatg gacagagaaa    540
atcgccaatt tgggtgatcc tgtctcacag ggtggttttag cggttttgta ctataatggc    600
aatggagttt tgaccgacag gaaaaaagcg cgttactggg cagaaaaagc ggccgcgcaa    660
ggcaatgata caggagaagt ggttcttggt atgttgaacc agtatagttt gcctcctgaa    720
    
```

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caaaatttgg aaaaagcagc agaatggtat ttgaaatcag cagctcaggg aaataccgca 780
gcacaacatc agttggcggg tatgtatgaa aagggagaag gggttccgca agatttgaaa 840
aaagcccgtt attattatga agaagcggca aagagtaagt atgaagaacc caaaaaagcc 900
cttgaggaat tcaatgccac gtatcaaaagt aaaaactaa 939

```

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<210> SEQ ID NO 88
<211> LENGTH: 312
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

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<400> SEQUENCE: 88

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```

Met Leu Lys Tyr Ile Leu Leu Ile Gln Ile Val Ser Ile Met Lys Lys
1           5           10          15
Phe Leu Phe Tyr Phe Leu Phe Leu Ser Ile Ser Leu Leu Ile Met Ser
20          25          30
Gly Cys Asp Asp Lys Asn Ile Leu Asp Glu Lys Glu Gly Val Glu Tyr
35          40          45
Tyr Lys Gln Lys Gln Tyr Glu Lys Ala Leu Pro Leu Leu Glu Lys Ser
50          55          60
Ala Ser Ser Gly Asn Ser Ile Ala Ser Tyr Tyr Leu Gly Glu Met Phe
65          70          75          80
Tyr Lys Gly Glu Gly Val Glu Lys Asn Glu Asn Thr Gly Cys Gln Arg
85          90          95
Tyr Leu Glu Ser Ser Lys Gly Gly Asn Lys Asn Ala Tyr Leu Val Thr
100         105        110
Ala Ile Cys Phe Tyr Thr Gly Arg Gly Leu Glu Lys Asn Asp Ala Glu
115        120        125
Ala Phe Lys Trp Ala Lys Lys Val Glu Asn Glu Ile Asn Glu Thr Gly
130        135        140
Leu Gly Glu Tyr Asp Arg Lys His Phe Tyr Phe Leu Met Gln Asn Ile
145        150        155        160
Tyr Leu Ser Gly Arg Gly Thr Leu Gln Asp Leu Ser Glu Ala Ala Lys
165        170        175
Trp Thr Glu Lys Ile Ala Asn Leu Gly Asp Pro Val Ser Gln Gly Gly
180        185        190
Leu Ala Val Leu Tyr Tyr Asn Gly Asn Gly Val Leu Thr Asp Arg Lys
195        200        205
Lys Ala Arg Tyr Trp Ala Glu Lys Ala Ala Ala Gln Gly Asn Asp Thr
210        215        220
Gly Glu Val Val Leu Gly Met Leu Asn Gln Tyr Ser Leu Pro Pro Glu
225        230        235        240
Gln Asn Leu Glu Lys Ala Ala Glu Trp Tyr Leu Lys Ser Ala Ala Gln
245        250        255
Gly Asn Thr Ala Ala Gln His Gln Leu Ala Val Met Tyr Glu Lys Gly
260        265        270
Glu Gly Val Pro Gln Asp Leu Lys Lys Ala Arg Tyr Tyr Tyr Glu Glu
275        280        285
Ala Ala Lys Ser Lys Tyr Glu Glu Pro Lys Lys Ala Leu Glu Glu Phe
290        295        300
Asn Ala Thr Tyr Gln Ser Lys Asn
305        310

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<210> SEQ ID NO 89
<211> LENGTH: 1104
<212> TYPE: DNA
<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 89

atgaaaaagt ttattgtttc agtgttttta ttgggatggc ttccgttata tgctgccggt    60
tcttctttgc aggaacaga agaagtcgct gtggcaaatg aatgagtga agccggttcg    120
gtcgatacag atcgtttcaa tgttcattg caaaaagaaa agaagcgcca gtcacctcgt    180
gaaaaacaaa aaagagtgat gaatacagag aaaacaggtg ttgcatcat cgcaccggc    240
ggttatgttc cggattcggg ccttcagcgg gcaatcggtg tgctgaagtc caggggatat    300
gaggttttca attatgttga tccccaaaaa cgtcatgaac gttttgcggc aaatgatgaa    360
gagcgaagcc gtcagattat ggaagcggca acgaatccgg atgtgaagat tgtgattgca    420
ttgccccggg gctacgggac gacgcggcct ttgcacgac ttgattttgc caaactggcc    480
aaaagcggaa aactctttgt cggtcacagc gatttcacgg ttttcgaaat ggcattgtta    540
aagcatggag ccgtcagttt ttccggccc atgattcaaa gcgattttac gcgaggtgat    600
ctgagcgcct ttacctttaa tcatttcgat gaaacctga catcgccgga aacatcggtc    660
aagtgggttt ccaagacaa tcccgatgtc gatgtcgagg gaacgctgtg gggcggaac    720
ctgacaatgc tggctcatat ggccgggact ccttgatgac cggacatttc cggtggcatt    780
ctgtttgtgg aagacattca cgaacatccc tatcgctcgc agcgtatgct gctccagctg    840
gatgaatcag gtatcctcaa gaaacagaaa gctctcgtgt tgggacattt ttccgaatc    900
aaactttccg actacgataa tggtacgat ttcaacgcca tgctttcctg gctccgttcg    960
cgtctgtcga taccctgtgt gaccggtttg ccttcgggc atacgaaaga caaagtcact   1020
ttgcctgtcg ggggcagagc gcatttgatg tccaaggcag gcaagattca actcgatatt   1080
ggggattatc caacggtaag gtaa                                           1104

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<210> SEQ ID NO 90
<211> LENGTH: 367
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 90

