

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
16 February 2006 (16.02.2006)

PCT

(10) International Publication Number
WO 2006/017688 A2

(51) International Patent Classification:
A61K 38/26 (2006.01)

(21) International Application Number:
PCT/US2005/027800

(22) International Filing Date: 3 August 2005 (03.08.2005)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/598,031 3 August 2004 (03.08.2004) US

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(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AE, AG, AL, AM,
AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,
KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ,
OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL,
SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC,
VN, YU, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every
kind of regional protection available): ARIPO (BW, GH,
GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,
ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI,
FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT,
RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA,
GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— without international search report and to be republished
upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: COMBINATION THERAPY USING TRANSFERRIN FUSION PROTEINS COMPRISING GLP-1

(57) Abstract: The present invention provides combination therapy comprising transferrin fusion protein and DPP-IV inhibitors and/or neutral endopeptidase (NEP) inhibitors. The transferrin fusion protein comprises therapeutic polypeptides or peptides useful in the treatment of diseases such as diabetes.

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TITLE: Combination Therapy Using Transferrin Fusion Proteins Comprising GLP-1

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RELATED APPLICATION

[0001] This application claims the benefit of U.S. Provisional Application 60/598,031, filed August 3, 2004, which is herein incorporated by reference in its entirety.

[0002] This application is related to a Continuation-in-Part of PCT/US03/26818, filed August 28, 2003, which is a Continuation-in-Part of U.S. Application No. 10/378,094, filed March 4, 2003, both of which are herein incorporated by reference in their entirety.

FIELD OF THE INVENTION

[0003] The present invention is related to transferrin fusion proteins comprising insulinotropic peptides with extended effective therapeutic *in vivo* half life. The present invention also relates to combination therapies using DPP-IV inhibitors and/or neural endopeptidase inhibitors (NEP) and insulinotropic peptides.

BACKGROUND OF THE INVENTION

Proteases

[0004] Proteolytic enzymes play an important role in regulating physiological processes such as cell proliferation, differentiation, and signaling processes by regulating protein turnover and processing. Proteolytic enzyme controls the levels of important structural proteins, enzymes, and regulatory proteins through proteolytic degradation. Uncontrolled proteolytic enzyme activity, either increased or decreased, has been implicated in a variety of disease conditions including inflammation, cancer, arteriosclerosis, and degenerative disorders.

[0005] The International Union of Biochemistry and Molecular Biology (IUBMB) has recommended the use of the term "peptidase" for the subset of peptide bond hydrolases (Subclass E.C 3.4.). The widely used term protease is synonymous with peptidase. Peptidases comprise two groups of enzymes: the endopeptidases and the exopeptidases, which cleave peptide bonds at points within the protein and remove amino acids sequentially from either N or C-terminus respectively. The term proteinase is synonymous

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with endopeptidase. Proteolytic enzymes are classified according to their catalytic mechanisms. Four mechanistic classes have been recognized by the IUBMB: the serine proteases, cysteine proteases, aspartic proteases, and metalloproteases.

[0006] Serine proteases are a large family of proteolytic enzymes containing a serine residue in the active catalytic site for protein cleavage. They are ubiquitous being found in viruses, bacteria, and eukaryotes. Serine proteases have a wide range of substrate specificities and can be subdivided into subfamilies on the basis of these specificities. There are over 20 subfamilies of serine proteases which are grouped into six clans (SA, SB, SC, SE, SF, and SG).

[0007] Prolyl oligopeptidase is a serine protease grouped in the SC clan. It hydrolyzes proline containing peptides at the carboxyl side of proline residues. Presumably, it is involved in the maturation and degradation of peptide hormones and neuropeptides (Wilk *et al.* 1983 *Life Sci.* 33, 2149-2157). Examples of prolyl oligopeptidase include dipeptidyl peptidase IV (DPP-IV), dipeptidyl peptidase II (DPP-II), fibroblast activation protein, and prolyl oligopeptidase. These enzymes display distinct specificities.

[0008] Proline is present in numerous peptide hormones. It determines certain structural properties of these peptides, such as conformation and stability of these peptides, preventing degradation by non-specific proteases. A number of peptidases exist which attack the proline bonds. These peptidases are not only involved in the cleavage of X-Pro or Pro-X bonds, but also in the degradation of corresponding alanyl bonds, with reduced activity. Peptidases having highly specific actions on proline-containing sequences are attractive targets of medicinal chemistry because some of them have been linked to the modulation of the biological activity of natural peptide substrates. For example, DPP-IV is linked to the treatment of diabetes through regulating the level of glucagon-like peptide-1 (GLP-1). DPP-IV activity is increased in various diseases such as rheumatoid arthritis, multiple sclerosis, Grave's disease, and Hashimoto's thyroiditis, sarcoidosis, and cancer. DPP-IV activity is also increased in AIDS, Down's syndrome, anorexia/bulimia, pregnancy and hypogammaglobulinemia.

Dipeptidyl Peptidases Including DPP-IV

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[0009] Dipeptidyl aminopeptidase activity is peptidase activity which catalyzes the removal of dipeptides from the N-terminus of peptides, polypeptides, and proteins. Generally, a dipeptidyl aminopeptidase is capable of cleaving the dipeptide XY from the unsubstituted N-terminal amino group of a peptide, polypeptide or protein, wherein X and Y represent any amino acid residue. Examples of dipeptidyl peptidases (DPPs) include dipeptidyl peptidase I (DPP-I), dipeptidyl peptidase II (DPP-II), dipeptidyl peptidase III (DPP-III), and dipeptidyl peptidase (DPP-IV).

[0010] DPP-I, also known as cathepsin C, is a lysosomal cysteine protease that is expressed in most tissues. DPP-I has been implicated in the processing of granzymes, which are neutral serine proteases expressed exclusively in the granules of activated cytotoxic lymphocytes. DPP-II is a serine protease found in lysosomes. Like DPP-IV, it cleaves proline containing peptide bonds. In fact, DPP-II has a similar substrate specificity to DPP-IV but is only active at acidic pH. Dipeptidyl peptidase III (DPP-III) is a metalloprotease.

[0011] DPP-IV is a serine protease comprising the serine protease motif GWSYG and having broad substrate specificity. It hydrolyzes a peptide in sequence from the amino terminus to release an amino acid. However, the hydrolysis is terminated when an amino acid residue followed by proline is reached. As a result, a peptide having a bond of X-Pro-Y- (X and Y are optional amino acids) will be cleaved to yield X-Pro and Y-. DPP-IV will also cleave dipeptides with alanine in the penultimate position, though less effectively than dipeptides with proline (Yaron *et al.*, 1993 Crit. Rev. Biochem. Mol. Biol. 28:31-81). The enzyme will also cleave other sequences, but with still lower efficiency.

[0012] DPP-IV has been shown to be highly specific in releasing dipeptides from the N-terminal end of biologically active peptides with proline or alanine in the penultimate position of the N-terminal sequence of the peptide substrate. A large number of potential peptide substrates for DPP-IV have been identified. DPP-IV substrates include peptide hormones and chemokines. Examples of some peptide hormones are endomorphin-2, GLP-1, GLP-2, gastric inhibitory peptide (GIP), neuropeptide Y, growth hormone releasing hormone (GHRH) and substance P, and examples of some chemokines are RANTES, GCP-2, SDF-1 α , SDF-2 β , MDC, MCP-1, MCP-2, and MCP-3. DPP-II possesses almost identical substrate specificity to DPP-IV.

DPP-IV and Diabetes

[0013] Insulin-dependent diabetes mellitus (IDDM, or type I diabetes) is currently treated through the administration of insulin to patients. Non-insulin-dependent diabetes mellitus (NIDDM, or type II diabetes) is treated by diet, administration of sulphonylureas to stimulate insulin secretion or with biguanides to increase glucose uptake. Resistant individuals may need insulin therapy. Standard therapy requires daily intravenous injection of insulin which will treat the acute symptoms, but prolonged therapy results in vascular disease and nerve damage. Modern methods such as transplantation are expensive and require risky surgical intervention. Thus, there is a need to develop a highly effective, low cost alternative to the treatment of diabetes.

[0014] In recent years, there has been a growing interest in DPP-IV as a target for lowering the level of blood glucose. The use of inhibitors to block DPP-IV enzyme or DPP-IV-like enzyme activity in the blood of subjects leads to reduced degradation of endogenous or exogenously administered insulinotropic peptides such as, GIP, GLP-1 or analogs thereof. GIP and GLP-1, hormones that stimulate glucose-induced secretion of insulin by the pancreas, are substrates of DPP-IV. Specifically, since DPP-IV removes the amino-terminal His-Ala dipeptide of GLP-1 to generate GLP-1-(9-36)-amide, which is unable to elicit glucose-dependent insulin secretion from the islets, the inhibition of such DPP-IV or DPP-IV-like enzyme activity *in vivo* would effectively suppress undesired enzyme activity in pathological conditions in mammalian organisms.

[0015] PCT/DE97/00820 discloses alanyl pyrrolidide and isoleucyl thiazolidide as inhibitors of DPP-IV or DPP-IV-like enzyme activity. DD 296075 discloses pyrrolidide and isoleucyl thiazolidide hydrochloride. U.S. Patent 6,548,481 discloses inhibitors analogous to dipeptide compounds formed from an amino acid and a thiazolidine or pyrrolidine group, and salts thereof. Although these are functional inhibitors of DPP-IV activities, the use of these inhibitors in certain patients or certain forms of the disease may be problematic since the enzyme is responsible for activation or inactivation of such a wide range of bioactive peptides, *i.e.* DPP-IV inhibitors lack specificity for the desired targets GIP and GLP-1.

Protection of Therapeutic Peptides by Modification

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[0016] An alternative way to prevent therapeutic proteins and peptides such as GIP or GLP-1 from being cleaved by proteolytic enzymes is to modify the proteins and peptides themselves to block their exposure to proteolytic enzymes. Protein modifications have been shown to increase therapeutic polypeptides' stability, circulation time, and biological activity. Some general methods of modifying amino acids and peptides are disclosed in Chemistry and Biochemistry of Amino Acids, Peptides, and Proteins --A Survey of Recent Developments (Weinstein, B., ed., Marcel Dekker, Inc., publ., New York 1983) which is incorporated herein by reference. Also, the review article of Francis (1992 Focus on Growth Factors 3:4-10, (Mediscript, London)) describes protein modification and fusion proteins, which is incorporated herein by reference.

[0017] With the advance of recombinant DNA technology and automated techniques, one may now easily prepare large quantities of modified polypeptides that are short, medium or long. A large number of modified small polypeptide hormones may be synthesized using automated peptide synthesizers, solid-state resin techniques, or recombinant techniques. For example, large quantities of modified substrates of dipeptidyl peptidase, for example, the substrates of DPP-IV such as GLP-1, GIP, neuropeptide Y, and bradykinin can be produced using an automated peptide synthesizer.

SUMMARY OF THE INVENTION

[0018] The present invention provides transferrin fusion proteins comprising therapeutic peptides or proteins that are susceptible to protease cleavage. The present invention also provides transferrin fusion proteins comprising therapeutic peptides or proteins that are sensitive, resistant or partially resistant to protease cleavage. The protease may be DPP-IV or neutral endopeptidase (NEP). Moreover, the present invention provides compositions comprising transferrin fusion proteins and a second agent such as, but not limited to, inhibitors of DPP-IV and/or NEP. Further, the compositions may be pharmaceutical compositions used in the treatment of various diseases.

[0019] The present invention provides combination therapies comprising administering a transferrin fusion protein and at least one second agent in the treatment of various diseases. The transferrin fusion protein may be administered concurrently with the one or more second agent. Alternatively the transferrin fusion protein is administered sequentially,

either prior to or after the administration of the second agent. Preferably, the second agent is an inhibitor of DPP-IV or NEP.

[0020] The GLP-1 peptide moieties of the present invention may be modified to contain one or more mutations so that they are partially or fully resistant to protease cleavage, such as DPP-IV cleavage. The GLP-1 peptide may be GLP-1(7-37) (SEQ ID NO: 32) or GLP-1(7-36) (amino acids 1-30 of SEQ ID NO: 2). For example, these peptides may be modified by mutating A8 to G and/or K34 A.

BRIEF DESCRIPTION OF THE DRAWINGS

[0021] **Figure 1** shows the restriction enzyme map of pREX0094.

[0022] **Figure 2** shows the restriction enzyme map of plasmid pREX0198.

[0023] **Figure 3** shows the restriction enzyme map of pSAC35.

[0024] **Figure 4** shows the restriction enzyme map of plasmid pREX0240.

[0025] **Figure 5** shows the restriction enzyme map of pREX0052.

[0026] **Figure 6** shows the restriction enzyme map of pREX0367.

[0027] **Figure 7** shows the restriction enzyme map of pREX0368.

[0028] **Figure 8** shows time course of incubation of GLP-1 and H-GLP-1 and DPP-IV. The graph shows the amount of active, full length peptide remaining, as measured by an ELISA specific for active GLP-1.

DETAILED DESCRIPTION

1. General Description

[0029] This invention is based, in part, on the need to develop a more effective, low cost alternative for the treatment of diabetes. Insulinotropic peptides, such as GLP-1, are promising therapeutic agents for the treatment of type 2 non-insulin-dependent diabetes mellitus as well as related metabolic disorders, such as pre-diabetes, metabolic syndromes, and obesity. Other useful insulinotropic peptides include exendin 3 and exendin 4. However, these insulinotropic peptides have short plasma half-lives *in vivo*, mainly due to rapid serum clearance and proteolytic degradation. Extensive work has been done to inhibit DPP-IV, the enzyme responsible for the degradation of GLP-1 or to modify GLP-1 in such

a way that its degradation is slowed down while still maintaining biological activity. Despite these extensive efforts, a long lasting, active GLP-1 has not been produced. There is thus a need to modify GLP-1, exendin 3, exendin 4 and other insulinotropic peptides to provide longer duration of action *in vivo*, while maintaining their low toxicity and therapeutic advantages.

2. Definitions

[0030] As used herein, the term “derivative” refers to a modification of one or more amino acid residues of a peptide by chemical means, either with or without an enzyme, *e.g.*, by alkylation, acylation, ester formation, or amide formation.

[0031] As used herein, the term “derived from” refers to obtaining a molecule from a specified source such as obtaining a molecule from a parent molecule.

[0032] As used herein, the term “dipeptidyl aminopeptidase activity” refers to a peptidase activity which cleaves dipeptides from the N-terminal end of a peptide, polypeptide, or protein sequence. Generally, the dipeptidyl aminopeptidase is capable of cleaving the dipeptide XY from the unsubstituted N-terminal amino group of a peptide, polypeptide, or protein, wherein X or Y may represent any amino acid residue selected from the group consisting of Ala, Arg, Asn, Asp, Cys, Gln, Glu, Gly, His, Ile, Leu, Lys, Met, Phe, Pro, Ser, Thr, Trp, Tyr, and Val, but at least Ala, Arg, Asp, and/or Gly. Preferably, Y is Pro or Ala. All of X and Y may be different or identical. Examples of dipeptidyl aminopeptidase include, but are not limited to DPP-I, DPP-II, DPP-III, and DPP-IV.

[0033] As used herein, the terms “Glucagon-Like Peptide-1 (GLP-1)” and “GLP-1 derivatives” refer to intestinal hormones which generally simulate insulin secretion during hyperglycemia, suppress glucagon secretion, stimulate (pro) insulin biosynthesis and decelerate gastric emptying and acid secretion. Some GLP-1s and GLP-1 derivatives promote glucose uptake by cells but do not simulate insulin expression as disclosed in U.S. Pat. No. 5,574,008 which is hereby incorporated by reference.

[0034] As used herein, the term “insulinotropic peptides” refers to peptides with insulinotropic activity. Insulinotropic peptides stimulate, or cause the stimulation of, the synthesis or expression of the hormone insulin. Such peptides include precursors,

analogues, fragments of peptides such as Glucagon-like peptide 1, exendin 3 and exendin 4 and other peptides with insulinotropic activity.

[0035] As used herein, “pharmaceutically acceptable” refers to materials and compositions that are physiologically tolerable and do not typically produce an allergic or similar untoward reaction, such as gastric upset, dizziness and the like, when administered to a human. Typically, as used herein, the term “pharmaceutically acceptable” means approved by a regulatory agency of the Federal or a state government or listed in the U.S. Pharmacopeia or other generally recognized pharmacopeias for use in animals, and more particularly in humans.

[0036] As used herein, the term “pharmaceutical composition” refers to a composition comprising an agent together with a pharmaceutically acceptable carrier or diluent when needed. Pharmaceutically acceptable carriers and additives are chosen such that side effects from the pharmaceutical compound are minimized and the performance of the compound is not canceled or inhibited to such an extent that treatment is ineffective.

[0037] As used herein, “physiologically effective amount” is that amount delivered to a subject to give the desired palliative or curative effect. This amount is specific for each drug and its ultimate approved dosage level.

[0038] As used herein, “therapeutically effective amount” refers to that amount of modified therapeutic polypeptide or peptide which, when administered to a subject in need thereof, is sufficient to effect treatment. The amount of modified therapeutic polypeptide or peptide which constitutes a “therapeutically effective amount” will vary depending on the therapeutic protein used, the severity of the condition or disease, and the age and body weight of the subject to be treated, but can be determined routinely by one of ordinary skill in the art having regard to his/her own knowledge and to this disclosure.