Met Lys Lys Phe Ile Val Ser Val Phe Leu Leu Gly Trp Leu Pro Leu
 1          5          10         15
Tyr Ala Ala Gly Ser Ser Leu Gln Glu Thr Glu Glu Val Ala Val Ala
 20         25         30
Asn Glu Met Ser Glu Ala Gly Ser Val Asp Thr Asp Arg Phe Asn Val
 35         40         45
Pro Leu Gln Lys Glu Lys Lys Arg Gln Ser Pro Arg Glu Lys Gln Lys
 50         55         60
Arg Val Met Asn Thr Glu Lys Thr Gly Ile Ala Ile Ile Ala Pro Gly
 65         70         75         80
Gly Tyr Val Pro Asp Ser Asp Leu Gln Arg Ala Ile Gly Val Leu Lys
 85         90         95
Ser Arg Gly Tyr Glu Val Phe Asn Tyr Val Asp Pro Gln Lys Arg His
100        105        110

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Glu Arg Phe Ala Ala Asn Asp Glu Glu Arg Ser Arg Gln Ile Met Glu
 115 120 125
 Ala Ala Thr Asn Pro Asp Val Lys Ile Val Ile Ala Leu Arg Gly Gly
 130 135 140
 Tyr Gly Thr Thr Arg Leu Leu His Asp Leu Asp Phe Ala Lys Leu Ala
 145 150 155 160
 Lys Ser Gly Lys Leu Phe Val Gly His Ser Asp Phe Thr Val Phe Glu
 165 170 175
 Met Ala Leu Leu Lys His Gly Ala Val Ser Phe Ser Gly Pro Met Ile
 180 185 190
 Gln Ser Asp Phe Thr Arg Gly Asp Leu Ser Ala Phe Thr Leu Asn His
 195 200 205
 Phe Asp Glu Thr Met Thr Ser Pro Glu Thr Ser Val Lys Trp Val Ser
 210 215 220
 Lys Asp Asn Pro Asp Val Asp Val Glu Gly Thr Leu Trp Gly Gly Asn
 225 230 235 240
 Leu Thr Met Leu Ala His Met Ala Gly Thr Pro Trp Met Pro Asp Ile
 245 250 255
 Ser Gly Gly Ile Leu Phe Val Glu Asp Ile His Glu His Pro Tyr Arg
 260 265 270
 Val Glu Arg Met Leu Leu Gln Leu Asp Glu Ser Gly Ile Leu Lys Lys
 275 280 285
 Gln Lys Ala Leu Val Leu Gly His Phe Ser Glu Phe Lys Leu Ser Asp
 290 295 300
 Tyr Asp Asn Gly Tyr Asp Phe Asn Ala Met Leu Ser Trp Leu Arg Ser
 305 310 315 320
 Arg Leu Ser Ile Pro Val Val Thr Gly Leu Pro Phe Gly His Thr Lys
 325 330 335
 Asp Lys Val Thr Leu Pro Val Gly Gly Arg Ala His Leu Met Ser Lys
 340 345 350
 Ala Gly Lys Ile Gln Leu Asp Ile Gly Asp Tyr Pro Thr Val Arg
 355 360 365

<210> SEQ ID NO 91

<211> LENGTH: 948

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 91

```

atgagtgaaa acggctggga acgtgaagtt ctcgaaaaac tgatgctgca aaccctgaaa   60
gagcaacggg cacgccgccc gtggggatc tttttcaagc tgacgacgat cattctggtt   120
atTTTTgtca ttttctcgat caagagcctt tctttttcca gcaaggaaac cgttcccgtc   180
cagaaacata cggcgatggg cgaatccgc gggacgatcg attcctccgg caattcgtcg   240
gcagccaata tcacaaaggc gctggacaag gcctatgacg agccactggc aacaggcggt   300
atcctgaaaa tcaacagccc gggcggagc cctgttcagg caggcatgat ttatgatgaa   360
atccggcggt tgcgtgaggt tcaccccgac aagcctcttt acgtcgtcgt tgaagaatta   420
tgcgcttccg gcggttatta tgttgccgca gcggcagaca agatctttgt cgacaaggcc   480
agtctggteg gttccatcgg tgtcatgatc aacggtttcg gagtcaccgg cctgatgcag   540
aaactcggcg tcgagcggc cctggtgacg gccggggaat acaagggttt tctcgaccgc   600

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ttttctcccc aaacgccaca gcaggtccag tatgcccaat ccatgctgaa ccagattcac    660
cagcaattca tcgaagtgt ccgtcagggc agggcgaca ggctgaaaga aaacagggaa    720
atctatagcg gtctggtttt tctcggtcct gaagcgatca aaatggggct ggctgacgaa    780
ctgggtacgg tcgaaagtgt tgcccgtgac gtgatcgggg aacctgtcat cgtcgattac    840
accgaacagg aaagactgtc tgaccgtttc ctgaagaaat tcggtgccct cgtcggttat    900
ggagcggttc aggcgggtct ggatatcaat cccgttcct tgcattga                948

```

<210> SEQ ID NO 92

<211> LENGTH: 315

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 92

```

Met Ser Glu Asn Gly Trp Glu Arg Glu Val Leu Glu Lys Leu Met Leu
1          5          10          15
Gln Thr Leu Lys Glu Gln Arg Ala Arg Arg Arg Trp Gly Ile Phe Phe
20          25          30
Lys Leu Thr Thr Ile Ile Leu Val Ile Phe Val Ile Phe Ser Ile Lys
35          40          45
Ser Leu Ser Phe Ser Ser Lys Glu Thr Val Pro Val Gln Lys His Thr
50          55          60
Ala Met Val Glu Ile Arg Gly Thr Ile Asp Ser Ser Gly Asn Ser Ser
65          70          75          80
Ala Ala Asn Ile Ile Lys Ala Leu Asp Lys Ala Tyr Asp Glu Pro Leu
85          90          95
Ala Thr Gly Val Ile Leu Lys Ile Asn Ser Pro Gly Gly Ser Pro Val
100         105         110
Gln Ala Gly Met Ile Tyr Asp Glu Ile Arg Arg Leu Arg Glu Val His
115         120         125
Pro Asp Lys Pro Leu Tyr Val Val Val Glu Glu Leu Cys Ala Ser Gly
130         135         140
Gly Tyr Tyr Val Ala Ala Ala Ala Asp Lys Ile Phe Val Asp Lys Ala
145         150         155         160
Ser Leu Val Gly Ser Ile Gly Val Met Ile Asn Gly Phe Gly Val Thr
165         170         175
Gly Leu Met Gln Lys Leu Gly Val Glu Arg Arg Leu Leu Thr Ala Gly
180         185         190
Glu Tyr Lys Gly Phe Leu Asp Pro Phe Ser Pro Gln Thr Pro Gln Gln
195         200         205
Val Gln Tyr Ala Gln Ser Met Leu Asn Gln Ile His Gln Gln Phe Ile
210         215         220
Glu Val Val Arg Gln Gly Arg Gly Asp Arg Leu Lys Glu Asn Arg Glu
225         230         235         240
Ile Tyr Ser Gly Leu Val Phe Leu Gly Pro Glu Ala Ile Lys Met Gly
245         250         255
Leu Ala Asp Glu Leu Gly Thr Val Glu Ser Val Ala Arg Asp Val Ile
260         265         270
Gly Glu Pro Val Ile Val Asp Tyr Thr Glu Gln Glu Arg Leu Ser Asp
275         280         285
Arg Phe Leu Lys Lys Phe Gly Ala Ser Val Gly Tyr Gly Ala Val Gln
290         295         300

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Met	Lys	Thr	Asp	Leu	Pro	Gly	Lys	Asn	Ala	Ile	Pro	Lys	Asn	Arg	Met
1				5					10					15	
Thr	Ile	Asn	Arg	Leu	Phe	Ala	Tyr	Thr	Arg	Ile	Pro	Arg	Ile	Arg	Pro
		20						25					30		
Ala	Leu	Thr	Met	Ala	Leu	Cys	Ala	Ala	Thr	Leu	Ala	Gly	Cys	Ala	Val
		35					40					45			
Gly	Pro	Asp	Tyr	Val	Gln	Pro	Pro	Ile	Asp	Val	Pro	Ala	Thr	Tyr	Lys
	50					55					60				
Glu	Asp	Ser	Leu	Trp	Lys	Thr	Ala	Lys	Pro	Ser	Asp	Glu	Tyr	Pro	Arg
65					70					75					80
Gly	Ala	Trp	Trp	Ser	Val	Phe	Lys	Asp	Pro	Glu	Leu	Asp	Arg	Leu	Met
				85					90					95	
Val	Leu	Leu	Asn	Lys	Gln	Asn	Leu	Thr	Ile	Ala	Gln	Ala	Glu	Ala	Gln
			100					105					110		
Tyr	Arg	Gln	Ala	Gln	Ala	Leu	Leu	Arg	Gln	Ala	Gln	Ser	Ser	Leu	Phe
		115						120					125		
Pro	Ser	Leu	Ser	Leu	Asp	Ala	Ser	Arg	Thr	Arg	Gly	Phe	Gln	Ser	Asn
	130					135					140				
Thr	Ser	Thr	Ser	Ala	Ser	Thr	Gln	Asn	Ala	Phe	Val	Gly	Asn	Leu	Ser
145					150					155					160
Trp	Glu	Val	Asp	Ile	Trp	Gly	Gly	Val	Arg	Arg	Asn	Val	Glu	Ala	Gly
				165					170					175	
Glu	Ala	Gly	Arg	Gln	Ala	Ser	Ala	Ala	Gln	Leu	Glu	Ala	Ile	Lys	Leu
			180					185					190		
Ser	Ser	Gln	Ala	Gln	Met	Ala	Thr	Ala	Tyr	Leu	Gln	Leu	Val	Ile	Thr
		195					200					205			
Asp	Arg	Gln	Val	Lys	Gln	Leu	Glu	Glu	Ser	Glu	Thr	Leu	Leu	Ala	Glu
	210					215					220				
Ser	Leu	Arg	Leu	Thr	Lys	Asn	Gln	Phe	Ala	Val	Gly	Ile	Val	Ser	Asp
225					230					235					240
Ala	Asp	Val	Ala	Gln	Ala	Glu	Ser	Gln	Leu	Lys	Thr	Ala	Gln	Ala	Ala
				245					250					255	
Thr	Val	Asp	Met	Lys	Leu	Ala	Arg	Ser	Gln	Leu	Glu	His	Ser	Ile	Ala
			260					265					270		
Val	Ser	Ile	Gly	Gln	Ala	Pro	Ser	Thr	Leu	Thr	Ile	Asp	Met	Ala	Lys
		275					280					285			
Ala	Asp	Pro	Tyr	Leu	Pro	Gln	Ile	Pro	Ala	Gly	Leu	Pro	Ser	Ala	Leu
	290					295					300				
Leu	Gln	Arg	Arg	Pro	Asp	Val	Ala	Ser	Ala	Glu	Arg	Lys	Val	Ala	Gln
305					310					315					320
Ala	Asn	Ala	Leu	Ile	Gly	Val	Ala	Lys	Ser	Ala	Phe	Phe	Pro	Ala	Leu
				325					330					335	
Ser	Ile	Ser	Ala	Ser	Gly	Gly	Phe	Arg	Asn	Ser	Ser	Phe	Ala	Asp	Leu
			340					345					350		
Phe	Thr	Val	Pro	Asn	Arg	Ile	Trp	Ser	Ile	Gly	Pro	Gln	Ile	Ala	Leu
			355				360					365			
Ser	Ile	Phe	Asp	Ala	Gly	Leu	Arg	Ser	Ala	Gln	Thr	Asp	Gln	Ala	Ile
	370				375						380				
Ala	Val	Tyr	Asp	Glu	Thr	Val	Ala	Ala	Tyr	Arg	Gln	Lys	Val	Leu	Ile
385					390					395					400
Ala	Phe	Gln	Glu	Val	Glu	Asp	Asn	Leu	Ala	Ala	Gln	Ala	Met	Leu	Gly

-continued

	405		410		415										
Asp	Ala	Ser	Asp	Met	Gln	Thr	Ala	Ala	Leu	Asn	Ala	Ala	Lys	Arg	Ala
	420								425					430	
Glu	Thr	Ile	Thr	Met	Asn	Gln	Tyr	Lys	Ala	Gly	Val	Val	Ser	Tyr	Ile
	435						440						445		
Asn	Val	Leu	Ile	Ala	Gln	Asn	Thr	Arg	Ile	Ala	Ala	Glu	Asn	Thr	Leu
	450					455						460			
Tyr	Asn	Val	Lys	Lys	Arg	Gln	Phe	Thr	Ser	Ser	Val	Ala	Leu	Ile	Ala
465					470					475					480
Ala	Ile	Gly	Gly	Gln	Trp	Glu	Thr	Pro	Lys	Gln	Thr	Ala	Lys	Tyr	Arg
				485					490						495
His	Gly	Ile	Thr	Gly	Lys	Asn	Gly	Ala	Glu	Asn	Gly	Asn	Glu	Lys	Leu
			500					505						510	
Gly	Thr	Phe	Cys	Tyr	Arg	Ile	Asn	Asp	Ser	Ser	Phe	Gly	Thr	Asp	Ser
		515					520							525	
Val	Thr	Tyr	Arg	Gln	Ile	Phe	Lys	Ser	Asp	Lys	Arg	Glu	Asn	Thr	Trp
	530					535						540			
Ile															
545															

<210> SEQ ID NO 95
 <211> LENGTH: 1476
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 95

atgaaagtca gaaggacaaa gaagatgatt aagaaattaa ccctcgtttt actggctgcc	60
ggtgtccttga gcagttgcag tatggcgccc aattacgtgc gaccggatgc cccgattgaa	120
agccagttcc cgggcaacgc ggatgacgct tctgcaaaaa cgcccgtgac gcaaatcgga	180
tggaatgaat ttttccatga accgcgattg aaagcgctga tcgctgccgc aattgaaaac	240
aatcgtgatt tgagagtagc agccttgcca attgaggaag cgcgtgccct gtatggcatc	300
cagtgggctg accgtctccc gaatttcgaa gctcagggag ccggaacccg tcaaagaacg	360
gtcggcacia cggcgggcat ggtaacacag ggcaattaca ctgttgccct gggctctggct	420
gccttcgagc tggacttctt cggaaggggtg aaaagcctgt cggacgctgc actggcagaa	480
tatctggcca cggaagaagc ccaacgtgcc gcctatatca gccttgtttc ggaagtggcc	540
aaaacctatc tgaccgaacg tgcacaagcg cgacagattg agttggcaaa agaactctac	600
gagtcttaca aacgcagcta tgagctgatg caaaagcgtt atgaagtcgg tgcactctcc	660
gcactggaat tgcgtcagta tgaaacgctg atgcaatcgg cactggtttc gctctccacc	720
ctgcaaagac aacgcgcca gacagaaaac gcaactgtcg tactgattgg tggaaagcag	780
atcaaggatc ttctgcggc gcacgacttg tcggaagacg acatcatgca ggacattccc	840
gccggattgc cgtccgatct gatcaacaac cggccggaca ttcgccagta tgaacaattg	900
ctcaagtcag cgaacgcaa tatcgcgca gccagagcag cctttttccc gcgtattacc	960
ctgacaggtt ttgccggaac agcaagcccg actttatccg gccttttcga tgccgggtcc	1020
ggtgcatgga catttatgcc gcaactgaca ctgccgattt tcgatgcagg ccgcaatate	1080
agcaatctgg atctggcaga ggctcgcaag aatatcgcg tggcccaata tgaaaaaacc	1140
gttcaggtcg ctttccgtga agtcgctgac gccctgatgg cacgcgactg gttgaaatgaa	1200

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caggtaagg ctcaggctgc tgttctgaaa tcagagacgg aacgcctgaa actgtccgaa 1260
gcgcggtaca acaacggat tgccagttcg ctggaagtat tcgattcaca acgccagcaa 1320
tttgcggcac agcagctgct ggttgacgca cgtctgttac gcctgatcaa tacagtggaa 1380
ctgtatcggt cactgggtgg cggcatcgtc gatgccaatg cgccgaaaac agccaaaaca 1440
actgctgaac cggtagcgcga aaatactgaa ggctga 1476

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<210> SEQ ID NO 96

<211> LENGTH: 491

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 96