[0039] As used herein, “therapeutic protein” refers to proteins, polypeptides, antibodies, peptide fragments or variants thereof, having one or more therapeutic and/or biological activities. Therapeutic proteins encompassed by the invention include but are not limited to proteins, polypeptides, peptides, antibodies and biologics. The terms peptides, proteins, and polypeptides are used interchangeably herein. Additionally, the term “therapeutic protein” may refer to the endogenous or naturally occurring correlate of a therapeutic protein. By a polypeptide or peptide displaying a “therapeutic activity” or a protein that is

“therapeutically active” is meant a polypeptide, peptide or protein that possesses one or more known biological and/or therapeutic activities associated with a therapeutic protein such as one or more of the therapeutic proteins described herein or otherwise known in the art. As a non-limiting example, a “therapeutic protein” is a protein, polypeptide, or peptide that is useful to treat, prevent or ameliorate a disease, condition or disorder. Such a disease, condition or disorder may be in humans or in a non-human animal, *e.g.*, veterinary use.

[0040] As used herein, the term “treatment” or “treating” refers to any administration of a compound of the present invention and includes: (1) preventing the disease from occurring in an animal which may be predisposed to the disease but does not yet experience or display the pathology or symptomatology of the disease; (2) inhibiting the disease in an animal that is experiencing or displaying the pathology or symptomatology of the disease (i.e., arresting further development of the pathology and/or symptomatology); or (3) ameliorating the disease in an animal that is experiencing or displaying the pathology or symptomatology of the disease (*i.e.*, reversing the pathology and/or symptomatology).

[0041] As used herein, the term “biological activity” refers to the ability to mediate a biological function. “Biological activity” includes functional activity as well as structural activity.

[0042] As used herein, the term “palliative” refers to the ability to relieve or soothe the symptoms of a disease or disorder without affecting a cure. For example, an agent that alleviates pain without curing the condition or disease is a palliative agent.

[0043] As used herein, the term “prophylactic” refers to the having protective effect such as acting to defend against or prevent something, especially disease or condition.

[0044] As used herein, “purified” protein or nucleic acid refers a protein or nucleic acid that has been separated from a cellular component. “Purified” proteins or nucleic acids have been purified to a level of purity not found in nature.

[0045] As used herein, the term "substantially pure" protein or nucleic acid refers to a protein or nucleic acid preparation that is lacking in all other cellular components.

[0046] As used herein, the term “therapeutic” refers to having a curative, restorative, or remedial effect. For example, a “therapeutic agent” or a “therapeutic composition” has a curative effect.

3. Specific Embodiments

Dipeptidyl Peptidases

[0047] Dipeptidyl peptidases are hydrolases that remove dipeptides from the unsubstituted N-terminal amino group of a peptide, polypeptide, or protein. Examples of dipeptidyl peptidases include but are not limited to DPP-I, DPP-II, DPP-III, DPP-IV, attractin, and fibroblast activation protein (FAP). New enzymes of this family or with similar function but different structure are emerging.

[0048] Dipeptidyl peptidase I (DPP-I), also known as cathepsin C, is a lysosomal cysteine protease belonging to the papain family. DPP-I is capable of sequentially removing dipeptides from the free amino terminus of various peptide and protein substrates, thus acting in the exopeptidase (specifically dipeptidyl peptidase) mode. The cleavage is ineffective if the fragmented bond has on either side a proline residue, or the N-terminal residue is lysine or arginine.

[0049] DPP-II is a serine protease found in lysosomes with unknown function. Like DPP-IV, it cleaves predominantly proline containing peptide bonds. In fact, DPP-II has a similar substrate specificity to DPP-IV but is only active at acidic pH. Mammalian DPP-II and DPP-IV can be distinguished using the inhibitors puromycin and bacitracin; puromycin will inhibit DPP-II only while bacitracin inhibits DPP-IV only (1988 J. Biol. Chem. 263, 6613-6618). Dipeptidyl peptidase III (DPP-III) is a metalloprotease. DPP-V releases N-terminal X-Ala, His-Ser, and Ser-Tyr dipeptides.

[0050] DPP-VII, also known as quiescent cell proline dipeptidase, is a proline-specific dipeptidase. It has been suggested that DPP-VII and DPP-II are identical proteases based on a sequence comparison of human DPP-VII and rat DPP-II (78% identity) (Araki *et al.* 2001 J. Biochem. 129, 279-288).

[0051] DPP-VIII is a human postproline dipeptidyl aminopeptidase that is homologous to DPP-IV and FAP (Abbott, C.A. *et al.*, 2000 European Journal of Biochemistry 267, 6140). Similar to DPP-IV, DPP-VIII is ubiquitous. The full-length DPP-VIII cDNA codes for an 882-amino-acid protein that has about 27% identity and 51% similarity to DPP-IV and FAP, but no transmembrane domain and no N-linked or O-linked glycosylation. Purified recombinant DPP-VIII hydrolyzed the DPP-IV substrates Ala-Pro, Arg-Pro and Gly-Pro. Thus recombinant DPP-VIII shares a postproline dipeptidyl aminopeptidase activity with

DPP-IV and FAP. DPP-VIII enzyme activity had a neutral pH optimum consistent with it being nonlysosomal. The similarities between DPP-VIII and DPP-IV in tissue expression pattern and substrates suggests a potential role for DPP-VIII in T-cell activation and immune function similar to DPP-IV.

[0052] Olsen C. *et al.* (2002 *Gene* 299, 185-93) report the identification and characterization of a novel DPP-IV-like molecule, termed dipeptidyl peptidase-like protein DPP-IX. Like DPP-IV, DPP-IX comprises the serine protease motif GWSYG (SEQ ID NO: 110). The presence of this motif and the conserved order and spacing of the Ser, Asp, and His residues that form the catalytic triad in DPP-IV, places DPP-IX in the DPP-IV gene family.

[0053] Attractin (DPPT-L) is a 175-kDa soluble glycoprotein reported to hydrolyze Gly-Pro. Attractin contains a kelch repeat domain and shares no significant sequence homology with DPP-IV or any other peptidase. Fibroblast activation protein (FAP) is a cell surface-bound protease of the prolyl oligopeptidase gene family expressed at sites of tissue remodelling.

[0054] Prolyl endopeptidase (PEP), also called proline oligopeptidase (PO), was first discovered by Walter and coworkers as an oxytocin-degrading enzyme in the human uterus (Walter *et al.*, *Science* 173, 827-829 (1971)). The enzyme cleaves peptide bonds at the carboxy-side of proline in peptides containing the sequence X-Pro-Y, where X is a peptide or N-terminal substituted amino-acid and Y is a peptide, amino acid, amide or alcohol (Yoshimoto *et al.*, *J. Biol. Chem.* 253, 3708-3716 (1979)). The enzyme has a high specificity for the trans-conformation of the peptide bond at the imino-side of proline (Lin & Brandts, *Biochemistry* 22, 4480-4485 (1983)).

[0055] Prolyl oligopeptidase hydrolyzes angiotensin I and angiotensin II which results in the release of angiotensin (1-7). Angiotensin (1-7) has vasodilator activity and modulates the release of vasopressin, which is able to influence the process of memory as was shown by injecting rats with specific PEP-inhibitors. The injection reverses the scopolamine induced amnesia. This experiment is not only an example which provides evidence for a possible physiologic function for the enzyme, but moreover it has led to the hypothesis that inhibitors for PEP can influence the memory process and counter dementia (Yoshimoto *et al.* 1987 *J. Pharmacobio-Dyn.* 10, 730-735).

Dipeptidyl Peptidase (DPP-IV) and Substrates

[0056] DPP-IV is a ubiquitously expressed molecule that has been implicated in the degradation of several peptides and hormones. Various types of DPP-IV have been purified and the enzymological properties have been revealed. For example, DPP-IV has been isolated from rat liver (Hopsu-Havu V. K. *et al.*, 1966 *Histochem.*, 7:197-201), swine kidney (Barth A. *et al.*, 1974 *Biol. Med. Chem.*, 32:157-174), small intestine (Svensson B. 1978 *Eur. J. Biochem.*, 90:489-498), liver (Fukasawa K. M. *et al.* 1981 *Biochim. Biophys. Acta*, 657:179-189), human submaxillary gland (Oya H., *et al.*, 1972 *Biochim. Biophys. Acta*, 258:591-599), sheep kidney (Yoshimoto T. *et al.*, 1977 *Biochim. Biophys. Acta*, 485:391-401; Yoshimoto T. *et al.*, 1978 *J. Biol. Chem.*, 253:3708-3716) or microorganisms (Fukasawa K. M. 1981 *Biochem. Biophys.*, 210:230-237; Yoshimoto T. 1982 *J. Biochem.*, 91:1899-1906 (1982)).

[0057] In the human immune system, DPP-IV is identical to the T-cell surface antigen CD26 which is expressed by activated lymphocytes (T-, B-, and natural killer cells). CD26/DPP-IV is a Type II membrane glycoprotein with intrinsic dipeptidyl peptidase IV activity and the ability to bind adenosine deaminase Type I (ADA-1). It is expressed on epithelial cells constitutively, but on T lymphocytes, it is expressed under tight cellular regulation, with expression upregulated upon cell activation. CD26/DPP-IV has been shown to have dipeptidyl peptidase IV activity in its extracellular domain (Hegen *et al.*, 1990 *J. Immunol* 144:2908-2914; Ulmer *et al.*, 1990 *Scand. J. Immunol.* 31:429-435) and the costimulatory activity appears to be partially dependent upon this enzyme activity (Tanaka *et al.*, 1993 *Proc. Natl. Acad. Sci. USA* 90:4586-4590). DPP-IV is involved in the regulation of chemokine function and may play an important role in HIV infection.

[0058] US Patent 6,265,551 discloses a circulating, soluble form of DPP-IV/CD26 isolated from human serum. The serum form shares similar enzymatic and antigenic properties with the ubiquitous membrane form; however, in several biochemical aspects there are distinct differences. In particular, the circulating serum form has a molecular weight of 175 kDa, in contrast to the 105 kDa molecular weight of the membrane form, and it does not bind ADA-1. Nevertheless, the circulating form expresses functional dipeptidylpeptidase IV activity and retains the ability to costimulate the T lymphocyte response to recall antigen.

[0059] The proteolytic activity of DPP-IV resides in a stretch of approximately 200 amino acids located at the C-terminal end of the protein. The catalytic residues (Ser-629, Asp-708, His-740) are arranged in a unique order which is different from the classical serine proteases such as chymotrypsin and subtilisin. Proline specific dipeptidyl peptidase activity alters the biological activity of a large number of bioactive proteins and polypeptides comprising, amongst others, GLP-1, the neurotransmitter substance P, human growth hormone-releasing factor, erythropoietin, interleukin 2 and many others. Potential DPP-IV substrates are listed in Tables 1, 2 and 3. Modulation of these polypeptides to affect DPP-IV cleavage may be useful in the treatment of clinical conditions including but not limited to diabetes, inflammation, vascular diseases, auto-immune disease, multiple sclerosis, joint diseases and diseases associated with benign and malign cell transformation.

TABLE 1: Human cytokines, growth factors, neuro- and vasoactive peptides with a penultimate proline, which are putative substrates for DPP-IV

Polypeptide	SEQ ID NO:	N-terminal sequence
Interleukin-1.beta.	1	Ala-Pro-Val-Arg-Ser-
Interleukin-2	2	Ala-Pro-Thr-Ser-Ser-
Interleukin-5	3	Ile-Pro-Thr-Glu-Ile-
Interleukin-6	4	Val-Pro-Pro-Gly-Glu-
Interleukin-10	5	Ser-Pro-Gly-Gln-Gly-
Interleukin-13 (recombinant)	6	Ser-Pro-Gly-Pro-Val-
Complement C4a	7	Lys-Pro-Arg-Leu-Leu-
Granulocyte chemotactic protein II	8	Gly-Pro-Val-Ser-Ala-
Granulocyte macrophage colony stimulating Factor	9	Ala-Pro-Ala-Arg-Ser-
Granulocyte colony stimulating factor	10	Thr-Pro-Leu-Gly-Pro-
Erythropoietin	11	Ala-Pro-Pro-Arg-Leu-
Gastrin releasing peptide growth hormone	12	Phe-Pro-Thr-Ile-Pro-
Interferon inducible peptide 10 (.gamma.IP10)	13	Val-Pro-Leu-Ser-Arg-
Interferon regulatory factor 1 (IRF-1)	14	Met-Pro-Ile-Thr-Arg
Interferon regulatory factor 2 (IRF-2)	15	Met-Pro-Val-Glu-Arg
Insulin-like growth factor-1	16	Gly-Pro-Glu-Thr-Leu-
Melanoma growth stimulating activity	17	Ala-Pro-Leu-Ala-Thr-
Migration inhibition factor	18	Met-Pro-Met-Phe-Ile-
Monocyte chemotactic protein I	19	Glu-Pro-Asp-Ala-Ile-
Neuropeptide Y	20	Tyr-Pro-Ser-Lys-Pro-
Pancreatic polypeptide	21	Ala-Pro-Leu-Glu-Pro-
Peptide YY	22	Try-Pro-Ile-Lys-Pro-
Prolactin	23	Leu-Pro-Ile-Cys-Pro-
RANTES	24	Ser-Pro-Tyr-Ser-Ser-
Substance P	25	Arg-Pro-Lys-Pro-Gln-

Polypeptide	SEQ ID NO:	N-terminal sequence
Thrombopoietin	26	Ser-Pro-Ala-Pro-Pro-
Transforming protein (N-myc) version 1	27	Met-Pro-Gly-Met-Ile-
Transforming protein (N-myc) version 2	28	Met-Pro-Ser-Cys-Ser-
Tumor necrosis factor .beta.	29	Leu-Pro-Gly-Val-Leu-
Vascular endothelial growth factor	30	Ala-Pro-Met-Ala-Glu-

TABLE 2: Human peptides and proteins with a penultimate alanine that are putative substrates for DPP IV

adenosine deaminase
Annexins
breast basic conserved protein
Cofilin
natural killer cell enhancing factor b
precursors of α -interferon
precursors of interleukin 1, α and 1, β and interleukin 13
precursors of macrophage inflammatory protein-2- α and 2- β
precursor of melanocyte stimulating hormone
precursor of oxytocin-neurophysin 1
growth hormone releasing hormone
β amyloid protein (1-28)
anxiety peptide
joining peptide of pro-opiomelanocortin

[0060] The present invention may utilize modified substrates of DPP comprising one or more additional amino acids at the N-terminus of the substrates to protect the substrates from DPP activity. The preferred substrates for modification according to the present invention are disclosed in Table 3.

[0061]

Table 3: Substrates for DPP-IV (CD26) Cleavage

DPP-IV Substrate	SEQ ID NO:	Sequence
GIP	31	YAEGTFISDY SIAMDKIHQQ DFNWLLAQK GKKNDWKHNI TQ
GLP-1	32 (Amino Acids 1- 29)	HAEGTFTSDV SSYLEGQAAK EFIAWL VKG
GLP-2	33	HADGSFSDEM NTILDNLAAR DFINWLIQTK ITD
growth hormone releasing hormone	34	YADAIFTNSY RKVLGQLSAR KLLQDIMSRQ QGESNQERGA RARL

DPP-IV Substrate	SEQ ID NO:	Sequence
Glucagon (slow inactivation, unlike GIP and the GLPs)	35	HSQGTFTSDY SKYLDSRRAQ DfvQWLMNT
peptide histidine-methionine	36	HADGVFTSDF SKLLGQLSAK KYLES LM
IGF-1	37	G PETLCGAELV DALQFVCGDR GFYFNKPTGY GSSRRAPQT GIVDECCFRS CDLRREMYC APLKPAKSAR SVRAQRHTDM PKAQKEVHLK NASRGSAGNK TY
Bradykinin	38	RPPGFSPFR
Substance P	39	RPKPQQFFGL M
CLIP	40	RPVKVYPNGA EDESAEAFPL EF
Neuropeptide Y	41	YPSKPDNPGE DAPAEDMARY YSALRHYINL ITRQRY
peptide YY (DPP-IV activates it)	42	YPIKPEAPGE DASPEELNRY YASLRHYLNL VTRQRY
Prolactin	43	LPICPGGAA RCQVTLRDLF DRAVVLSHYI HNLSEMFSFSE FDKRYTHGRG FITKAINSCH TSSLATPEDK EQAQQMNQKD FLSLIVSILR SWNEPLYHLV TEVRGMQEAP EAILSKAVEI EEQTKRLLEG MELIVSQVHP ETKENEIYPV WSGPLSLQMA DEESRLSA YY NLLHCLRRDS HKIDNYLKLK KCRHNNNC
human chorionic gonadotropin (HCG)	44	(alpha subunit) APDVQDCPEC TLQEDPFFSQ PGAPILQCMG CCFSRAYPTP LRSKKTMLVQ KNVTSSESTCC VAKSYNRVTV MGGFKVEDHT ACHCSTCYH KS
human chorionic gonadotropin (HCG)	45	(beta subunit) SKEPLRPRCR PINATLAVEK EGCPVCITVN TTICAGYCPT MTRVLQGVLP ALPQVVCNYR NVRFESIRLP GCPRGVNPVV SYAVALSCQC ALCRRSTDC GGPKDHPLTC DDPRFQDSSS SKAPPSLPS PSRLPKPSDT PILPQ
enterostatin	46	APGPR
gastrin-releasing peptide	47	VPLPAGGGTV LTKMYPRGNH WAVGHLM
IL-2	48	APTSSSTKKT QLQLEHLLLD LQMILNGINN YKNPKLTRML TFKFYMPKKA TELKHLQCLE EELKPLEEVL NLAQSKNFHL RPRDLISNIN VIVLELKGSE TTFMCEYADE TATIVEFLNR WITFCQSIIS TLT
IL-1b	49	APVR SLNCTLRDSQ QKSLVMSGPY ELKALHLQGG DMEQQVVFSM SFVQGEESND KIPVALGLKE KNLYLSCVLK DDKPTLQLES VDPKNYPKKK MEKRFVFNKI EINNKLFEFES AQFPNWIYST SQAENMPVFL GGTKGGQDIT DFTMQFVSS
endomorphin-2	50	YFFF
tyr-melanostatin	51	YPLG