```

Met Lys Val Arg Arg Thr Lys Lys Met Ile Lys Lys Leu Thr Leu Val
 1           5           10          15
Leu Leu Ala Ala Gly Val Leu Ser Ser Cys Ser Met Ala Pro Asn Tyr
          20           25           30
Val Arg Pro Asp Ala Pro Ile Glu Ser Gln Phe Pro Gly Asn Ala Asp
          35           40           45
Asp Ala Ser Ala Lys Thr Pro Val Thr Gln Ile Gly Trp Asn Glu Phe
 50           55           60
Phe His Glu Pro Arg Leu Lys Ala Leu Ile Ala Ala Ala Ile Glu Asn
 65           70           75           80
Asn Arg Asp Leu Arg Val Ala Ala Leu Arg Ile Glu Glu Ala Arg Ala
          85           90           95
Leu Tyr Gly Ile Gln Trp Ala Asp Arg Leu Pro Asn Phe Glu Ala Gln
          100          105          110
Gly Ala Gly Thr Arg Gln Arg Thr Val Gly Thr Thr Gly Gly Met Val
          115          120          125
Thr Gln Gly Asn Tyr Thr Val Gly Leu Gly Leu Ala Ala Phe Glu Leu
          130          135          140
Asp Phe Phe Gly Arg Val Lys Ser Leu Ser Asp Ala Ala Leu Ala Glu
          145          150          155          160
Tyr Leu Ala Thr Glu Glu Ala Gln Arg Ala Ala Tyr Ile Ser Leu Val
          165          170          175
Ser Glu Val Ala Lys Thr Tyr Leu Thr Glu Arg Ala Gln Ala Arg Gln
          180          185          190
Ile Glu Leu Ala Lys Glu Ser Tyr Glu Ser Tyr Lys Arg Ser Tyr Glu
          195          200          205
Leu Met Gln Lys Arg Tyr Glu Val Gly Ala Ser Ser Ala Leu Glu Leu
          210          215          220
Arg Gln Tyr Glu Thr Leu Met Gln Ser Ala Leu Val Ser Leu Ser Thr
          225          230          235          240
Leu Gln Arg Gln Arg Ala Gln Thr Glu Asn Ala Leu Val Val Leu Ile
          245          250          255
Gly Gly Lys Gln Ile Lys Asp Leu Pro Ala Ala His Asp Leu Ser Glu
          260          265          270
Asp Asp Ile Met Gln Asp Ile Pro Ala Gly Leu Pro Ser Asp Leu Ile
          275          280          285
Asn Asn Arg Pro Asp Ile Arg Gln Tyr Glu Gln Leu Leu Lys Ser Ala
          290          295          300