DPP-IV Substrate	SEQ ID NO:	Sequence
aprotinin	52	RPDFCLEPPY TGPCCKARIIR YFYNAKAGLC QTFVYGGCRA KRNNFKSAED CMRTCGGA
RANTES	53	SPYSSDTTPC CFAYIARPLP RAHIKEYFYT SGKCSNPAVV FVTRKNRQVC ANPEKKWVRE YINSLEMS
trypsinogen	54	NPLLILTFV AAALAAPFDD DDKIVGGYNC EENSVPYQVS LNSGYHFCGG SLINEQWVVS AGHCYKSRIQ VRLGEHNIEV LEGNEQFINA AKIIRHPQYD RKTLNNDIML IKLSSRAVIN ARVSTISLPT APPATGTKCL ISGWGNTASS GADYPDELQC LDAPVLSQAK CEASYPGKIT SNMFCVGFLE GGKDCSQGDS GGPVVCGNQL QGVVSWGDGC AQKNKPGVYT KVYNYVKWIK NTIAANS
alpha1-microglobulin	55	G PVPTPPDNIQ VQENFNISRI YGKWYNLAIG STCPWLKKIM DRMTVSTLVL GEGATEAEIS MTSTRWRKGV CEETSGAYEK TDTDGKFLYH KSKWNITMES YVVHTNYDEY AIFLTKKFSR HHGPTITAKL YGRAPQLRET LLQDFRVVAQ GVGIPEDSIF TMADRGEVCP GEQEPEPILI PRV
interferon-inducible protein 10 (IP10)	56	VPLSRTVRCT CISISNQPVN PRSLEKLEII PASQFCPRVE IIATMKKKGE KRCLNPESKA IKNLLKAVSK ERSKRSP
Eotaxin	57	GPASVPTTCC FNLANRKIPL QRLESYRRIT SGKCPQKAVI FLTCLKAKDIC ADPKKKYVQD SMKYLDQKSP TPKP
Monocyte chemoattractant protein 1 (MCP-1)	58	QPDAINAPVT CCYNFTNRKI SVQRLASYRR ITSSKCPKEA VIFKTIVAKE ICADPKQKWV QDSMDHLDKQ TQTP
Monocyte chemoattractant protein 2 (MCP-2)	59	QPDSVSIPIT CCFNVINRKI PIQRLESYTR ITNIQCPKEA VIFKTKRGKE VCADPKERWV RDSMKHLDQI FQNLKP
Monocyte chemoattractant protein 3 (MCP-3)	60	QPVGINTSTT CCYRFINKKI PKQRLESYRR TTSSHCPREA VIFKTKLDKE ICADPTQKWV QDFMKHLDKK TQTPKL
Granulocyte chemotactic protein-2	61	GPV SAVLTELRCT CLRVTLRVNP KTIGKLQVFP AGPQCSKVEVV ASLKNKGQVC LDPEAPFLKK VIQKILDSGN KKN
SDF-1a	62	KPVSLSYRCP CRFFESHVAR ANVKHLKILN TPNCALQIVA RLKNNNRQVC IDPKLKWIQE YLEKALNK
SDF-1b	63	KPVSLSYRCP CRFFESHVAR ANVKHLKILN TPNCALQIVA RLKNNNRQVC IDPKLKWIQE YLEKALNKRF KM
Macrophage-derived chemokine	64	GPYGANMEDS VCCR DYVRYR LPLRVVKH FY WTSDSCPRPG VLLTFRDKE ICADPRVPWV KMILNKLSQ
b-casomorphin	65	YPFVEPI
Procolipase	66	APG PRGIIINLEN GELCMNSAQC KSNCCQHSSA LGLARCTSMA SENSECSVKT LYGIYYKCPC ERGLTCEGDK TIVGSITNTN FGICH DAGRS KQ

<u>DPP-IV Substrate</u>	<u>SEQ ID NO:</u>	<u>Sequence</u>
Vasoactive Intestinal Peptide (VIP)	67	HSDAVFTDNYTRLRKQMAVKKYLNSILN
Pituitary Adenylyl Cyclase-Activating Peptide 38 (PACAP38)	68	HSDGIFTDSYSRYRKQMAVKKYLA AVL GKRYKQRVKNK
Oxntomodulin	69	HSQGTFTSDYSKYLDSSRAQDFVQWLMNTKRNRNNIA
Growth hormone (1-43)	70	FPTIPLSRLFDNAMLRAHRLHQLAFDTYQEFEEAYIPKEQKYS
Secretin	71	HSDGFTSELSRLREGARLQRLQLV
Brain-derived natriuretic peptide		SPKMQVQSGCFGRKMDRISSSSGLGCKVLRH

[0062] The substrates for modification comprise X-ProY, X-Ala-Y, X-Ser-Y, or X-Gly-Y at the amino terminus. Preferably, the substrate for modification is GLP-1.

Modified Polypeptides Protected from DPP Activity

[0063] The present invention provides modified polypeptides, such as modified polypeptide substrates of DPP, comprising one or more additional amino acids at the N-terminus to protect the polypeptide substrates from DPP activity. In one embodiment, the modified polypeptides have one additional amino acid at their N-terminus as compared to the wild-type polypeptides. In another embodiment, the modified polypeptides have five additional amino acids at their N-terminus. Alternatively, the modified polypeptides have between one and five additional amino acids at their N-terminus. Any one of the 20 amino acids may be added to the N-terminus of the polypeptide substrate or non-natural amino acids may be added.

[0064] It is expected that any pharmaceutical polypeptide having peptide bonds which would be subject to cleavage in the circulation or anywhere *in vivo* after administration would benefit from modification in accordance with the present invention because of the protection from DPP cleavage that is afforded by the present invention.

[0065] In accordance with this aspect of the invention, it is possible to remove at least about 30%, preferably at least about 50%, more preferably at least about 70%, still more preferably at least about 90%, and most preferably at least about 99% of the dipeptidyl peptidase activity. It is also possible to completely remove the dipeptidyl aminopeptidase activity using the methods of the present invention.

[0066] Likewise, it is possible to reduce the substrate's dipeptidyl peptidase sensitivity by at least about 30%, preferably at least about 50%, more preferably at least about 70%, still more preferably at least about 90%, and most preferably at least about 99% of the dipeptidyl peptidase sensitivity. It is also possible to completely remove the dipeptidyl aminopeptidase sensitivity using the methods of the present invention.

[0067] Although the modified polypeptide or peptide substrates of the present invention are partially or substantially protected from DPP activity, the modified polypeptide substrates have retained at least about 10%, preferably at least about 30%, more preferably at least about 50%, more preferably at least about 70%, and still more preferably at least about 90%, and most preferably at least about 99% of their functional activity and potency. In some instances, the modified polypeptide or peptide substrates with lowered functional activity or potency will be useful. For example, when the modified polypeptide or peptide is fused to another polypeptide, such as transferrin, to form a fusion protein with increased serum stability and *in vivo* circulatory half-life, a modified polypeptide peptide substrate with lowered functional activity or potency may be useful.

[0068] In other instances, the modified polypeptides or peptides may have increased potency as compared to the non-modified polypeptides or peptides.

[0069] Modified polypeptide molecules of the invention are substantially protected from dipeptidyl peptidase cleavage as compared to an unmodified version of the same polypeptide. Qualification of this substantial protection may vary by the assay used to compare the modified versus unmodified polypeptide. In order to exhibit substantial protection, however, the modified polypeptide will exhibit a detectable level of resistance to dipeptidyl peptidase cleavage in the assay. Such assays include but are not limited to those disclosed in Doyle *et al.* (2002 *Endocrinology* 142, 4462-4468), O'Harte *et al.* (1999 *Diabetes* 48, 758-765) and Siegel *et al.* (1999 *Regulatory Peptides* 79, 93-102).

[0070] DPP stabilized polypeptide substrates of the present invention are also more stable in the presence of DPP *in vivo* than a non-stabilized polypeptide substrates. A DPP stabilized therapeutic polypeptide substrate generally has an increased activity half-life as compared to a non-stabilized peptide of identical sequence. Peptidase stability may be determined by comparing the half-life of the unmodified polypeptide substrate in serum or blood to the half-life of a modified counterpart therapeutic peptide in serum or blood. Half-life may be determined by sampling the serum or blood after administration of the modified

and non-modified peptides and determining the activity of the peptide. In addition to determining the activity, the length of the polypeptide substrates may also be measured by HPLC or Mass Spectrometry.

[0071] The present invention also provides modified polypeptides or peptides having an altered amino terminus according to the invention to protect against DPP cleavage and having internal and/or C-terminus amino acid alterations that do not affect the functional activity or potency of the polypeptide. These modified polypeptides would have minor amino acid changes that are usually conservative amino acid substitutions, although non-conservative substitutions are also contemplated.

[0072] The modified polypeptides or peptides of the present invention may also have altered functional activity. For instance, a modified polypeptide or peptide with increased functional activity may be useful. Alternatively, a modified polypeptide or peptide with decreased functional activity may be used. Thus, the modified polypeptides or peptides of the present invention also contain amino acid changes that do affect functional activity or potency. For example, the analogs of GLP-1 with altered functional activity may be modified at its amino terminus to protect against DPP cleavage.

[0073] Examples of conservative amino acid substitutions are substitutions made within the same group such as within the group of basic amino acids (such as arginine, lysine, histidine), acidic amino acids (such as glutamic acid and aspartic acid), polar amino acids (such as glutamine and asparagine), hydrophobic amino acids (such as leucine, isoleucine, valine), aromatic amino acids (such as phenylalanine, tryptophan, tyrosine) and small amino acids (such as glycine, alanine, serine, threonine, methionine).

[0074] Non-conservative substitutions encompass substitutions of amino acids in one group by amino acids in another group. For example, a non-conservative substitution would include the substitution of a polar amino acid for a hydrophobic amino acid. For a general description of nucleotide substitution, see *e.g.* Ford *et al.* (1991), *Prot. Exp. Pur.* 2: 95-107.

[0075] The present invention provides obvious variants of the amino acid sequence of the modified polypeptides and peptides, such as naturally occurring mature forms of the polypeptides or peptides, allelic/sequence variants of the polypeptides, non-naturally occurring recombinantly derived variants of the peptides, and orthologs and paralogs of the polypeptides or peptides. Such variants can readily be generated using art-known

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techniques in the fields of recombinant nucleic acid technology and protein biochemistry. Such variants can readily be identified/made using molecular techniques and the sequence information. Further, such variants can readily be distinguished from other peptides based on sequence and/or structural homology to the modified polypeptides or peptides of the present invention.

[0076] Preferably, the modified peptides of the present invention are GLP-1 and analogs thereof comprising one or more additional amino acids at their N-terminus.

[0077] In some instances, the DPP such as DPP-IV may activate a peptide instead of inactivating it through cleavage. In such instances, modification of the peptide could substantially reduce, delay, or prevent peptide activation.

Nucleic Acids Encoding Modified Polypeptides

[0078] The present invention provides nucleic acid molecules encoding modified polypeptides and peptides that are partially or substantially protected from DPP cleavage and have functional activity and potency. In one embodiment, nucleic acid molecules provided by the present invention encode modified polypeptides and peptides having at least one additional amino acid at its N-terminus as compared to their wild-type unmodified polypeptide. In another embodiment, the nucleic acid molecules encode modified polypeptides and peptides having five additional amino acids at their N-terminus. Alternatively, the nucleic acid molecules encode modified polypeptides and peptides having between one and five additional amino acids at their N-terminus. Preferably, the nucleic acid molecules encoding modified GLP-1 comprise sequence encoding one or more additional amino acids at its N-terminus.

[0079] The nucleic acid molecules of the invention include deoxyribonucleic acids (DNAs), both single- and double-stranded deoxyribonucleic acids. However, they can also be ribonucleic acids (RNAs), as well as hybrid RNA:DNA double-stranded molecules. Contemplated nucleic acid molecules also include genomic DNA, cDNA, mRNA, and antisense molecules. The nucleic acids molecules of the present invention also include native or synthetic RNA, DNA, or cDNA that encode a modified polypeptide, or the complementary strand thereof.

[0080] To construct modified polypeptides that are partially or substantially protected from DPP activity but having functional activity and/or potency compared to wild-type unmodified polypeptides, the nucleic acid encoding the wild-type unmodified polypeptide can be used as a starting point and modified to encode the desired modified polypeptide. Numerous methods are known to add sequences or to mutate nucleic acid sequences that encode a polypeptide and to confirm the function of the polypeptides encoded by these modified sequences.

[0081] The present invention also provides nucleic acids encoding polypeptides and peptides having a modified amino terminus for protection against DPP cleavage and having internal and C-terminus amino acid alterations that do not substantially affect the functional activity or potency of the polypeptide. These modified polypeptides would have minor amino acid changes that are usually conservative amino acid substitutions, although non-conservative substitutions are also contemplated. Nucleotide substitutions using techniques for accomplishing site-specific mutagenesis are well-known in the art. Preferably, the nucleic acids encode GLP-1 analogs having one or more additional amino acids at their N-terminus.

[0082] As known in the art "similarity" between two polynucleotides or polypeptides is determined by comparing the nucleotide or amino acid sequence and the conserved nucleotide or amino acid substitutes of one polynucleotide or polypeptide to the sequence of a second polynucleotide or polypeptide. Also known in the art is "identity" which means the degree of sequence relatedness between two polypeptide or two polynucleotide sequences as determined by the identity of the match between two strings of such sequences. Both identity and similarity can be readily calculated (Computational Molecular Biology, Lesk, A. M., ed., Oxford University Press, New York, 1988; Biocomputing: Informatics and Genome Projects, Smith, D. W., ed., Academic Press, New York, 1993; Computer Analysis of Sequence Data, Part I, Griffin, A. M., and Griffin, H. G., eds., Humana Press, New Jersey, 1994; Sequence Analysis in Molecular Biology, von Heinje, G., Academic Press, 1987; and Sequence Analysis Primer, Gribskov, M. and Devereux, J., eds., M Stockton Press, New York, 1991).

[0083] While there exist a number of methods to measure identity and similarity between two polynucleotide or polypeptide sequences, the terms "identity" and "similarity" are well known to skilled artisans (Sequence Analysis in Molecular Biology, von Heinje, G.,

Academic Press, 1987; Sequence Analysis Primer, Gribskov, M. and Devereux, J., eds., M Stockton Press, New York, 1991; and Carillo, H., and Lipman, D., SIAM J. Applied Math., 48: 1073 (1988). Methods commonly employed to determine identity or similarity between two sequences include, but are not limited to those disclosed in Guide to Huge Computers, Martin J. Bishop, ed., Academic Press, San Diego, 1994, and Carillo, H., and Lipman, D., SIAM J. Applied Math. 48:1073 (1988).

[0084] Preferred methods to determine identity are designed to give the largest match between the two sequences tested. Methods to determine identity and similarity are codified in computer programs. Preferred computer program methods to determine identity and similarity between two sequences include, but are not limited to, GCG program package (Devereux, *et al.*, Nucleic Acids Research 12(1):387 (1984)), BLASTP, BLASTN, FASTA (Atschul, *et al.*, J. Molec. Biol. 215:403 (1990)). The degree of similarity or identity referred to above is determined as the degree of identity between the two sequences indicating a derivation of the first sequence from the second. The degree of identity between two nucleic acid sequences may be determined by means of computer programs known in the art such as GAP provided in the GCG program package (Needleman and Wunsch (1970) Journal of Molecular Biology 48:443-453). For purposes of determining the degree of identity between two nucleic acid sequences for the present invention, GAP is used with the following settings: GAP creation penalty of 5.0 and GAP extension penalty of 0.3.

Codon Optimization

[0085] The degeneracy of the genetic code permits variations of the nucleotide sequence of polypeptides, while still producing a modified polypeptide comprising an identical amino acid sequence as the polypeptide encoded by a first DNA sequence. The procedure, known as "codon optimization" (described in U.S. Patent 5,547,871 which is incorporated herein by reference in its entirety) provides one with a means of designing such an altered DNA sequence. The design of codon optimized genes should take into account a variety of factors, including the frequency of codon usage in an organism, nearest neighbor frequencies, RNA stability, the potential for secondary structure formation, the route of synthesis and the intended future DNA manipulations of that gene. In particular, available

methods may be used to alter the codons encoding a given fusion protein with those most readily recognized by yeast when yeast expression systems are used.