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Asn Ala Asn Ile Gly Ala Ala Arg Ala Ala Phe Phe Pro Arg Ile Thr
 305 310 315 320

Leu Thr Gly Phe Ala Gly Thr Ala Ser Pro Thr Leu Ser Gly Leu Phe
 325 330 335

Asp Ala Gly Ser Gly Ala Trp Thr Phe Met Pro Gln Leu Thr Leu Pro
 340 345 350

Ile Phe Asp Ala Gly Arg Asn Ile Ser Asn Leu Asp Leu Ala Glu Ala
 355 360 365

Arg Lys Asn Ile Ala Val Ala Gln Tyr Glu Lys Thr Val Gln Val Ala
 370 375 380

Phe Arg Glu Val Ala Asp Ala Leu Met Ala Arg Asp Trp Leu Asn Glu
 385 390 395 400

Gln Val Lys Ala Gln Ala Ala Val Leu Lys Ser Glu Thr Glu Arg Leu
 405 410 415

Lys Leu Ser Glu Ala Arg Tyr Asn Asn Gly Ile Ala Ser Ser Leu Glu
 420 425 430

Val Phe Asp Ser Gln Arg Gln Gln Phe Ala Ala Gln Gln Ser Leu Val
 435 440 445

Asp Ala Arg Leu Leu Arg Leu Ile Asn Thr Val Glu Leu Tyr Arg Ser
 450 455 460

Leu Gly Gly Gly Ile Val Asp Ala Asn Ala Pro Lys Thr Ala Lys Thr
 465 470 475 480

Thr Ala Glu Pro Val Ala Gln Asn Thr Glu Gly
 485 490

<210> SEQ ID NO 97
 <211> LENGTH: 602
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 97

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taacgtgagg aacagatcac atcgacaagc cttgccgatt gtctttccat ctcgatcaga    60
aaactgtcga caagcgcccg gaaatgacga taaaaaacga gagtcggcat ggcaatcagc    120
aaaccgaagc cggatttgta aagcgccaca gaaatcccgt gagccagttc cgcaggattg    180
gcccctccgg catttttgta accgaaaatc tcgatcatcc cgacaaccgt accgaacagg    240
cccatcaggg gagccagcgt ggcaatcgtc ccgatcgtcg tcagaaagcg ttccatctgg    300
agagcgacac tgcggccggg ttcttccatc gcttccttca tattgtcgcg ggaaacatcg    360
gaatgctgca atccggctgc cagcaactgtc cccaaaggcg aaccttttc cagtttgctg    420
attgtgcgcy catccacagc accttcacga tactcactga tgacctgacg cagaagatgt    480
ggcggcaaaa tctttcggcg tctcagataa atgagacggt cgataatcag ggccaatgca    540
ataacagagg cgatcagcaa aaaccagatc ggccatcctg ccgcttgaat gattgaaac    600
ac                                                                                   602
    
```

<210> SEQ ID NO 98
 <211> LENGTH: 659
 <212> TYPE: DNA
 <213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 98

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tgtccaaaca acaatttaac ctgatcgttt tcgattggga cggcaccctg atggacagta    60
    
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cgctccgceat	cgctcaagagc	cttcaggatt	cggcagccga	tctgggcttg	cccacaccag	120
acaaaaagaa	agcggcgcat	gtgatcggcc	tggccttcg	agaagcctg	gaaaccgtcc	180
tgccggacgt	cagcccgaa	tactatccga	aactcatcga	gcgctatcgt	accattttcc	240
tgaaaaacag	cgtatcgctt	tccctgttcg	acgggtgcag	ggaaatgctc	gacgacctga	300
agaatcagga	ttattttctg	gcagtgcca	cgggaaaaag	cggggccggt	ctgaaccgcg	360
ccatcgacga	gggtggcttg	cagggatatt	tcgatgccag	cgcacggct	gatgaaacgc	420
attccaaacc	tcaccggcc	atgttgctgg	aactgaccga	acagttgggc	gaaccgatga	480
aacgcacggt	catgatcggg	gataccaccc	atgatttgc	gatggcaaac	aatgccgggtg	540
cctcgggcat	cgctgtacaa	tacgggtcac	attcacccaa	agaactgcaa	ttgatgcatc	600
cgatgtattc	agccgatacg	gtcgaagaac	tgcatcgtg	gctgaatgaa	aatgcctga	659

<210> SEQ ID NO 99

<211> LENGTH: 2747

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 99

tatcgttcaa	gccggttoga	tcaagcagt	ttttcaaatt	cgcgcctgt	caggggttca	60
gagaaaagat	atccctgacc	ataattgcat	cggcaccgg	ccaacaacaa	tctctgtcca	120
tcgtcctgga	caccttggc	aatcactttc	atgtccagtt	tgtgcgccat	aacgataatc	180
gcctcacaca	aggccatata	cttgggttca	ttttcgatat	ggcgaacgaa	agcctgatcg	240
attttcagat	agtcgatcgg	aaatttttc	agataagaca	atgaagaata	tcccgteccg	300
aaatcgtcca	gtgccacttc	cattccatat	tgatgaaatt	cgcaccgccc	atcaagcacc	360
tgggtattcg	aaacaatcaa	aacaccttct	gttattttca	tcacgatgct	gctgccaggt	420
acattcagtt	tttctagcac	ttcgaaccag	tcgctgtaat	cattottggt	ctggaacacg	480
ataggcgaga	tattgaaact	gatttgcaag	tcaggaaga	gatgttctct	ccatcggctg	540
agctccctgg	caacctcccc	gaaagcccag	tccgagatcg	tcaaaatctg	accggtttct	600
tccgcaatga	aaatgaattc	tgtgggactg	ataatgccac	gaaccggatg	gtgccagcgg	660
atcagtgett	cggtttctg	gatgcggcca	gtctcaatct	cgacaatcgg	ctgatacagc	720
atttcaaatt	gatgttcatt	tatagctata	cgcctgcat	tggccagcct	tatccggttt	780
tgcgcagccg	cctgcatgga	tgccgtaaag	taactcatcc	ggttacgtcc	ggccagtttg	840
gcggcatata	aggcctgatc	ggcattttct	atgagtgtat	cgatatccgt	gccgtcatcc	900
ggataaacgg	taatgccgat	acttctctgc	acatacgcaa	tttcgctacc	caactggtaa	960
ggctgggcca	gggaacgcag	gatgctgtgc	gcaatttttt	ccattgtcgg	caaatcggac	1020
aactctgtca	gaataatggc	aaactcatcg	ccaccagac	gtgacacggg	atcgaattca	1080
cggacgcaat	ccttcagacg	tttcgagacg	tccagtaaca	gggtgatcgcc	cgcategtgc	1140
ccaaggggat	cgttgatata	cttgaaatta	tccagatcga	gaaacagcag	cgcaggtttt	1200
tttccggact	ggttgacctt	gactagttcc	tgtttcatcc	tgtccatcag	catgcgacga	1260
ttcggcaaat	tcgtcaggaa	atcgaaatc	gcctgacgcc	atatcatgct	ttcgccttct	1320
ttttctccg	tgaaatcatt	ggagagttca	acgtaataac	gaacggatct	gtcttcggtg	1380
taaaccacat	tgattgtcag	ccatgatgca	tacgattcgc	cattottgcg	cgggttcag	1440

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agttccccac gccagtagcc tttctcatcc aatgcatccc acatcgctt gtaaaatgcc 1500
ttgtcctggc gtcccgaact caacagtttc ggatccctgc cgatcacttc ggcagcggta 1560
tagcctgtaa ttttttcaaa ggcgggattg actgcaagaa tatgattgtc ggcattactg 1620
atcaaaatcg cttcactgga gttacggtag accagtgtct ccagttccag tttttcctcg 1680
tcttcctttc ttcgcgtaat atcccgtccg attcccagaa caccaccag ttcgcccgtcc 1740
ggcgtttctga cagggtgcctt gatcacatcg aaaaaccccc tgttgccaga caatgcataa 1800
gtgccccact gcttgatcac cagtggtttg ccttccttca gaatacgttt gtcctgttct 1860
tcaaaaagct cggcttcgtc atggctgaat atacgaaat cggttttacc gataaatatgc 1920
tcgtcagaaa gcccgacata atccttgtaa gcttgattac aggccagata aacgcccga 1980
gaatTTTTca tccataccag atcgggaata ttttcaaaaa cgtttttcag tcgcccctcg 2040
ctttgggtca gaacgacatt gacctgttc agatcgctcg ttctgatctc aaccagtttc 2100
tccagattga caagcaactg gttccggatt tcttgcaagc ggaccataaa tagaatggcc 2160
aaaaagaaca gcgccatgag ggcgattgaa atcccgatga ctttgggtgcc ccaataaaaa 2220
aggctcgcac tatatgcgcc cagcggcaaa agcgtgatga ctttccattt ataggcactg 2280
acatcacggg gacctgttc tccgaaaaaa atttcgttac tgcctttact ggttttggtta 2340
tcggccgaga gagtgaaag cgttttgaaac gtccataggc cttcccgggt aatgaactgg 2400
ccttcttctc gatttttcat ccgctcccag gcagcaggat atcgatgagc cataactcaag 2460
tcatttttat tgaacatgaa tccgaattca tcgttgaaa cgggtccttt cagccagaat 2520
ccgttctgat tgacgagcca ggcagaattg ccacgcacgc tcgtcaactg ctggaaccgg 2580
ccgatcattg aagtggcata gtagtcaac agaagaatgc cctttttctc cccttttttg 2640
ttgaagagag gcacccgaa acggatcgtc gggacataag gctcctggat atggtcactc 2700
tcaatattca ggtcaagagg tgaacatag acttcccctt ctttcaa 2747

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<210> SEQ ID NO 100

<211> LENGTH: 659

<212> TYPE: DNA

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 100

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tgtccaaaca acaatttaac ctgatcgtt tcgattggga cggcaccctg atggacagta 60
cgtccgccat cgtcaagagc cttcaggatt cggcagccga tctgggcttg cccacaccag 120
acaaaaagaa agcggcgcat gtgatcggcc tcggccttcg agaagcgtcg gaaaccgtcc 180
tgccggacgt cagcccggaa tactatccga aactcactga gcgctatcgt acccatttcc 240
tgaaaaacag cgtatcgtt tccctgttcg acggtgtcag gaaatgctc gacgacctga 300
agaatcagga ttattttctg gcagtgcca ccgaaaaag ccgggcccgt ctgaaccgcg 360
ccatcgacga ggtgggcttg caggatatt tcgatgccag ccgcacggct gatgaaacgc 420
attccaaacc tcatccggcc atgttgctgg aactgaccga acagttgggc gaaccgatga 480