[0086] The degeneracy of the genetic code permits the same amino acid sequence to be encoded and translated in many different ways. For example, leucine, serine and arginine are each encoded by six different codons, while valine, proline, threonine, alanine and glycine are each encoded by four different codons. However, the frequency of use of such synonymous codons varies from genome to genome among eukaryotes and prokaryotes. For example, synonymous codon-choice patterns among mammals are very similar, while evolutionarily distant organisms such as yeast (*S. cerevisiae*), bacteria (such as *E. coli*) and insects (such as *D. melanogaster*) reveal a clearly different pattern of genomic codon use frequencies (Grantham, R., *et al.*, Nucl. Acids Res., 8, 49-62 (1980); Grantham, R., *et al.*, Nucl. Acids Res., 9, 43-74 (1981); Maroyama, T., *et al.*, Nucl. Acids Res., 14, 151-197 (1986); Aota, S., *et al.*, Nucl. Acids Res., 16, 315-402 (1988); Wada, K., *et al.*, Nucl. Acids Res., 19 Supp., 1981-1985 (1991); Kurland, C. G., FEBS Letters, 285, 165-169 (1991)). These differences in codon-choice patterns appear to contribute to the overall expression levels of individual genes by modulating peptide elongation rates. (Kurland, C. G., FEBS Letters, 285, 165-169 (1991); Pedersen, S., EMBO J., 3, 2895-2898 (1984); Sorensen, M. A., J. Mol. Biol., 207, 365-377 (1989); Randall, L. L., *et al.*, Eur. J. Biochem., 107, 375-379 (1980); Curran, J. F., and Yarus, M., J. Mol. Biol., 209, 65-77 (1989); Varenne, S., *et al.*, J. Mol. Biol., 180, 549-576 (1984), Varenne, S., *et al.*, J. Mol. Biol., 180, 549-576 (1984); Garesl, J.-P., J. Theor. Biol., 43, 211-225 (1974); Ikemura, T., J. Mol. Biol., 146, 1-21 (1981); Ikemura, T., J. Mol. Biol., 151, 389-409 (1981)).

[0087] The preferred codon usage frequencies for a synthetic gene should reflect the codon usages of nuclear genes derived from the exact (or as closely related as possible) genome of the cell/organism that is intended to be used for recombinant protein expression, particularly that of yeast species. As discussed above, in one preferred embodiment the modified polypeptide is codon optimized, before or after modification as herein described for yeast expression.

Vectors

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[0088] Expression units for use in the present invention will generally comprise the following elements, operably linked in a 5' to 3' orientation: a transcriptional promoter, a secretory signal sequence, a DNA sequence encoding a modified polypeptide and a transcriptional terminator. As discussed above, any arrangement of the modified polypeptide and peptide may be used in the vectors of the invention. The selection of suitable promoters, signal sequences and terminators will be determined by the selected host cell and will be evident to one skilled in the art and are discussed more specifically below.

[0089] Suitable yeast vectors for use in the present invention are described in U.S. Patent 6,291,212 and include YRp7 (Struhl *et al.*, Proc. Natl. Acad. Sci. USA 76: 1035-1039, 1978), YEp13 (Broach *et al.*, Gene 8: 121-133, 1979), pJDB249 and pJDB219 (Beggs, Nature 275:104-108, 1978), pPPC0005, pSeCHSA, pScNHSA, pC4 and derivatives thereof. Useful yeast plasmid vectors also include pRS403-406, pRS413-416 and the *Pichia* vectors available from Stratagene Cloning Systems, La Jolla, CA 92037, USA. Plasmids pRS403, pRS404, pRS405 and pRS406 are Yeast Integrating plasmids (YIps) and incorporate the yeast selectable markers *HIS3*, *TRP1*, *LEU2* and *URA3*. Plasmids pRS413~41.6 are Yeast Centromere plasmids (YCps).

[0090] Such vectors will generally include a selectable marker, which may be one of any number of genes that exhibit a dominant phenotype for which a phenotypic assay exists to enable transformants to be selected. Preferred selectable markers are those that complement host cell auxotrophy, provide antibiotic resistance or enable a cell to utilize specific carbon sources, and include *LEU2* (Broach *et al.* *ibid.*), *URA3* (Botstein *et al.*, Gene 8: 17, 1979), *HIS3* (Struhl *et al.*, *ibid.*) or *POT1* (Kawasaki and Bell, EP 171,142). Other suitable selectable markers include the CAT gene, which confers chloramphenicol resistance on yeast cells. Preferred promoters for use in yeast include promoters from yeast glycolytic genes (Hitzeman *et al.*, J Biol. Chem. 225: 12073-12080, 1980; Alber and Kawasaki, J. Mol. Appl. Genet. 1: 419-434, 1982; Kawasaki, U.S. Pat. No. 4,599,311) or alcohol dehydrogenase genes (Young *et al.*, in Genetic Engineering of Microorganisms for Chemicals, Hollaender *et al.*, (eds.), p. 355, Plenum, N.Y., 1982; Ammerer, Meth. Enzymol. 101: 192-201, 1983). In this regard, particularly preferred promoters are the *TP11* promoter (Kawasaki, U.S. Pat. No. 4,599,311) and the *ADH2-4^C* (see U.S. Patent 6,291,212) promoter (Russell *et al.*, Nature 304: 652-654, 1983). The expression units may also include a transcriptional terminator. A preferred transcriptional terminator is the *TP11*

terminator (Alber and Kawasaki, *ibid.*). More preferably, the promoter is the *PRB1* promoter disclosed in EP 431880 and the terminator is the *ADH1* terminator disclosed in EP 60057, which are herein incorporated by reference in their entirety.

[0091] In addition to yeast, modified polypeptides and peptides of the present invention can be expressed in filamentous fungi, for example, species of the genus *Aspergillus*. Examples of useful promoters include those derived from *Aspergillus nidulans* glycolytic genes, such as the *ADH3* promoter (McKnight *et al.*, EMBO J. 4: 2093-2099, 1985) and the *tpiA* promoter. An example of a suitable terminator is the *ADH3* terminator (McKnight *et al.*, *ibid.*). The expression units utilizing such components may be cloned into vectors that are capable of insertion into the chromosomal DNA of *Aspergillus*, for example.

[0092] Mammalian expression vectors for use in carrying out the present invention will include a promoter capable of directing the transcription of the modified polypeptides and peptides. Preferred promoters include viral promoters and cellular promoters. Preferred viral promoters include the major late promoter from adenovirus 2 (Kaufman and Sharp, Mol. Cell. Biol. 2: 1304-13199, 1982) and the SV40 promoter (Subramani *et al.*, Mol. Cell. Biol. 1: 854-864, 1981). Preferred cellular promoters include the mouse metallothionein-1 promoter (Palmiter *et al.*, Science 222: 809-814, 1983) and a mouse V κ (see U.S. Patent 6,291,212) promoter (Grant *et al.*, Nuc. Acids Res. 15: 5496, 1987). A particularly preferred promoter is a mouse V H (see U.S. Patent 6,291,212) promoter. Such expression vectors may also contain a set of RNA splice sites located downstream from the promoter and upstream from the DNA sequence encoding the modified polypeptide or peptide. Preferred RNA splice sites may be obtained from adenovirus and/or immunoglobulin genes.

[0093] Also contained in the expression vectors is a polyadenylation signal located downstream of the coding sequence of interest. Polyadenylation signals include the early or late polyadenylation signals from SV40 (Kaufman and Sharp, *ibid.*), the polyadenylation signal from the adenovirus 5 E1B region and the human growth hormone gene terminator (DeNoto *et al.*, Nuc. Acids Res. 9: 3719-3730, 1981). A particularly preferred polyadenylation signal is the V H (see U.S. Patent 6,291,212) gene terminator. The expression vectors may include a noncoding viral leader sequence, such as the adenovirus 2 tripartite leader, located between the promoter and the RNA splice sites. Preferred vectors may also include enhancer sequences, such as the SV40 enhancer and the mouse μ (see U.S.

Patent 6,291,212) enhancer (Gillies, Cell 33: 717-728, 1983). Expression vectors may also include sequences encoding the adenovirus VA RNAs.

[0094] The expression vectors are also used for expressing fusion proteins comprising the modified polypeptide or peptide of the present invention fused to a second polypeptide or peptide, for example transferrin, to enhance the half-life of the modified polypeptide or peptide, as described below. Also, the modified polypeptide or peptide may be fused to a tag and/or a cleavage site for expression and release of the modified polypeptide or peptide.

Transformation

[0095] Techniques for transforming fungi are well known in the literature, and have been described, for instance, by Beggs (*ibid.*), Hinnen *et al.* (Proc. Natl. Acad. Sci. USA 75: 1929-1933, 1978), Yelton *et al.*, (Proc. Natl. Acad. Sci. USA 81: 1740-1747, 1984), and Russell (Nature 301: 167-169, 1983). The genotype of the host cell will generally contain a genetic defect that is complemented by the selectable marker present on the expression vector. Choice of a particular host and selectable marker is well within the level of ordinary skill in the art.

[0096] Cloned DNA sequences comprising modified polypeptides and peptides of the invention may be introduced into cultured mammalian cells by, for example, calcium phosphate-mediated transfection (Wigler *et al.*, Cell 14: 725, 1978; Corsaro and Pearson, Somatic Cell Genetics 7: 603, 1981; Graham and Van der Eb, Virology 52: 456, 1973.) Other techniques for introducing cloned DNA sequences into mammalian cells, such as electroporation (Neumann *et al.*, EMBO J. 1: 841-845, 1982), or lipofection may also be used. In order to identify cells that have integrated the cloned DNA, a selectable marker is generally introduced into the cells along with the gene or cDNA of interest. Preferred selectable markers for use in cultured mammalian cells include genes that confer resistance to drugs, such as neomycin, hygromycin, and methotrexate. The selectable marker may be an amplifiable selectable marker. A preferred amplifiable selectable marker is the DHFR gene. A particularly preferred amplifiable marker is the DHFR^r (see U.S. Patent 6,291,212) cDNA (Simonsen and Levinson, Proc. Natl. Acad. Sci. USA 80: 2495-2499, 1983). Selectable markers are reviewed by Thilly (Mammalian Cell Technology, Butterworth

Publishers, Stoneham, Mass.) and the choice of selectable markers is well within the level of ordinary skill in the art.

Host Cells

[0097] The present invention also includes a cell, preferably a yeast cell transformed to express a modified polypeptides or peptides of the invention. In addition to the transformed host cells themselves, the present invention also includes a culture of those cells, preferably a monoclonal (clonally homogeneous) culture, or a culture derived from a monoclonal culture, in a nutrient medium. If the polypeptide is secreted, the medium will contain the polypeptide, with the cells, or without the cells if they have been filtered or centrifuged away.

[0098] Host cells for use in practicing the present invention include eukaryotic cells, and in some cases prokaryotic cells, capable of being transformed or transfected with exogenous DNA and grown in culture, such as cultured mammalian, insect, fungal, plant and bacterial cells. A vector comprising a nucleic acid sequence of the present invention is introduced into a host cell so that the vector is maintained as a chromosomal integrant or as a self-replicating extra-chromosomal vector. Integration is generally considered to be an advantage as the nucleic acid sequence is more likely to be stably maintained in the cell. Integration of the vector into the host chromosome may occur by homologous or non-homologous recombination.

[0099] The choice of a host cell will to a large extent depend upon the gene encoding the polypeptide and its source. The host cell may be a unicellular microorganism, *e.g.*, a prokaryote, or a non-unicellular microorganism, *e.g.*, a eukaryote. Either prokaryotes or eukaryotes can be used. As prokaryotic host cells, generally used cells such as *Escherichia coli* or *Bacillus subtilis* can be used.

[00100] When prokaryotic cells are used as host cells, a vector replicable in the host cells may be used. An expression plasmid can be preferably used in which a promoter, an SD sequence (Shine-Dalgarno sequence), and an initiation codon (*e.g.* ATG) required for starting protein synthesis are provided in the vector upstream of the gene of the present invention to facilitate expression of the gene. Examples of the above vector include generally-used plasmids derived from *E. coli* such as pBR322, pBR325, pUC12, pUC13

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and the like. However, applicable vectors are not limited to these examples and various known vectors can also be used. Examples of commercially available vectors usable in expression system using *E. coli* include pGEX-4T (Amersham Pharmacia Biotech), pMAL-C2, pMAL-P2 (New England Biolabs), pET21/lacq (Invitrogen), pBAD/His (Invitrogen) and the like.

[00101] Examples of eukaryotic host cells include yeast cells and the like. Examples of preferably used craniate cells include COS cell (cell from monkey) (1981 Cell, 23, 175), Chinese Hamster Ovary cells and the dihydrofolate reductase defective strain derived therefrom (1980 Proc. Natl. Acad. Sci., USA., 77, 4216) and the like, and examples of preferably used yeast cells include *Saccharomyces cerevisiae* or the like. However, cells to be used are not limited to these examples. Preferably, a yeast cell is used to express the modified polypeptide or peptide.

[00102] Fungal cells, including species of yeast (*e.g.*, *Saccharomyces* spp., *Schizosaccharomyces* spp., *Pichia* spp.) may be used as host cells within the present invention. Examples of fungi including yeasts contemplated to be useful in the practice, of the present invention as hosts for expressing the modified polypeptide or peptides of the inventions are *Pichia* (including species formerly classified as *Hansenula*), *Saccharomyces*, *Kluyveromyces*, *Aspergillus*, *Candida*, *Torulopsis*, *Torulaspora*, *Schizosaccharomyces*, *Citeromyces*, *Pachysolen*, *Zygosaccharomyces*, *Debaromyces*, *Trichoderma*, *Cephalosporium*, *Humicola*, *Mucor*, *Neurospora*, *Yarrowia*, *Metschnikowia*, *Rhodosporidium*, *Leucosporidium*, *Botryosascus*, *Sporidiobolus*, *Endomycopsis*, and the like. Examples of *Saccharomyces* spp. are *S. cerevisiae*, *S. italicus* and *S. rouxii*. Examples of *Kluyveromyces* spp. are *K. fragilis*, *K. lactis* and *K. marxianus*. A suitable *Torulaspora* species is *T. delbrueckii*. Examples of *Pichia* spp. are *P. angusta* (formerly *H. polymorpha*), *P. anomala* (formerly *H. anomala*) and *P. pastoris*.

[00103] Particularly useful host cells to produce the modified polypeptide or peptide of the invention are the methanoltrophic *Pichia pastoris* (Steinlein *et al.* (1995) *Protein Express. Purif.* 6:619-624). *Pichia pastoris* has been developed to be an outstanding host for the production of foreign proteins since its alcohol oxidase promoter was isolated and cloned; its transformation was first reported in 1985. *P. pastoris* can utilize methanol as a carbon source in the absence of glucose. The *P. pastoris* expression system can use the methanol-induced alcohol oxidase (AOX1) promoter, which controls the gene that codes for the

expression of alcohol oxidase, the enzyme which catalyzes the first step in the metabolism of methanol. This promoter has been characterized and incorporated into a series of *P. pastoris* expression vectors. Since the proteins produced in *P. pastoris* are typically folded correctly and secreted into the medium, the fermentation of genetically engineered *P. pastoris* provides an excellent alternative to *E. coli* expression systems.

[00104] Strains of the yeast *Saccharomyces cerevisiae* are another preferred host. In a preferred embodiment, a yeast cell, or more specifically, a *Saccharomyces cerevisiae* host cell that contains a genetic deficiency in a gene required for asparagine-linked glycosylation of glycoproteins is used. *S. cerevisiae* host cells having such defects may be prepared using standard techniques of mutation and selection, although many available yeast strains have been modified to prevent or reduce glycosylation or hypermannosylation.

[00105] To optimize production of the heterologous proteins, it is also preferred that the host strain carry a mutation, such as the *S. cerevisiae pep4* mutation (Jones, Genetics 85: 23-33, 1977), which results in reduced proteolytic activity. It is particularly advantageous to use a host that carries a mutation in the gene encoding the aspartyl protease yapsin 1 (*YAP3*) or the gene encoding yapsin 2 (*MKC7*), or both (Copley *et al.* 1998 *Biochem. J.* 330, 1333-1340), such that the proteolytic activity directed to basic residues is reduced or eliminated. Host strains containing mutations in other protease encoding regions are particularly useful to produce large quantities of the modified therapeutic polypeptides or peptides of the invention.

[00106] Host cells containing DNA constructs of the present invention are grown in an appropriate growth medium. As used herein, the term "appropriate growth medium" means a medium containing nutrients required for the growth of cells. Nutrients required for cell growth may include a carbon source, a nitrogen source, essential amino acids, vitamins, minerals and growth factors. The growth medium will generally select for cells containing the DNA construct by, for example, drug selection or deficiency in an essential nutrient which is complemented by the selectable marker on the DNA construct or co-transfected with the DNA construct. Yeast cells, for example, are preferably grown in a chemically defined medium, comprising a non-amino acid nitrogen source, inorganic salts, vitamins and essential amino acid supplements. The pH of the medium is preferably maintained at a pH greater than 2 and less than 8, preferably at pH 5.5 to 6.5. Methods for maintaining a stable pH include buffering and constant pH control, preferably through the addition of

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ammonia, ammonium hydroxide or sodium hydroxide. Preferred buffering agents include citric acid, phosphate, succinic acid and Bis-Tris (Sigma Chemical Co., St. Louis, Mo.). Yeast cells having a defect in a gene required for asparagine-linked glycosylation are preferably grown in a medium containing an osmotic stabilizer. A preferred osmotic stabilizer is sorbitol supplemented into the medium at a concentration between 0.1 M and 1.5 M., preferably at 0.5 M or 1.0 M.

[00107] Cultured mammalian cells are generally grown in commercially available serum-containing or serum-free medium. Selection of a medium appropriate for the particular cell line used is within the level of ordinary skill in the art. Transfected mammalian cells are allowed to grow for a period of time, typically 1-2 days, to begin expressing the DNA sequence(s) of interest. Drug selection is then applied to select for growth of cells that are expressing the selectable marker in a stable fashion. For cells that have been transfected with an amplifiable selectable marker the drug concentration may be increased in a stepwise manner to select for increased copy number of the cloned sequences, thereby increasing expression levels.