aacgcacggc catgatcggc gataccacc atgatttgct gatggcaaac aatgccggtg 540
cctcgggcac cgtgtacaa tacggtgcac attcacccaa agaactgcaa ttgatgcac 600
cgatgtattc agccgatcgc gtcgaagaac tgcacgctg gctgaatgaa aatgcctga 659

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<210> SEQ ID NO 101

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<211> LENGTH: 190
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 101
Leu Ile Val Phe Asp Trp Asp Gly Thr Leu Met Asp Ser Thr Ser Ala
1           5           10          15
Ile Val Lys Ser Leu Gln Asp Ser Ala Ala Asp Leu Gly Leu Pro Thr
20          25
Pro Asp Lys Lys Lys Ala Ala His Val Ile Gly Leu Gly Leu Arg Glu
35          40          45
Ala Leu Glu Thr Val Leu Pro Asp Val Ser Pro Glu Tyr Tyr Pro Lys
50          55          60
Leu Ile Glu Arg Tyr Arg Thr His Phe Leu Lys Asn Ser Val Ser Leu
65          70          75          80
Ser Leu Phe Asp Gly Val Arg Glu Met Leu Asp Asp Leu Lys Asn Gln
85          90          95
Asp Tyr Phe Leu Ala Val Ala Thr Gly Lys Ser Arg Ala Gly Leu Asn
100         105         110
Arg Ala Ile Asp Glu Val Gly Leu Gln Gly Tyr Phe Asp Ala Ser Arg
115         120         125
Thr Ala Asp Glu Thr His Ser Lys Pro His Pro Ala Met Leu Leu Glu
130         135         140
Leu Thr Glu Gln Leu Gly Glu Pro Met Lys Arg Thr Val Met Ile Gly
145         150         155         160
Asp Thr Thr His Asp Leu Leu Met Ala Asn Asn Ala Gly Ala Ser Gly
165         170         175
Ile Ala Val Gln Tyr Gly Ala His Ser Pro Lys Glu Leu Gln
180         185         190

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<210> SEQ ID NO 102
<211> LENGTH: 112
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 102
Phe Glu Asn Ile Pro Asp Leu Val Trp Met Lys Asn Ser Ala Gly Val
1           5           10          15
Tyr Leu Ala Cys Asn Gln Ala Tyr Lys Asp Tyr Val Gly Leu Ser Asp
20          25          30
Glu His Ile Ile Gly Lys Thr Asp Phe Asp Ile Phe Ser His Asp Glu
35          40          45
Ala Glu Leu Phe Glu Glu Gln Asp Lys Arg Ile Leu Lys Glu Gly Lys
50          55          60
Pro Leu Val Ile Lys Gln Trp Gly Thr Tyr Ala Leu Ser Gly Asn Arg
65          70          75          80
Val Phe Phe Asp Val Ile Lys Ala Pro Val Arg Thr Pro Asp Gly Glu
85          90          95
Leu Val Gly Val Leu Gly Ile Gly Arg Asp Ile Thr Arg Arg Lys Glu
100         105         110

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<210> SEQ ID NO 103
<211> LENGTH: 190
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

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-continued

<400> SEQUENCE: 103

Lys Ala Ala Lys Ser Gly Asn Ala Glu Ala Gln Tyr Leu Phe Gly Met
 1 5 10 15
 Leu Val Tyr Asp Gly Arg Gly Val Gln Gln Asp Asn Cys Val Ala Met
 20 25 30
 Leu Trp Trp Met Lys Ala Ala Glu Gln Asn His Ala Lys Ala Leu Val
 35 40 45
 Met Leu Gly Asn Leu His Arg Lys Gly Gln Cys Ile Ala Glu Asn Tyr
 50 55 60
 Pro Lys Ala Ile Ala Tyr Trp Lys Arg Ala Ala Val Gln Asn Asn Val
 65 70 75 80
 Trp Ala Tyr His Asn Leu Gly Thr Ala Tyr Tyr Asp Gly Ile Gly Val
 85 90 95
 Asp Lys Asn Pro His Glu Ala Val Arg Trp Trp Lys Lys Ala Ala Glu
 100 105 110
 Leu Gly Phe Pro Glu Ser Gln Asn Asn Leu Gly Ala Leu Tyr Asn Asp
 115 120 125
 Gly Asn Gly Val Asp Arg Asp Tyr Gln Glu Ala Val Phe Trp Tyr Arg
 130 135 140
 Lys Ser Ala Leu Gln Gly Asp Glu Leu Gly Gln Tyr Asn Leu Gly Val
 145 150 155 160
 Ala Tyr Tyr Tyr Gly Arg Gly Ile Lys Lys Asp Phe Ser Glu Ala Val
 165 170 175
 Ser Trp Tyr Lys Lys Ser Ala Glu Gln Asp Tyr Ala Gln Ala
 180 185 190

<210> SEQ ID NO 104

<211> LENGTH: 249

<212> TYPE: PRT

<213> ORGANISM: Oxalobacter formigenes

<400> SEQUENCE: 104

Leu Gly Thr Ala Tyr Tyr Asp Gly Ile Gly Val Asp Lys Asn Pro His
 1 5 10 15
 Glu Ala Val Arg Trp Trp Lys Lys Ala Ala Glu Leu Gly Phe Pro Glu
 20 25 30
 Ser Gln Asn Asn Leu Gly Ala Leu Tyr Asn Asp Gly Asn Gly Val Asp
 35 40 45
 Arg Asp Tyr Gln Glu Ala Val Phe Trp Tyr Arg Lys Ser Ala Leu Gln
 50 55 60
 Gly Asp Glu Leu Gly Gln Tyr Asn Leu Gly Val Ala Tyr Tyr Tyr Gly
 65 70 75 80
 Arg Gly Ile Lys Lys Asp Phe Ser Glu Ala Val Ser Trp Tyr Lys Lys
 85 90 95
 Ser Ala Glu Gln Asp Tyr Ala Gln Ala Gln His Asn Leu Gly Val Thr
 100 105 110
 Tyr Tyr Glu Gly Glu Gly Ile Lys Lys Asp Tyr Ala Lys Ala Val Tyr
 115 120 125
 Trp Trp Lys Lys Ala Ala Glu Gln Gly Ile Pro Gln Ser Gln Tyr Asn
 130 135 140
 Leu Gly Ile Ala Tyr Glu Glu Gly Trp Gly Ala Glu Lys Asn Pro Glu
 145 150 155 160

-continued

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Asn Ala Val Phe Trp Tyr Arg Lys Ala Ala Glu Gln Gly His Ala Asp
      165                               170                               175
Ala Gln Asn Arg Leu Gly Ile Ala Tyr Arg Tyr Gly Thr Gly Val Arg
      180                               185                               190
Lys Asn Pro Ala Leu Ser Val Lys Trp Leu Glu Lys Ala Ala Lys Gln
      195                               200                               205
Gly Leu Ala Arg Ala Gln Phe Asn Leu Gly Lys Thr Phe Tyr Ile Gly
      210                               215                               220
Ala Gly Ile Asn Lys Asn Thr Asp Lys Ala Val Tyr Trp Phe Ile Lys
      225                               230                               235                               240