[00108] Baculovirus/insect cell expression systems may also be used to produce the modified therapeutic polypeptides or peptides of the invention. The BacPAK™ Baculovirus Expression System (BD Biosciences (Clontech) expresses recombinant proteins at high levels in insect host cells. The target gene is inserted into a transfer vector, which is cotransfected into insect host cells with the linearized BacPAK6 viral DNA. The BacPAK6 DNA is missing an essential portion of the baculovirus genome. When the DNA recombines with the vector, the essential element is restored and the target gene is transferred to the baculovirus genome. Following recombination, a few viral plaques are picked and purified, and the recombinant phenotype is verified. The newly isolated recombinant virus can then be amplified and used to infect insect cell cultures to produce large amounts of the desired protein.

Secretory Signal Sequences

[00109] The terms "secretory signal sequence" or "signal sequence" or "secretion leader sequence" are used interchangeably and are described, for example in U.S. Pat. 6,291,212 and U.S. Pat 5,547,871, both of which are herein incorporated by reference in their entirety. Secretory signal sequences or signal sequences or secretion leader sequences encode

secretory peptides. A secretory peptide is an amino acid sequence that acts to direct the secretion of a mature polypeptide or protein from a cell. Secretory peptides are generally characterized by a core of hydrophobic amino acids and are typically (but not exclusively) found at the amino termini of newly synthesized proteins. Very often the secretory peptide is cleaved from the mature protein during secretion. Secretory peptides may contain processing sites that allow cleavage of the signal peptide from the mature protein as it passes through the secretory pathway. Processing sites may be encoded within the signal peptide or may be added to the signal peptide by, for example, *in vitro* mutagenesis.

[00110] Secretory peptides may be used to direct the secretion of modified polypeptides and peptides of the invention. One such secretory peptide that may be used in combination with other secretory peptides is the third domain of the yeast Barrier protein. Secretory signal sequences or signal sequences or secretion leader sequences are required for a complex series of post-translational processing steps which result in secretion of a protein. If an intact signal sequence is present, the protein being expressed enters the lumen of the rough endoplasmic reticulum and is then transported through the Golgi apparatus to secretory vesicles and is finally transported out of the cell. Generally, the signal sequence immediately follows the initiation codon and encodes a signal peptide at the amino-terminal end of the protein to be secreted. In most cases, the signal sequence is cleaved off by a specific protease, called a signal peptidase. Preferred signal sequences improve the processing and export efficiency of recombinant protein expression using viral, mammalian or yeast expression vectors. A preferred signal sequence is a mammalian or human transferrin signal sequence. In some cases, the native substrate signal sequence may be used to express and secrete modified polypeptide or peptides of the invention. In order to ensure efficient removal of the signal sequence, in some cases it may be preferable to include a short pro-peptide sequence between the signal sequence and the mature protein in which the C-terminal portion of the pro-peptide comprises a recognition site for a protease, such as the yeast *kex2p* protease. Preferably, the pro-peptide sequence is about 2-12 amino acids in length, more preferably about 4-8 amino acids in length. Examples of such pro-peptides are Arg-Ser-Leu-Asp-Lys-Arg, Arg-Ser-Leu-Asp-Arg-Arg, Arg-Ser-Leu-Glu-Lys-Arg, and Arg-Ser-Leu-Glu-Arg-Arg (SEQ ID NOS: 111-114, respectively).

Production of Modified Polypeptide Substrates Protected from DPP Cleavage

[00111] The modified polypeptides of this invention that are partially or substantially resistant to DPP activity, may be prepared by standard synthetic methods, recombinant DNA techniques, or any other methods of preparing peptides and fusion proteins.

[00112] The solid phase peptide synthesis method is generally described in the following references: Merrifield, *J. Am. Chem. Soc.*, 888:2149, 1963; Barany and Merrifield, In the *Peptides*, E. Gross and J. Meinenhofer, Eds., Academic Press, New York, 3:285 (1980); S. B. H. Kent. *Annu. Rev. Biochem.*, 57:957 (1988). By the solid phase peptide synthesis method, a peptide of a desired length and sequence can be produced through the stepwise addition of amino acids to a growing peptide chain which is covalently bound to a solid resin particle. Automated synthesis may be employed in this method.

[00113] As discussed above, the modified polypeptide of the present invention may also be obtained using molecular biology techniques, employing nucleic acid sequences that encode those polypeptides. Those sequences may be RNA or DNA and may be associated with control sequences and/or inserted into vectors. The latter are then transfected into host cells, for example bacteria. The preparation of the vectors and their production or expression in a host is carried out by conventional molecular biology and genetic engineering techniques.

[00114] Moreover, the modified polypeptides of the present invention can also be made by recombinant techniques using readily synthesized DNA sequences in commercially available expression systems.

[00115] The modified polypeptides of the present invention may be obtained by recombinant means comprising (a) cultivating a host cell under conditions conducive to production of the polypeptide; and (b) recovering the polypeptide. The cells are cultivated in a nutrient medium suitable for production of the polypeptide using methods known in the art. For example, the cell may be cultivated by shake flask cultivation, small-scale or large-scale fermentation (including continuous, batch, fed-batch, or solid state fermentations) in laboratory or industrial fermentors performed in a suitable medium and under conditions allowing the polypeptide to be expressed and/or isolated. The cultivation takes place in a suitable nutrient medium comprising carbon and nitrogen sources and inorganic salts, using procedures known in the art (see, *e.g.*, references for bacteria and yeast; Bennett, J. W. and LaSure, L., editors, *More Gene Manipulations in Fungi*, Academic Press, California, 1991). Suitable media are available from commercial suppliers or may be prepared according to

published compositions (e.g., in catalogues of the American Type Culture Collection). If the modified polypeptide is secreted into the nutrient medium, the polypeptide can be recovered directly from the medium. If the modified polypeptide is not secreted, it can be recovered from cell lysates.

[00116] As an example, the modified polypeptides or peptides of the present invention including the modified polypeptide or peptide fusion protein may be made by the fermentation methodology disclosed in WO 0044772, which is herein incorporated by reference in its entirety.

[00117] The modified polypeptides may be detected using methods known in the art that are specific for the polypeptides. These detection methods may include use of specific antibodies, formation of an enzyme product, or disappearance of an enzyme substrate, binding to a specific receptor, or by detection of activation of a specific receptor in a cell-based assay. For example, an enzyme assay may be used to determine the activity of the modified polypeptide. The resulting modified polypeptide may be recovered by methods known in the art. For example, the modified polypeptide may be recovered from the nutrient medium by conventional procedures including, but not limited to, centrifugation, filtration, extraction, spray-drying, evaporation, or precipitation.

[00118] The polypeptides of the present invention may be purified by a variety of procedures known in the art including, but not limited to, chromatography (e.g., ion exchange, affinity, hydrophobic, chromatofocusing, and size exclusion), electrophoretic procedures (e.g., preparative isoelectric focusing, differential solubility (e.g., ammonium sulfate precipitation), SDS-PAGE, or extraction (see, e.g., Protein Purification, J. -C. Janson and Lars Ryden, editors, VCH Publishers, New York, 1989).

Fusion Proteins and Protein Conjugates.

[00119] The present invention provides modified polypeptides or peptides attached to a heterologous molecule via recombinant means or covalent attachment. The attachment to a heterologous molecule, for example a plasma protein, extends the activity of the modified polypeptides or peptides for days to weeks. In some instances, only one administration of such modified therapeutic polypeptide or peptide need be given during this period of time. Greater specificity can be achieved, since the active compound will be primarily bound to

large molecules, where it is less likely to be taken up intracellularly to interfere with other physiological processes.

[00120] In another embodiment, the modified polypeptides or peptides of the present invention can be attached to heterologous sequences to form chimeric or fusion proteins via recombinant means. Such chimeric or fusion proteins comprise a modified polypeptide or peptide, partially or substantially protected from DPP cleavage, operatively linked to a heterologous protein having an amino acid sequence not substantially homologous to the modified polypeptide or peptide. "Operatively linked" indicates that the modified polypeptide or peptide and the heterologous protein are fused in-frame. The heterologous protein can be fused to the N-terminus or C-terminus of the modified polypeptide or peptide.

[00121] In one embodiment, the fusion protein does not affect the activity of the modified polypeptide of the invention per se. For example, the fusion protein can include, but is not limited to, enzymatic fusion proteins, for example beta-galactosidase fusions, yeast two-hybrid GAL fusions, poly-His fusions, MYC-tagged, HI-tagged and Ig fusions. Such fusion proteins, particularly poly-His fusions, can facilitate the purification of recombinant modified polypeptide. In a further example, the fusion protein comprises an amino acid sequence between the modified peptide of the invention and the other moiety, said amino acid sequence providing a recognition sequence that enables release of the modified peptide of the invention following chemical or enzymatic cleavage. In certain host cells (*e.g.*, mammalian host cells), expression and/or secretion of a protein can be increased by using a heterologous signal sequence. In another embodiment, the modified polypeptide or peptide is fused to a molecule that will extend its serum stability or serum half-life, such as a plasma protein. Preferably, the modified polypeptide or protein is fused to serum albumin, immunoglobulin, or a portion thereof such as the Fc domain. More preferably, the modified polypeptide or peptide is fused to transferrin, lactotransferrin, melanotransferrin, or hybrids thereof. Methods for making such fusion proteins are provided by U.S. Applications 10/231,494 and 10/378,094, and International Application PCT/US03/26818, which are herein incorporated by reference in their entirety.

[00122] As discussed in these applications, the transferrin to be attached to the modified polypeptide or peptide may be modified. It may exhibit reduced glycosylation. The modified transferrin polypeptide may be selected from the group consisting of a single

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transferrin N domain, a single transferrin C domain, a transferrin N and C domain, two transferrin N domains, and two transferrin C domains.

[00123] When the C domain of Tf is part of the fusion protein, the two N-linked glycosylation sites, amino acid residues corresponding to N413 and N611 (SEQ ID NO: 3 of PCT/US03/26818, which is incorporated by reference herein in its entirety) may be mutated for expression in a yeast system to prevent glycosylation or hypermannosylation and extend the serum half-life of the fusion protein and/or therapeutic protein (to produce asialo-, or in some instances, monosialo-Tf or disialo-Tf). In addition to Tf amino acids corresponding to N413 and N611, mutations may be to the adjacent residues within the N-X-S/T glycosylation site to prevent or substantially reduce glycosylation. See U.S. Patent 5,986,067 of Funk *et al.* It has also been reported that the N domain of Tf expressed in *Pichia pastoris* becomes O-linked glycosylated with a single hexose at S32 which also may be mutated or modified to prevent such glycosylation.

[00124] Accordingly, in one embodiment of the invention, the transferrin fusion protein includes a modified transferrin molecule wherein the transferrin exhibits reduced glycosylation, including but not limited to asialo- monosialo- and disialo- forms of Tf. In another embodiment, the transferrin portion of the transferrin fusion protein includes a recombinant transferrin mutant that is mutated to prevent glycosylation. In another embodiment, the transferrin portion of the transferrin fusion protein includes a recombinant transferrin mutant that is fully glycosylated. In a further embodiment, the transferrin portion of the transferrin fusion protein includes a recombinant human serum transferrin mutant that is mutated to prevent glycosylation, wherein at least one of Asn413 and Asn611 (SEQ ID NO: 3 of PCT/US03/26818, which is incorporated by reference herein in its entirety) are mutated to an amino acid which does not allow glycosylation. In another embodiment, the transferrin portion of the transferrin fusion protein includes a recombinant human serum transferrin mutant that is mutated to prevent or substantially reduce glycosylation, wherein mutations may be to the adjacent residues within the N-X-S/T glycosylation site. Moreover, glycosylation may be reduced or prevented by mutating the serine or threonine residue. Further, changing the X to proline is known to inhibit glycosylation.

[00125] A chimeric or fusion protein can be produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different protein sequences are

ligated together in-frame in accordance with conventional techniques. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers which give rise to complementary overhangs between two consecutive gene fragments which can subsequently be annealed and re-amplified to generate a chimeric gene sequence (see Ausubel *et al.* 1992 Current Protocols in Molecular Biology). Moreover, many expression vectors are commercially available that already encode a fusion moiety (*e.g.*, a GST protein). A modified polypeptide or peptide encoding nucleic acid can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the modified polypeptide or peptide.

[00126] In another embodiment, the modified therapeutic polypeptide or peptide is conjugated via a covalent bond to a heterologous molecule via a covalent bond to increase its stability and protection from DPP activity.

[00127] As an example, the modified polypeptide or peptide is conjugated to a blood component via a covalent bond formed between the reactive group of the modified peptide and a blood component, with or without a linking group. Blood components may be either fixed or mobile. Examples of fixed blood components are non-mobile blood components and include tissues, membrane receptors, interstitial proteins, fibrin proteins, collagens, platelets, endothelial cells, epithelial cells and their associated membrane and membranous receptors, somatic body cells, skeletal and smooth muscle cells, neuronal components, osteocytes and osteoclasts and all body tissues especially those associated with the circulatory and lymphatic systems. Example of mobile blood components are blood components that do not have a fixed situs for any extended period of time, generally not exceeding 5, more usually one minute. These blood components are not membrane-associated and are present in the blood for extended periods of time and are present in a minimum concentration of at least 0.1 $\mu\text{g/ml}$. Mobile blood components include serum albumin, transferrin, immunoglobulins such as IgM and IgG, α_1 protease inhibitor, antithrombin III and α_2 -antiplasmin. The half-life of mobile blood components is typically at least about 12 hours.

[00128] The formation of the covalent bond between the blood component and the modified therapeutic polypeptide or peptide may occur *in vivo* or *ex vivo*. For *ex vivo* covalent bond formation, the modified polypeptide or peptide is added to blood, serum or

saline solution containing the blood component, *e.g.* human serum albumin or IgG to permit covalent bond formation between the modified polypeptide or peptide and the blood component. Also, the modified polypeptide or peptide may be modified with maleimide or a similarly reactive chemical group and reacted with a blood component in saline solution. Once the modified therapeutic polypeptide or peptide is reacted with the blood component to form a modified polypeptide or peptide conjugate, the conjugate may be administered to the patient. Alternatively, the modified therapeutic polypeptide or peptide may be administered to the patient directly so that the covalent bond forms between the modified therapeutic polypeptide or peptide and the blood component *in vivo*. Also, the same reaction may be carried out with a recombinant protein, for example, albumin.

[00129] The various sites with which the chemically reactive groups of the non-specific modified therapeutic polypeptide or peptide may react *in vivo* include cells, particularly red blood cells (erythrocytes) and platelets, and proteins, such as immunoglobulins, including IgG and IgM, serum albumin, ferritin, steroid binding proteins, transferrin, thyroxin binding protein, α -2-macroglobulin, and the like.

[00130] The modified polypeptide or peptide may contain or may be chemically modified to contain a reactive group for binding to thiol. In one embodiment of the invention the modified polypeptide or peptide may be conjugated to polyethylene glycol. Alternatively, the modified polypeptide or peptide may be conjugated to a polyethylene glycol modified glycolipid or polyethylene glycol modified fatty acid.

[00131] In one aspect, the modified polypeptide or peptide may be conjugated to a fatty acid or fatty acid derivative to improve its stability. Examples of fatty acids include, but are not limited to, lauric, palmitic, oleic, and stearic acids. Examples of fatty acid derivatives include ethyl esters, propyl esters, cholesteryl esters, coenzyme A esters, nitrophenyl esters, naphthyl esters, monoglycerides, diglycerides, and triglycerides, fatty alcohols, fatty alcohol acetates, and the like.

[00132] In another aspect, the modified polypeptide or peptide may be engineered into a drug affinity complex (DACTM). A drug affinity complex has three parts: a drug component which is responsible for biological activity; a connector attaching the drug component to the reactive chemistry group; and a reactive chemistry group, at the opposite end of the connector, which is responsible for the permanent bonding of the construct to certain target proteins in the body. For example, Kim *et al.* (2003, Diabetes 52(3):751) disclose a GLP-1-

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albumin drug affinity complex. Kim *et al.* show that the albumin-conjugated DAC:GLP-1 bound to the GLP-1 receptor (GLP-1R) and activated cAMP formation in heterologous fibroblasts expressing the receptor. The results suggest that the albumin-conjugated DAC:GLP-1 mimics the native GLP-1. Kim *et al.* provide a new approach for prolonged activation of GLP-1R signaling.

[00133] The modified polypeptide or peptide drug affinity complex is designed to be administered by subcutaneous injection and then rapidly and selectively bonds *in vivo* to albumin. The bioconjugate formed has the same therapeutic activity and similar potency as endogenous polypeptide or peptide but has a pharmacokinetic profile in animals that is closer to that of albumin.

Pharmaceutical Composition

[00134] The present invention provides pharmaceutical compositions comprising modified therapeutic polypeptides and peptides partially or substantially protected from DPP cleavage, but substantially retaining their functional activity and potency. Such pharmaceutical compositions may be administered orally, parenterally, such as intravascularly (IV), intraarterially (IA), intramuscularly (IM), subcutaneously (SC), intraperitoneally, transdermally, or the like. Administration may in appropriate situations be by transfusion. In some instances, administration may be oral, nasal, rectal, transdermal or aerosol, where the modified polypeptide allows for transfer to the vascular system. For example, fusion or conjugation of a modified polypeptide of the invention to a transferrin moiety allows for transport of the modified polypeptide to the vascular system or across the blood-brain barrier via binding to the transferrin receptor, as described in International Application PCT/US03/26778, which is herein incorporated by reference in its entirety. Usually a single injection will be employed although more than one injection may be used, if desired. The modified therapeutic polypeptides or peptides may be administered by any convenient means, including syringe, trocar, catheter, or the like. The particular manner of administration will vary depending upon the amount to be administered, whether a single bolus or continuous administration, or the like. Preferably, the administration will be intravascularly, where the site of introduction is not critical to this invention, preferably at a site where there is rapid blood flow, *e.g.*, intravenously, peripheral or central vein. More preferably, the pharmaceutical compositions will be administered subcutaneously. Other

routes may find use where the administration is coupled with slow release techniques or a protective matrix. The intent is that the modified therapeutic peptides or polypeptides be effectively distributed, for example, in the blood, so as to be able to react with the blood or tissue components.

[00135] Generally, the invention encompasses pharmaceutical compositions comprising effective amounts of modified therapeutic polypeptide or peptide of the invention together with pharmaceutically acceptable diluents, preservatives, solubilizers, emulsifiers, adjuvants and/or carriers. Such compositions may include diluents of various buffer content (*e.g.*, Tris-HCl, acetate, phosphate), pH and ionic strength; additives such as detergents and solubilizing agents (*e.g.*, Polysorbate 80), anti-oxidants (*e.g.*, ascorbic acid, sodium metabisulfite), preservatives (*e.g.*, Thimersol, benzyl alcohol) and bulking substances (*e.g.*, lactose, mannitol); incorporation of the material into particulate preparations of polymeric compounds such as polylactic acid, polyglycolic acid, *etc.* or into liposomes. Hyaluronic acid may also be used, and this may have the effect of promoting sustained duration in the circulation. Such compositions may influence the physical state, stability, rate of *in vivo* release, and rate of *in vivo* clearance of the present proteins and derivatives. See, *e.g.*, Remington's Pharmaceutical Sciences, 18th Ed. (1990, Mack Publishing Co., Easton, Pa. 18042) pages 1435-1712 which are herein incorporated by reference.

[00136] For example, the modified therapeutic polypeptides or peptides may be administered in a physiologically acceptable medium, *e.g.*, deionized water, phosphate buffered saline (PBS), saline, aqueous ethanol or other alcohol, plasma, proteinaceous solutions, mannitol, aqueous glucose, alcohol, vegetable oil, or the like. Other additives which may be included include buffers, where the media are generally buffered at a pH in the range of about 5 to 10, where the buffer will generally range in concentration from about 50 to 250 mM, salt, where the concentration of salt will generally range from about 5 to 500 mM, physiologically acceptable stabilizers, and the like. Examples of physiological buffers, especially for injection, include Hank's solution and Ringer's solution. Transdermal formulations may contain penetrants such as bile salts or fusidates.

[00137] The pharmaceutical compositions may be prepared as tablets or dragees, sublingual tablets, sachets, paquets, soft gelatin capsules, suppositories, creams, ointments, dermal gels, transdermal devices, aerosols, drinkable and injectable ampoules. The compositions may also be prepared in liquid form, or may be in dried powder, such as lyophilized form

convenient for storage and transport. Implantable sustained release formulations are also contemplated.

Oral Dosage Forms

[00138] In one embodiment, the present invention provides pharmaceutical compositions comprising the modified therapeutic polypeptides or peptides in oral solid dosage forms, which are described generally in Remington's Pharmaceutical Sciences (1990), 18th Ed., Mack Publishing Co. Easton Pa. 18042, which is herein incorporated by reference. Solid dosage forms include tablets, capsules, pills, troches or lozenges, cachets or pellets. Also, liposomal or proteinoid encapsulation may be used to formulate the present compositions (as, for example, proteinoid microspheres reported in U.S. Pat. No. 4,925,673). Liposomal encapsulation may be used and the liposomes may be derivatized with various polymers (e.g., U.S. Pat. No. 5,013,556). A description of possible solid dosage forms for the therapeutic is given in Chapter 10 of Marshall, K., Modern Pharmaceutics (1979), edited by G. S. Banker and C. T. Rhodes, herein incorporated by reference. In general, the formulation will include the modified therapeutic polypeptide or peptide, and inert ingredients which allow for protection against the stomach environment, and release of the biologically active material in the intestine.

[00139] If necessary, the modified therapeutic polypeptide or peptide may be chemically modified so that oral delivery is efficacious. Generally, the chemical modification contemplated is the attachment of at least one moiety to the modified therapeutic polypeptide or peptide itself, where said moiety permits uptake into the blood stream from the stomach or intestine. Also desired is the increase in overall stability of the compound and increase in circulation time in the body. Moieties useful as covalently attached vehicles in this invention may also be used for this purpose. Examples of such moieties include: PEG, copolymers of ethylene glycol and propylene glycol, carboxymethyl cellulose, dextran, polyvinyl alcohol, polyvinyl pyrrolidone and polyproline. See, for example, Abuchowski and Davis, Soluble Polymer-Enzyme Adducts, Enzymes as Drugs (1981), Hocenberg and Roberts, eds., Wiley-Interscience, New York, N.Y., pp 367-83; Newmark, *et al.* (1982), J. Appl. Biochem. 4:185-9. Other polymers that could be used are poly-1,3-dioxolane and poly-1,3,6-tioxocane. Preferred for pharmaceutical usage, as indicated above, are PEG moieties.

[00140] Likewise, the modified therapeutic polypeptide or peptide may be recombinantly fused to another polypeptide to increase its overall stability or improve oral delivery. For example, the modified therapeutic polypeptide or peptide may be fused to transferrin, melanotransferrin, or lactoferrin. Methods for making such fusion proteins are described in U.S. Application 10/378,094, which is herein incorporated by reference in its entirety.

[00141] For oral delivery dosage forms, it is also possible to use a salt of a modified aliphatic amino acid, such as sodium N-(8-[2-hydroxybenzoyl] amino) caprylate (SNAC), as a carrier to enhance absorption of the therapeutic compounds of this invention. The clinical efficacy of a heparin formulation using SNAC has been demonstrated in a Phase II trial conducted by Emisphere Technologies. See U.S. Pat. No. 5,792,451, "Oral drug delivery composition and methods" which is herein incorporated by reference in its entirety.

[00142] The modified therapeutic polypeptides or peptides of this invention can be included in the formulation as fine multiparticulates in the form of granules or pellets of particle size about 1 mm. The formulation of the material for capsule administration could also be as a powder, lightly compressed plugs or even as tablets. The therapeutic could be prepared by compression.

[00143] Colorants and flavoring agents may all be included. For example, the modified therapeutic polypeptide or peptide may be formulated (such as by liposome or microsphere encapsulation) and then further contained within an edible product, such as a refrigerated beverage containing colorants and flavoring agents.

[00144] One may dilute or increase the volume of the pharmaceutical composition of the invention with an inert material. These diluents could include carbohydrates, especially mannitol, cc-lactose, anhydrous lactose, cellulose, sucrose, modified dextrans and starch. Certain inorganic salts may also be used as fillers including calcium triphosphate, magnesium carbonate and sodium chloride. Some commercially available diluents are Fast-Flo, Emdex, STA-Rx 1500, Emcompress and Avicell.

[00145] Disintegrants may be included in the formulation of the therapeutic into a solid dosage form. Materials used as disintegrants include but are not limited to starch including the commercial disintegrant based on starch, Explotab. Sodium starch glycolate, Amberlite, sodium carboxymethylcellulose, ultramylopectin, sodium alginate, gelatin, orange peel, acid carboxymethyl cellulose, natural sponge and bentonite may all be used. Another form of

the disintegrants are the insoluble cationic exchange resins. Powdered gums may be used as disintegrants and as binders and these can include powdered gums such as agar, Karaya or tragacanth. Alginic acid and its sodium salt are also useful as disintegrants.

[00146] Binders may be used to hold the modified therapeutic polypeptide or peptide together to form a hard tablet and include materials from natural products such as acacia, tragacanth, starch and gelatin. Others include methyl cellulose (MC), ethyl cellulose (EC) and carboxymethyl cellulose (CMC). Polyvinyl pyrrolidone (PVP) and hydroxypropylmethyl cellulose (HPMC) could both be used in alcoholic solutions to granulate the therapeutic.

[00147] An antifrictional agent may be included in the formulation of the pharmaceutical composition of the invention to prevent sticking during the formulation process. Lubricants may be used as a layer between the modified therapeutic polypeptide or peptide and the die wall, and these can include but are not limited to; stearic acid including its magnesium and calcium salts, polytetrafluoroethylene (PTFE), liquid paraffin, vegetable oils and waxes. Soluble lubricants may also be used such as sodium lauryl sulfate, magnesium lauryl sulfate, polyethylene glycol of various molecular weights, Carbowax 4000 and 6000.

[00148] Glidants that might improve the flow properties of the modified therapeutic polypeptide or peptide during formulation and to aid rearrangement during compression might be added. The glidants may include starch, talc, pyrogenic silica and hydrated silicoaluminate.

[00149] To aid dissolution of the modified therapeutic polypeptide or peptide of this invention into the aqueous environment a surfactant might be added as a wetting agent. Surfactants may include anionic detergents such as sodium lauryl sulfate, dioctyl sodium sulfosuccinate and dioctyl sodium sulfonate. Cationic detergents might be used and could include benzalkonium chloride or benzethonium chloride. The list of potential nonionic detergents that could be included in the formulation as surfactants are lauromacrogol 400, polyoxyl 40 stearate, polyoxyethylene hydrogenated castor oil 10, 50 and 60, glycerol monostearate, polysorbate 40, 60, 65 and 80, sucrose fatty acid ester, methyl cellulose and carboxymethyl cellulose. These surfactants could be present in the formulation of the protein or derivative either alone or as a mixture in different ratios.

[00150] Additives may also be included in the formulation to enhance uptake of the modified therapeutic polypeptide and peptide. Additives potentially having this property are for instance the fatty acids oleic acid, linoleic acid and linolenic acid.

[00151] Controlled release formulation also may be desirable. The modified therapeutic polypeptide or peptide of this invention could be incorporated into an inert matrix which permits release by either diffusion or leaching mechanisms *e.g.*, gums. Slowly degenerating matrices may also be incorporated into the formulation, *e.g.*, alginates, polysaccharides. Another form of a controlled release of the compounds of this invention is by a method based on the Oros therapeutic system (Alza Corp.), *i.e.*, the drug is enclosed in a semipermeable membrane which allows water to enter and push drug out through a single small opening due to osmotic effects. Some enteric coatings also have a delayed release effect.

[00152] Other coatings may be used for the formulation. These include a variety of sugars which could be applied in a coating pan. The modified therapeutic polypeptide or peptide could also be given in a film coated tablet and the materials used in this instance are divided into 2 groups. The first are the nonenteric materials and include methyl cellulose, ethyl cellulose, hydroxyethyl cellulose, methylhydroxy-ethyl cellulose, hydroxypropyl cellulose, hydroxypropyl-methyl cellulose, sodium carboxy-methyl cellulose, providone and the polyethylene glycols. The second group consists of the enteric materials that are commonly esters of phthalic acid.

[00153] A mix of materials might be used to provide the optimum film coating. Film coating may be carried out in a pan coater or in a fluidized bed or by compression coating.

Pulmonary Delivery Forms

[00154] In another embodiment, the present invention also provides pharmaceutical compositions comprising the modified therapeutic polypeptides or peptides for pulmonary delivery. The pharmaceutical composition is delivered to the lungs of a mammal while inhaling and traverses across the lung epithelial lining to the blood stream.

[00155] The present invention provides the use of a wide range of mechanical devices designed for pulmonary delivery of therapeutic products, including but not limited to nebulizers, metered dose inhalers, and powder inhalers, all of which are familiar to those

skilled in the art. Some specific examples of commercially available devices suitable for the practice of this invention are the Ultravent nebulizer, manufactured by Mallinckrodt, Inc., St. Louis, Mo.; the Acorn II nebulizer, manufactured by Marquest Medical Products, Englewood, Colo.; the Ventolin metered dose inhaler, manufactured by Glaxo Inc., Research Triangle Park, N.C.; and the Spinhaler powder inhaler, manufactured by Fisons Corp., Bedford, Mass.

[00156] All such devices require the use of formulations suitable for the dispensing of the modified therapeutic polypeptide and peptide. Typically, each formulation is specific to the type of device employed and may involve the use of an appropriate propellant material, in addition to diluents, adjuvants and/or carriers useful in therapy.

[00157] The modified therapeutic polypeptide or peptide should most advantageously be prepared in particulate form with an average particle size of less than 10 μm , most preferably 0.5 to 5 μm , for most effective delivery to the distal lung.

[00158] Pharmaceutically acceptable carriers include carbohydrates such as trehalose, mannitol, xylitol, sucrose, lactose, and sorbitol. Other ingredients for use in formulations may include DPPC, DOPE, DSPC and DOPC. Natural or synthetic surfactants may be used. PEG may be used (even apart from its use in derivatizing the protein or analog). Dextrans, such as cyclodextran, may be used. Bile salts and other related enhancers may be used. Cellulose and cellulose derivatives may be used. Amino acids may be used, such as use in a buffer formulation.

[00159] Also, the use of liposomes, microcapsules or microspheres, inclusion complexes, or other types of carriers is contemplated.

[00160] Formulations suitable for use with a nebulizer, either jet or ultrasonic, will typically comprise the inventive compound dissolved in water at a concentration of about 0.1 to 25 mg of biologically active protein per mL of solution. The formulation may also include a buffer and a simple sugar (*e.g.*, for protein stabilization and regulation of osmotic pressure). The nebulizer formulation may also contain a surfactant, to reduce or prevent surface induced aggregation of the protein caused by atomization of the solution in forming the aerosol.

[00161] Formulations for use with a metered-dose inhaler device will generally comprise a finely divided powder containing the inventive compound suspended in a propellant with

the aid of a surfactant. The propellant may be any conventional material employed for this purpose, such as a chlorofluorocarbon, a hydrochlorofluorocarbon, a hydrofluorocarbon, or a hydrocarbon, including trichlorofluoromethane, dichlorodifluoromethane, dichlorotetrafluoroethanol, and 1,1,1,2-tetrafluoroethane, or combinations thereof. Suitable surfactants include sorbitan trioleate and soya lecithin. Oleic acid may also be useful as a surfactant.

[00162] Formulations for dispensing from a powder inhaler device will comprise a finely divided dry powder containing the inventive compound and may also include a bulking agent, such as lactose, sorbitol, sucrose, mannitol, trehalose, or xylitol in amounts which facilitate dispersal of the powder from the device, *e.g.*, about 50 to 90% by weight of the formulation.

Nasal Delivery Forms

[00163] Nasal delivery of the pharmaceutical composition of the modified polypeptide or peptide of the present invention is also contemplated. Nasal delivery allows the passage of the protein to the blood stream directly after administering the modified therapeutic polypeptide or peptide to the nose, without the necessity for deposition of the product in the lung. Formulations for nasal delivery include those with dextran or cyclodextran. Delivery via transport across other mucous membranes is also disclosed.

Dosages

[00164] The dosage regimen involved in a method for treating the above-described conditions will be determined by the attending physician, considering various factors which modify the action of drugs, *e.g.* the age, condition, body weight, sex and diet of the patient, the severity of any infection, time of administration and other clinical factors. Generally, the daily regimen should be in the range of 0.01-1000 micrograms of the inventive compound per kilogram of body weight, preferably 0.1-150 micrograms per kilogram.

Treatment of Diseases with Modified Therapeutic Proteins

[00165] The present invention provides various transferrin fusion proteins that could be used in the treatment of a variety of diseases. For example, the pharmaceutical

compositions comprising the fusion polypeptides or peptides of the present invention could be used to treat diseases such as, but not limited, to insulin resistance, hyperglycemia, hyperinsulinemia, or elevated blood levels of free fatty acids or glycerol, hyperlipidemia, obesity, Syndrome X, dysmetabolic syndrome, inflammation, diabetic complications, impaired glucose homeostasis, impaired glucose tolerance, type II diabetes, prediabetes, hypertriglyceridemia atherosclerosis, nervous system disorders, congestive heart failure, dyspepsia, and irritable bowel syndrome. The modified polypeptides and peptides could also be used to induce an anxiolytic effect on the CNS, to activate the CNS or for post surgery treatment.

[00166] The modified therapeutic polypeptides and peptides of the present invention are more stable *in vivo* than the nonmodified therapeutic polypeptides and peptides because they are fused to transferrin or modified transferrin or are partially or substantially protected from DPP activity. Accordingly, smaller amounts of the molecule may be administered for effective treatment. A lower dosage amount may in some instances alleviate side effects.

[00167] In one embodiment, the modified therapeutic polypeptides and peptides of the present invention may be used as a sedative. Accordingly, the present invention provides a method of sedating a mammalian subject with an abnormality resulting in increased activation of the central or peripheral nervous system using the modified polypeptides or peptides of the invention. The method comprises administering a modified therapeutic polypeptides or peptides to the subject in an amount sufficient to produce a sedative or anxiolytic effect on the subject. The modified therapeutic polypeptides or peptides may be administered intracerebroventricularly, orally, subcutaneously, intramuscularly, or intravenously. Such methods are useful to treat or ameliorate nervous system conditions such as anxiety, movement disorder, aggression, psychosis, seizures, panic attacks, hysteria and sleep disorders.

[00168] Moreover, the present invention encompasses a method of increasing the activity of a mammalian subject, comprising administering a modified therapeutic polypeptides or peptides to the subject in an amount sufficient to produce an activating effect on the subject. The subject has a condition resulting in decreased activation of the central or peripheral nervous system. The modified therapeutic polypeptides or peptides are useful in the treatment or amelioration of depression, schizoaffective disorders, sleep apnea, attention deficit syndromes with poor concentration, memory loss, forgetfulness, and narcolepsy, to

name just a few conditions in which arousal of the central nervous system may be advantageous.