Ala Ala Asn Gln Gly Phe Thr Glu Ala
      245

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<210> SEQ ID NO 105
<211> LENGTH: 172
<212> TYPE: PRT
<213> ORGANISM: Oxalobacter formigenes

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<400> SEQUENCE: 105

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Tyr Arg Lys Ala Ala Glu Gln Gly His Ala Asp Ala Gln Asn Arg Leu
 1      5      10      15
Gly Ile Ala Tyr Arg Tyr Gly Thr Gly Val Arg Lys Asn Pro Ala Leu
 20     25     30
Ser Val Lys Trp Leu Glu Lys Ala Ala Lys Gln Gly Leu Ala Arg Ala
 35     40     45
Gln Phe Asn Leu Gly Lys Thr Phe Tyr Ile Gly Ala Gly Ile Asn Lys
 50     55     60
Asn Thr Asp Lys Ala Val Tyr Trp Phe Ile Lys Ala Ala Asn Gln Gly
 65     70     75     80
Phe Thr Glu Ala Gln Ala Tyr Ile Gly Met Ile Tyr Phe Lys Gly Lys
 85     90     95
Tyr Val Ala Lys Asn Glu Lys Lys Gly Phe Tyr Trp Leu Lys Lys Ala
100    105    110
Ala Glu Lys Asp Ser Ala Lys Ala Gln Ala Phe Leu Gly Ala Leu Tyr
115    120    125
Ile Ala Gly Asn Glu Val Lys Pro Asn Ile Lys Glu Gly Val Ala Leu
130    135    140
Thr Lys Lys Ala Ala Leu Gln Gly Asn Tyr Glu Ala Gln Thr Leu Leu
145    150    155    160
Gly Phe Cys Tyr Glu Asn Gly Leu Glu Val Lys Lys
165    170

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1. A method comprising administering to the subject one or more *Oxalobacter formigenes* (Of)-derived factors, and/or bioactive variants and/or fragments thereof, that result in stimulation of oxalate transport.

2. (canceled)

3. The method of claim 1, wherein the one or more Of-derived factors are administered by administering Of conditioned media (CM) to the subject.

4. The method of claim 3, wherein the Of CM is fractionated, purified, and/or otherwise processed prior to administration.

5.-6. (canceled)

7. The method of claim 1, wherein one or more of the Of-derived factors are Sell1-like repeat (SLR) proteins or bioactive variants and/or fragments of Sell1 proteins.

8. (canceled)

9. The method of claim 7, wherein one or more of the Of-derived factors comprises 70% or greater sequence identity to all or a portion of an Of-derived Sell1 protein.

10. The method of claim 7, wherein the Sell1 protein or proteins are selected from the group consisting of Sell1, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, and MerG.

11. The method of claim 10, wherein the Of-derived factor is a peptide fragment of one of Sell1, Hrd3, Chs4, Nif1, PodJ,

ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, and MerG, and/or a bioactive peptide having at least 70% sequence identity to a fragment thereof.

12. The method of claim 7, wherein the Of-derived factor comprises SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, and SEQ ID NO: 102.

13. The method of claim 7, wherein the Of-derived factor comprises a bioactive variant having 70% sequence identity to SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, and SEQ ID NO: 102.

14. (canceled)

15. The method of claim 1, wherein administration comprises rectal administration, oral administration, and/or injection.

16.-17. (canceled)

18. A pharmaceutical composition comprising one or more *Oxalobacter formigenes* (O)-derived factors, and/or bioactive variants and/or fragments thereof.

19. The pharmaceutical composition of claim 1, comprising Of conditioned media (CM).

20. The pharmaceutical composition of claim 3, wherein the Of CM is fractionated, purified, and/or otherwise processed.

21.-22. (canceled)

23. The pharmaceutical composition of claim 20, wherein one or more of the Of-derived factors are selected from polypeptides comprising all or a portion of SEQ ID NOS; 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, 36,

38, 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, 60, 62, 64, 66, 68, 70, 72, 74, 76, 78, 80, 82, 84, 86, 88, 90, 92, 94, 96, 101, or 102.

24. The pharmaceutical composition of claim 20, wherein one or more of the Of-derived factors are Sell-like repeat (SLR) proteins or bioactive variants and/or fragments of Sell proteins.

25. (canceled)

26. The pharmaceutical composition of claim 24, wherein one or more of the Of-derived factors comprises 70% or greater sequence identity to all or a portion of an Of-derived Sell protein.

27. The pharmaceutical composition of claim 24, wherein the Sell protein or proteins are selected from the group consisting of Sell, Hrd3, Chs4, Nif1, PodJ, ExoR, AlgK, HcpA, Hsp12, EnhC, LpnE, MotX, and MerG.

28. (canceled)

29. The pharmaceutical composition of claim 24, wherein the Of-derived factor comprises SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, or SEQ ID NO: 102.

30. The pharmaceutical composition of claim 24, wherein the Of-derived factor comprises a bioactive variant having 70% or more sequence identity to SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 90, SEQ ID NO: 92, SEQ ID NO: 94, SEQ ID NO: 96, SEQ ID NO: 101, or SEQ ID NO: 102.

31. (canceled)

32. The pharmaceutical composition of claim 18, wherein the pharmaceutical composition is formulated for rectal administration, oral administration, and/or injection.

33.-34. (canceled)

* * * * *