[00169] Also, insulin resistance following a particular type of surgery, elective abdominal surgery, is most profound on the first post-operative day, lasts at least five days, and may take up to three weeks to normalize. Thus, the post-operative patient may be in need of administration of the modified insulintropic peptides of the present invention for a period of time following the trauma of surgery. Accordingly, the modified therapeutic polypeptides or peptides of the invention may be utilized for post surgery treatments. A patient is in need of the modified insulintropic peptides of the present invention for about 1-16 hours before surgery is performed on the patient, during surgery on the patient, and after the patient's surgery for a period of not more than about 5 days.

[00170] Moreover, the modified therapeutic polypeptides and peptides, such as the insulintropic peptides, of the invention may be utilized to treat insulin resistance independently from their use in post surgery treatment. Insulin resistance may be due to a decrease in binding of insulin to cell-surface receptors, or to alterations in intracellular metabolism. The first type, characterized as a decrease in insulin sensitivity, can typically be overcome by increased insulin concentration. The second type, characterized as a decrease in insulin responsiveness, cannot be overcome by large quantities of insulin. Insulin resistance following trauma can be overcome by doses of insulin that are proportional to the degree of insulin resistance, and thus is apparently caused by a decrease in insulin sensitivity.

[00171] Preferably, the present invention provides modified insulintropic peptides to normalize hyperglycemia through glucose-dependent, insulin-dependent and insulin-independent mechanisms. As such, the modified insulintropic peptides are useful as primary agents for the treatment of diabetes, especially type II diabetes mellitus. The present invention is especially suited for the treatment of patients with diabetes, both type I and type II, in that the action of the peptide is dependent on the glucose concentration of the blood, and thus the risk of hypoglycemic side effects are greatly reduced over the risks in using current methods of treatment

[00172] The dose of modified insulintropic peptides effective to normalize a patient's blood glucose level will depend on a number of factors, among which are included, without limitation, the patient's sex, weight and age, the severity of inability to regulate blood

glucose, the underlying causes of inability to regulate blood glucose, whether glucose, or another carbohydrate source, is simultaneously administered, the route of administration and bioavailability, the persistence in the body, the formulation, and the potency.

[00173] Preferably, the modified therapeutic peptides such as the insulintropic peptides, of the present invention are used for the treatment of impaired glucose tolerance, glycosuria, hyperlipidaemia, metabolic acidoses, diabetes mellitus, diabetic neuropathy, and nephropathy. More preferably, the modified peptides are modified GLP-1 and analogs thereof for the treatment of type II diabetes.

Monitoring the Presence of Modified Therapeutic Polypeptides and Peptides

[00174] The modified therapeutic polypeptides and peptides may be monitored using assays for determining functional activity, HPLC-MS, or antibodies directed against the polypeptide or peptide. For example, the blood of the mammalian host may be monitored for the activity of the modified therapeutic polypeptide or peptide and/or presence of the modified therapeutic polypeptide or peptide. By taking a portion or sample of the blood of the host at different times, one may determine whether the modified therapeutic polypeptide or peptide has become bound to the long-lived blood components in sufficient amount to be therapeutically active and, thereafter, the level of modified therapeutic polypeptide or peptide in the blood. If desired, one may also determine to which of the blood components the modified therapeutic polypeptide or peptide, such as a modified insulintropic peptide, is bound.

[00175] As an example, assays for insulintropic activity may be used to monitor the modified insulintropic peptides of the present invention. The modified insulintropic peptides of the present invention have an insulintropic activity that at least equals the insulintropic activity of the non-modified insulintropic peptides. The insulintropic property of a modified insulintropic peptide may be determined by providing that modified peptide to animal cells, or injecting that peptide into animals and monitoring the release of immunoreactive insulin into the media or circulatory system of the animal, respectively. The presence of immunoreactive insulin is detected through the use of a radioimmunoassay which can specifically detect insulin. Although any radioimmunoassay capable of detecting the presence of IRI may be employed, it is preferable to use a modification of the assay

method of Albano, J. D. M., *et al.*, (1972 *Acta Endocrinol.* 70:487-509), which is herein incorporated by reference in its entirety.

[00176] The insulintropic property of a modified therapeutic polypeptide or peptide may also be determined by pancreatic infusion (Penhos, J. C., *et al.* 1969 *Diabetes* 18:733-738, which is hereby incorporated by reference). The manner in which perfusion is performed, modified, and analyzed preferably follows the methods of Weir, G. C., *et al.*, (*J. Clin. Investigat.* 54:1403-1412 (1974)), which is hereby incorporated by reference.

[00177] HPLC coupled with mass spectrometry (MS) can be utilized to assay for the presence of modified therapeutic polypeptide and peptides as is well known to the skilled artisan. Typically two mobile phases are utilized, such as 0.1% TFA/water and 0.1% TFA/acetonitrile. Column temperatures can be varied as well as gradient conditions.

[00178] Another method to monitor the presence of modified therapeutic polypeptides and peptides is to use antibodies specific to the modified therapeutic polypeptides and peptides. The use of antibodies, either monoclonal or polyclonal, having specificity for particular modified therapeutic polypeptides or peptides, can assist in mediating any such problem. The antibody may be generated or derived from a host immunized with the particular modified therapeutic polypeptide or peptide, or with an immunogenic fragment of the agent, or a synthesized immunogen corresponding to an antigenic determinant of the agent. Preferred antibodies will have high specificity and affinity for the modified therapeutic polypeptide or peptide. Such antibodies can also be labeled with enzymes, fluorochromes, or radiolabels.

[00179] The antibodies may be used to monitor the presence of modified therapeutic polypeptides and peptides in the blood stream. Blood and/or serum samples may be analyzed by SDS-PAGE and western blotting. Such techniques permit the analysis of the blood or serum to determine the bonding of the modified therapeutic polypeptides or peptides to blood components.

Glucagon-Like Peptide-1 (GLP-1)

[00180] Recombinant DNA technology has been used to create new molecules with increased the stability and biological activity. These molecules are combinations of biologically active proteins and peptides fused to a stabilizing protein with naturally long

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half-life such as immunoglobulin Fc portion, albumin and transferrin. These fusion molecules retain the biological activity of the active moiety with much longer pharmacokinetics than their natural unfused protein or peptide counterparts. Increase in the pharmacokinetics also improves the biological activity, reduces unwanted side effects and improves convenience to the patients. There are many examples of such fusion proteins like interferon-albumin, interferon-Fc, BNP-albumin, GLP-1-albumin, GLP-1-Transferrin,

[00181] Although the fusion proteins are stable and resistant to degradation, the underlying protease mechanism that degrades the active moiety may result in a slow inactivation of the molecule. Specifically, many peptides such as GLP-1, dynorphin (Berman YL, Juliano L, Devi LA *J Biol Chem.* 1995 Oct 6;270:23845-50), enkephalin (Gu ZF, Menozzi D, Okamoto A, Maton PN, Bunnett NW *Exp Physiol.* 1993 Jan;78:35-48), BNP, ANP, angiotensin, bradykinin, and PYY are very susceptible to proteases such as dipeptidyl-peptidase IV, neutral endopeptidase. These proteases individually or in combination cause rapid inactivation of the peptides in the circulation. The fusion of peptides to large proteins such as albumin, Fc and transferrin confers a significant resistance to proteases. However, it may not totally eliminate the effect of the inactivation by proteases. Therefore, combination of the fusion proteins and protease inhibitors can have better PK and PD than the fusion protein alone.

[00182] The present invention provides transferrin fusion proteins comprising therapeutic peptides that are resistant to proteases. Preferably, the modified therapeutic peptides of the present invention are modified insulinotropic peptides partially or substantially protected from DPP activity. More preferably, the modified insulinotropic peptides are modified GLP-1 peptides and analogs and fragments thereof. The modified GLP-1 peptides and analogs and fragments thereof are useful for treating diabetes, specifically type II diabetes. The N-terminal sequence of wild-type GLP-1 is His-Ala-Glu; modified GLP-1 polypeptides of the invention may comprise an N-terminal sequence selected from the group consisting of: His-His-Ala-Glu (SEQ ID NO: 115), Gly-His-Ala-Glu (SEQ ID NO: 116), His-Gly-Glu, His-Ser-Glu, His-Ala-Glu, His-Gly-Glu, His-Ser-Glu, His-His-Ala-Glu (SEQ ID NO: 82), His-His-Gly-Glu (SEQ ID NO: 83), His-His-Ser-Glu (SEQ ID NO: 84), Gly-His-Ala-Glu (SEQ ID NO: 85), Gly-His-Gly-Glu (SEQ ID NO: 86), Gly-His-Ser-Glu (SEQ ID NO: 87), His-X-Ala-Glu, His-X-Gly-Glu, and His-X-Ser-Glu, wherein X is any amino acid. As described below, other modifications may be made to decrease and prevent protease

degradation and these molecules may be used in the methods and compositions of the invention. Further, any GLP-1 moiety or GLP-1 analogs, derivatives, or mimetic may be used (as a fusion protein) in the methods and compositions of the invention.

[00183] The addition of an amino acid to the N-terminus of GLP-1 may prevent dipeptidyl peptidase from cleaving at the second amino acid of GLP-1 due to steric hindrance. Therefore, GLP-1 will remain functionally active. Any one of the 20 amino acids or a non-natural amino acid may be added to the N-terminus of GLP-1. Histidine is also a preferred amino acid. In some instances, an uncharged or positively charged amino acid may be used and preferably, a smaller amino acid such as Glycine is added. The modified GLP-1 with the extra amino acid can then be fused to transferrin to make a fusion protein. In one embodiment, the GLP-1 peptide is modified to contain at least one additional amino acid at its amino terminus. In another embodiment, the GLP-1 peptide is modified to contain at least five additional amino acids at its amino terminus. Alternatively, the GLP-1 peptide is modified to contain between one and five additional amino acids at its amino terminus.

[00184] Glucagon-Like Peptide-1 (GLP-1) is a gastrointestinal hormone that regulates insulin secretion belonging to the so-called enteroinsular axis. The enteroinsular axis designates a group of hormones, released from the gastrointestinal mucosa in response to the presence and absorption of nutrients in the gut, which promote an early and potentiated release of insulin. The incretin effect which is the enhancing effect on insulin secretion is probably essential for a normal glucose tolerance. GLP-1 is a physiologically important insulinotropic hormone because it is responsible for the incretin effect.

[00185] GLP-1 is a product of the proglucagon gene (Bell, *et al.*, Nature, 1983, 304: 368-371). It is synthesized in intestinal endocrine cells in two principal major molecular forms, as GLP-1(7-36)amide and GLP-1(7-37). The peptide was first identified following the cloning of cDNAs and genes for proglucagon in the early 1980s.

[00186] Initial studies done on the full length peptide GLP-1(1-37 and 1-36^{amide}) concluded that the larger GLP-1 molecules are devoid of biological activity. In 1987, three independent research groups demonstrated that removal of the first six amino acids resulted in a GLP-1 molecule with enhanced biological activity.

[00187] The amino acid sequence of GLP-1 is disclosed by Schmidt *et al.* (1985 Diabetologia 28 704-707). Human GLP-1 is a 37 amino acid residue peptide originating

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from preproglucagon which is synthesized in the L-cells in the distal ileum, in the pancreas, and in the brain. Processing of preproglucagon to GLP-1(7-36)amide, GLP-1(7-37) and GLP-2 occurs mainly in the L-cells. The amino acid sequence of GLP-1(7-37) is SEQ ID NO: 32 (X = Gly):

His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-Gly.

In GLP-1(7-36)amide, the terminal Gly is replaced by NH₂.

[00188] GLP-1 like molecules possesses anti-diabetic activity in human subjects suffering from Type II (non-insulin-dependent diabetes mellitus (NIDDM)) and, in some cases, even Type I diabetes. Treatment with GLP-1 elicits activity, such as increased insulin secretion and biosynthesis, reduced glucagon secretion, delayed gastric emptying, only at elevated glucose levels, and thus provides a potentially much safer therapy than insulin or sulfonylureas. Post-prandial and glucose levels in patients can be moved toward normal levels with proper GLP-1 therapy. There are also reports suggesting GLP-1-like molecules possess the ability to preserve and even restore pancreatic beta cell function in Type-II patients.

[00189] Any GLP-1 sequence may be modified by adding one or more amino acids at its amino terminus, including GLP-1(7-34), GLP-1(7-35), GLP-1(7-36), and GLP-1(7-37). GLP-1 also has powerful actions on the gastrointestinal tract. Infused in physiological amounts, GLP-1 potently inhibits pentagastrin-induced as well as meal-induced gastric acid secretion (Schjoldager *et al.*, *Dig. Dis. Sci.* 1989, 35:703-708; Wettergren *et al.*, *Dig Dis Sci* 1993; 38:665-673). It also inhibits gastric emptying rate and pancreatic enzyme secretion (Wettergren *et al.*, *Dig Dis Sci* 1993; 38:665-673). Similar inhibitory effects on gastric and pancreatic secretion and motility may be elicited in humans upon perfusion of the ileum with carbohydrate- or lipid-containing solutions (Layer *et al.*, *Dig Dis Sci* 1995, 40:1074-1082; Layer *et al.*, *Digestion* 1993, 54: 385-38). Concomitantly, GLP-1 secretion is greatly stimulated, and it has been speculated that GLP-1 may be at least partly responsible for this so-called "ileal-brake" effect (Layer *et al.*, *Digestion* 1993; 54: 385-38). In fact, recent studies suggest that, physiologically, the ileal-brake effects of GLP-1 may be more important than its effects on the pancreatic islets. Thus, in dose response studies GLP-1 influences gastric emptying rate at infusion rates at least as low as those required to influence islet secretion (Nauck *et al.*, *Gut* 1995; 37 (suppl. 2): A124).

[00190] GLP-1 seems to have an effect on food intake. Intraventricular administration of GLP-1 profoundly inhibits food intake in rats (Schick *et al.* in Ditschuneit *et al.* (eds.), Obesity in Europe, John Libbey & Company Ltd, 1994; pp. 363-367; Turton *et al.*, Nature 1996, 379: 69-72). This effect seems to be highly specific. Thus, N-terminally extended GLP-1(1-36^{amide}) is inactive and appropriate doses of the GLP-1 antagonist, exendin 9-39, abolish the effects of GLP-1 (Tang-Christensen *et al.*, Am. J. Physiol., 1996, 271(4 Pt 2):R848-56). Acute, peripheral administration of GLP-1 does not inhibit food intake acutely in rats (Tang-Christensen *et al.*, Am. J. Physiol., 1996, 271(4 Pt 2):R848-56; Turton *et al.*, Nature 1996, 379: 69-72). However, it remains possible that GLP-1 secreted from the intestinal L-cells may also act as a satiety signal.

[00191] In diabetic patients, GLP-1's insulintropic effects and the effects of GLP-1 on the gastrointestinal tract are preserved (Willms *et al.*, Diabetologia 1994; 37, suppl.1: A118), which may help curtail meal-induced glucose excursions, but, more importantly, may also influence food intake. Administered intravenously, continuously for one week, GLP-1 at 4 ng/kg/min has been demonstrated to dramatically improve glycaemic control in NIDDM patients without significant side effects (Larsen *et al.*, Diabetes 1996; 45, suppl. 2: 233A.).

[00192] Modified GLP-1 partially or substantially protected from DPP activity and modified GLP-1 analogs are useful in the treatment of Type 1 and Type 2 diabetes and obesity.

[00193] As used herein, the term "GLP-1 molecule" means GLP-1, a GLP-1 analog, or GLP-1 derivative.

[00194] As used herein, the term "GLP-1 analog" is defined as a molecule having one or more amino acid substitutions, deletions, inversions, or additions compared with GLP-1. Many GLP-1 analogs are known in the art and include, for example, GLP-1(7-34), GLP-1(7-35), GLP-1(7-36), Val⁸-GLP-1(7-37), Gln⁹-GLP1(7-37), D-Gln⁹-GLP-1(7-37), Thr¹⁶-Lys¹⁸-GLP-1(7-37), and Lys¹⁸-GLP-1(7-37) (SEQ ID NO: 72). U.S. Patent 5,118,666 discloses examples of GLP-1 analogs such as GLP-1(7-34) and GLP-1(7-35).

[00195] The term "GLP-1 derivative" is defined as a molecule having the amino acid sequence of GLP-1 or a GLP-1 analog, but additionally having chemical modification of one or more of its amino acid side groups, α -carbon atoms, terminal amino group, or

terminal carboxylic acid group. A chemical modification includes, but is not limited to, adding chemical moieties, creating new bonds, and removing chemical moieties.

[00196] As used herein, the term "GLP-1 related compound" refers to any compound falling within the GLP-1, GLP-1 analog, or GLP-1 derivative definition.

[00197] WO 91/11457 discloses analogs of the active GLP-1 peptides 7-34, 7-35, 7-36, and 7-37 which can also be useful as GLP-1 moieties.

[00198] EP 0708179-A2 (Eli Lilly & Co.) discloses GLP-1 analogs and derivatives that include an N-terminal imidazole group and optionally an unbranched C₆-C₁₀ acyl group in attached to the lysine residue in position 34.

[00199] EP 0699686-A2 (Eli Lilly & Co.) discloses certain N-terminal truncated fragments of GLP-1 that are reported to be biologically active.

[00200] U.S. Patent 5,545,618 discloses GLP-1 molecules consisting essentially of GLP-1(7-34), GLP1(7-35), GLP-1(7-36), or GLP-1(7-37), or the amide forms thereof, and pharmaceutically-acceptable salts thereof, having at least one modification selected from the group consisting of: (a) substitution of glycine, serine, cysteine, threonine, asparagine, glutamine, tyrosine, alanine, valine, isoleucine, leucine, methionine, phenylalanine, arginine, or D-lysine for lysine at position 26 and/or position 34; or substitution of glycine, serine, cysteine, threonine, asparagine, glutamine, tyrosine, alanine, valine, isoleucine, leucine, methionine, phenylalanine, lysine, or a D-arginine for arginine at position 36 (SEQ ID NO: 73); (b) substitution of an oxidation-resistant amino acid for tryptophan at position 31 (SEQ ID NO: 74); (c) substitution of at least one of: tyrosine for valine at position 16; lysine for serine at position 18; aspartic acid for glutamic acid at position 21; serine for glycine at position 22; arginine for glutamine at position 23; arginine for alanine at position 24; and glutamine for lysine at position 26 (SEQ ID NO: 75); and (d) substitution of at least one of: glycine, serine, or cysteine for alanine at position 8; aspartic acid, glycine, serine, cysteine, threonine, asparagine, glutamine, tyrosine, alanine, valine, isoleucine, leucine, methionine, or phenylalanine for glutamic acid at position 9; serine, cysteine, threonine, asparagine, glutamine, tyrosine, alanine, valine, isoleucine, leucine, methionine, or phenylalanine for glycine at position 10; and glutamic acid for aspartic acid at position 15 (SEQ ID NO: 76); and (e) substitution of glycine, serine, cysteine, threonine, asparagine, glutamine, tyrosine, alanine, valine, isoleucine, leucine, methionine, or phenylalanine, or the

D- or N-acylated or alkylated form of histidine for histidine at position 7 (SEQ ID NO: 77); wherein, in the substitutions is (a), (b), (d), and (e), the substituted amino acids can optionally be in the D-form and the amino acids substituted at position 7 can optionally be in the N-acylated or N-alkylated form.

[00201] U.S. Pat. No. 5,118,666 discloses a GLP-1 molecule having insulinotropic activity. Such molecule is selected from the group consisting of a peptide having the amino acid sequence His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys (GLP-1, 7-34, see SEQ ID NO: 32) or His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly (GLP-1, 7-35, see SEQ ID NO: 32); and a derivative of said peptide and wherein said peptide is selected from the group consisting of: a pharmaceutically-acceptable acid addition salt of said peptide; a pharmaceutically-acceptable carboxylate salt of said peptide; a pharmaceutically-acceptable lower alkylester of said peptide; and a pharmaceutically-acceptable amide of said peptide selected from the group consisting of amide, lower alkyl amide, and lower dialkyl amide.

[00202] U.S. Patent 6,277,819 teaches a method of reducing mortality and morbidity after myocardial infarction comprising administering GLP-1, GLP-1 analogs, and GLP-1 derivatives to the patient. The GLP-1 analog being represented by the following structural formula (SEQ ID NO: **): R_1 - X_1 -Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu- X_2 -Gly-Gln-Ala-Ala-Lys- X_3 -Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg- R_2 (SEQ ID NO: 78) and pharmaceutically-acceptable salts thereof, wherein: R_1 is selected from the group consisting of L-histidine, D-histidine, desamino-histidine, 2-amino-histidine, .beta.-hydroxy-histidine, homohistidine, alpha-fluoromethyl-histidine, and alpha-methyl-histidine; X_1 is selected from the group consisting of Ala, Gly, Val, Thr, Ile, and alpha-methyl-Ala; X_2 is selected from the group consisting of Glu, Gln, Ala, Thr, Ser, and Gly; X_3 is selected from the group consisting of Glu, Gln, Ala, Thr, Ser, and Gly; R_2 is selected from the group consisting of NH_2 , and Gly--OH; provided that the GLP-1 analog has an isoelectric point in the range from about 6.0 to about 9.0 and further providing that when R_1 is His, X_1 is Ala, X_2 is Glu, and X_3 is Glu, R_2 must be NH_2 .

[00203] Ritzel *et al.* (Journal of Endocrinology, 1998, 159: 93-102) disclose a GLP-1 analog, [Ser⁸]GLP-1, in which the second N-terminal alanine is replaced with serine. The

modification did not impair the insulinotropic action of the peptide but produced an analog with increased plasma stability as compared to GLP-1.

[00204] U.S. Patent 6,429,197 teaches that GLP-1 treatment after acute stroke or hemorrhage, preferably intravenous administration, can be an ideal treatment because it provides a means for optimizing insulin secretion, increasing brain anabolism, enhancing insulin effectiveness by suppressing glucagon, and maintaining euglycemia or mild hypoglycemia with no risk of severe hypoglycemia or other adverse side effects. The present invention provides a method of treating the ischemic or reperfused brain with GLP-1 or its biologically active analogues after acute stroke or hemorrhage to optimize insulin secretion, to enhance insulin effectiveness by suppressing glucagon antagonism, and to maintain euglycemia or mild hypoglycemia with no risk of severe hypoglycemia.

[00205] U.S. Patent 6,277,819 provides a method of reducing mortality and morbidity after myocardial infarction, comprising administering to a patient in need thereof, a compound selected from the group consisting of GLP-1, GLP-1 analogs, GLP-1 derivatives and pharmaceutically-acceptable salts thereof, at a dose effective to normalize blood glucose.

[00206] U.S. Patent 6,191,102 discloses a method of reducing body weight in a subject in need of body weight reduction by administering to the subject a composition comprising a glucagon-like peptide-1 (GLP-1), a glucagon-like peptide analog (GLP-1 analog), a glucagon-like peptide derivative (GLP-1 derivative) or a pharmaceutically acceptable salt thereof in a dose sufficient to cause reduction in body weight for a period of time effective to produce weight loss, said time being at least 4 weeks.

[00207] GLP-1 is fully active after subcutaneous administration (Ritzel *et al.*, *Diabetologia* 1995; 38: 720-725), but is rapidly degraded mainly due to degradation by dipeptidyl peptidase IV-like enzymes (Deacon *et al.*, *J Clin Endocrinol Metab* 1995, 80: 952-957; Deacon *et al.*, 1995, *Diabetes* 44: 1126-1131). Thus, unfortunately, GLP-1 and many of its analogues have a short plasma half-life in humans (Orskov *et al.*, *Diabetes* 1993; 42:658-661). Accordingly, it is an objective of the present invention to provide modified GLP-1 or analogues thereof which have a protracted profile of action relative to GLP-1(7-37). It is a further object of the invention to provide derivatives of GLP-1 and analogues thereof which have a lower clearance than GLP-1(7-37). Moreover, it is an object of the invention to provide pharmaceutical compositions comprising modified GLP-1 or GLP-1 analogs with improved stability. Additionally, the present invention includes the use of modified GLP-1

or GLP-1 analogs to treat diseases associated with GLP-1 such as but not limited to those described above.

[00208] In one aspect of the present invention, the pharmaceutical compositions comprising modified GLP-1 and GLP-1 analogs may be formulated by any of the established methods of formulating pharmaceutical compositions, *e.g.* as described in Remington's Pharmaceutical Sciences, 1985. The composition may be in a form suited for systemic injection or infusion and may, as such, be formulated with a suitable liquid vehicle such as sterile water or an isotonic saline or glucose solution. The compositions may be sterilized by conventional sterilization techniques which are well known in the art. The resulting aqueous solutions may be packaged for use or filtered under aseptic conditions and lyophilized, the lyophilized preparation being combined with the sterile aqueous solution prior to administration. The composition may contain pharmaceutically acceptable auxiliary substances as required to approximate physiological conditions, such as buffering agents, tonicity adjusting agents and the like, for instance sodium acetate, sodium lactate, sodium chloride, potassium chloride, calcium chloride, *etc.*

[00209] The modified GLP-1 and GLP-1 analogs of the present invention may also be adapted for nasal, transdermal, pulmonal or rectal administration. The pharmaceutically acceptable carrier or diluent employed in the composition may be any conventional solid carrier. Examples of solid carriers are lactose, terra alba, sucrose, talc, gelatin, agar, pectin, acacia, magnesium stearate and stearic acid. Similarly, the carrier or diluent may include any sustained release material known in the art, such as glyceryl monostearate or glyceryl distearate, alone or mixed with a wax.

[00210] It may be of particular advantage to provide the composition of the invention in the form of a sustained release formulation. As such, the composition may be formulated as microcapsules or microparticles containing the modified GLP-1 or GLP-1 analogs encapsulated by or dispersed in a suitable pharmaceutically acceptable biodegradable polymer such as polylactic acid, polyglycolic acid or a lactic acid/glycolic acid copolymer.

[00211] For nasal administration, the preparation may contain modified GLP-1 or GLP-1 analogs dissolved or suspended in a liquid carrier, in particular an aqueous carrier, for aerosol application. The carrier may contain additives such as solubilizing agents, *e.g.* propylene glycol, surfactants, absorption enhancers such as lecithin (phosphatidylcholine) or cyclodextrin, or preservatives such as parabenes.

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[00212] Generally, the modified polypeptides or peptides of the present invention are dispensed in unit dosage form together with a pharmaceutically acceptable carrier per unit dosage.

[00213] Moreover, the present invention contemplates the use of the modified GLP-1 and GLP-1 analogs for the manufacture of a medicinal product which can be used in the treatment of diseases associated with elevated glucose level (metabolic disease), such as but not limited to those described above. Specifically, the present invention contemplates the use of modified GLP-1 and GLP-1 analogs for the treatment of diabetes including type II diabetes, obesity, severe burns, and heart failure, including congestive heart failure and acute coronary syndrome.

[00214] The present invention also provides modified Exendin-3 and Exendin-4 peptides partially and substantially protected from DPP activity. Exendin-3 and Exendin-4 are insulinotropic peptides comprising 39 amino acids (differing at residues 2 and 3) which are approximately 53% homologous to GLP-1. The Exendin-3 sequence is HSDGTFTSDLSKQMEEEEAVRLFIEWLKNNGG PSSGAPPPS (SEQ ID NO: 79), and the Exendin-4 sequence is HEGTFTSDLSKQMEEEEAVRLFIEWLKNNGG PSSGAPPPS (SEQ ID NO: 80). The invention also encompasses the modified exendin-4 fragments comprising the amino acid sequences such as Exendin-4 (1-31) HEGTFTSDLSKQMEEAVR LFIIEWLKNNGGPY (SEQ ID NO: 81). Additionally, the present invention includes modified analogs of Exendin-3 and Exendin-4 peptides.

Modified GLP-1 Fusion Protein or Conjugate for Treating Diabetes, Prediabetes, or Obesity

[00215] The modified GLP-1 may be fused to a heterologous molecule for increased overall stability *in vivo*. The modified GLP-1 may be fused to a heterologous molecule by recombinant means or covalently attached to a heterologous molecule by methods well known in the art. Modified GLP-1 may be fused or covalently attached, for example to a plasma protein such as serum albumin or transferrin, an immunoglobulin, or a portion thereof such as the Fc domain. More preferably, the modified polypeptide or peptide is fused to transferrin, lactotransferrin, or melanotransferrin. Methods for making such fusion proteins are provided by U.S. Application 10/378,094, which is herein incorporated by reference in its entirety.

[00216] The GLP-1 molecule may be attached to the heterologous protein via a linker of variable length to provide greater physical separation and allow more spatial mobility between the fused proteins and thus maximize the accessibility of the therapeutic protein, for instance, for binding to its cognate receptor. The linker peptide may consist of amino acids that are flexible or more rigid. For example, a linker such as a poly-glycine stretch may be used. The linker can be less than about 50, 40, 30, 20, 10, or 5 amino acid residues. The linker can be covalently linked to and between the heterologous protein and GLP-1. Preferably, the linker may be one Ser residue, two Ser residues, the peptide Ser-Ser-Gly, the peptide PEAPTD, the peptide (PEAPTD)₂, the peptide PEAPTD in combination with IgG hinge linker, and the peptide (PEAPTD)₂ in combination with IgG hinge linker. These linkers may be used to link GLP-1 to transferrin.

[00217] The transferrin to be attached to the modified polypeptide or peptide may be modified. It may exhibit reduced glycosylation. The modified transferrin polypeptide may be selected from the group consisting of a single transferrin N domain, a single transferrin C domain, a transferrin N and C domain, two transferrin N domains, and two transferrin C domains.

[00218] As discussed above, GLP-1 activates and regulates important endocrine hormone systems in the body and plays a critical management role in the metabolism of glucose. Unlike all other diabetic treatments on the market GLP-1 has the potential to be restorative by acting as a growth factor for β -cells thus improving the ability of the pancreas to secrete insulin and also, to make the existing insulin levels act more efficiently by improving sensitivity and better stabilizing glucose levels. This reduces the burden on daily monitoring of glucose levels and potentially offers a delay in the serious long term side effects caused by fluctuations in blood glucose due to diabetes. Furthermore, GLP-1 can reduce appetite and reduce weight. Obesity is an inherent consequence of poor control of glucose metabolism and this only serves to aggravate the diabetic condition.

[00219] Clinical application of natural GLP-1 is limited because it is rapidly degraded in the circulation (half-life is several minutes). To maintain therapeutic levels in the circulation requires constant administration of high doses using pumps or patch devices which adds to the cost of treatment. This is inconvenient for long term chronic use especially in conjunction with all the other medications for treating diabetes and monitoring of glucose levels. The modified GLP-1 fusion proteins retain the activity of GLP-1 but

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have the long half-life (14-17 days), solubility, and biodistribution properties of transferrin. These properties could provide for a low cost, small volume, monthly s.c. (subcutaneous) injection and this type of product is absolutely needed for long term chronic use.

[00220] The modified GLP-1 also may be covalently attached to a blood component to increase its stability. For example, the modified GLP-1 may be covalently attached to serum albumin, transferrin, immunoglobulin, or the Fc portion of the immunoglobulin. In one embodiment, the modified GLP-1 may be attached to a fatty acid or a fatty acid derivative. In another embodiment, the modified GLP-1 may be engineered into a drug affinity complex (DAC). As discussed earlier, Kim *et al.* (2003, Diabetes 52(3):751) disclose a GLP-1-albumin drug affinity complex. Kim *et al.* show that the albumin-conjugated DAC:GLP-1 mimics the native GLP-1. Kim *et al.* provide a new approach for prolonged activation of GLP-1R signaling.

[00221] Upon subcutaneous administration, the DAC:modified GLP-1 rapidly and selectively bonds *in vivo* to albumin. The bioconjugate formed has the same therapeutic activity and similar potency as endogenous GLP-1 but has a pharmacokinetic profile that is closer to albumin.

Modified GLP-1 and its Fusion Protein in Combination with Other Therapeutic Agents

[00222] In one aspect of the invention, the modified GLP-1 peptide and its fusion protein, for example, GLP-1-Tf fusion protein, of the present invention are used in combination with at least one second therapeutic molecule such as Glucophage® (metformin hydrochloride tablets) or Glucophage® XR (metformin hydrochloride extended-release tablets) to treat type II diabetes, obesity, and other diseases or conditions associated with abnormal glucose levels.

[00223] Glucophage® and Glucophage® XR are oral antihyperglycemic drugs for the management of type II diabetes. Glucophage® XR is an extended release formulation of Glucophage. Accordingly, Glucophage® XR may be taken once daily because the drug is released slowly from the dosage form. Glucophage® helps the body produce less glucose from the liver. Accordingly, Glucophage® is effective in controlling blood sugar level in a

patient. Glucophage® rarely causes low blood glucose (hypoglycemia) because it does not cause the body to make more insulin.

[00224] Glucophage® also helps lower the fatty blood components, triglycerides and cholesterol, that are often high in people with Type II diabetes. Metformin has been shown to decrease the appetite and help people lose a few pounds when they starting taking the medicine.

[00225] Metformin has been approved for treatment with sulfonylureas, or with insulin, or as monotherapy (by itself). Metformin has been suggested for use in treating various cardiovascular diseases such as hypertension in insulin resistant patients (WO 9112003-Upjohn), for dissolving blood clots (in combination with a t-PA-derivative) (WO 9108763, WO 9108766, WO 9108767 and WO 9108765-Boehringer Mannheim), ischemia and tissue anoxia (EP 283369-Lipha), atherosclerosis (DE 1936274-Brunnengraber & Co., DE 2357875-Hurka, and U.S. Pat. No. 4,205,087-ICI). In addition, it has been suggested to use metformin in combination with prostaglandin-analogous cyclopentane derivatives as coronary dilators and for blood pressure lowering (U.S. Pat. No. 4,182,772-Hoechst). Metformin has also been suggested for use in cholesterol lowering when used in combination with 2-hydroxy-3,3,3-trifluoropropionic acid derivatives (U.S. Pat. No. 4,107,329-ICI), 1,2-diarylethylene derivatives (U.S. Pat. No. 4,061,772-Hoechst), substituted aryloxy-3,3,3-trifluoro-2-propionic acids, esters and salts (U.S. Pat. No. 4,055,595-ICI), substituted hydroxyphenyl-piperidones (U.S. Pat. No. 4,024,267-Hoechst), and partially hydrogenated 1H-indeno-[1,2B]-pyridine derivatives (U.S. Pat. No. 3,980,656-Hoechst).

[00226] Montanari *et al.* (Pharmacological Research, Vol. 25, No. 1, 1992) disclose that use of metformin in amounts of 500 mg twice a day (b.i.d.) increased post-ischemia blood flow in a manner similar to 850 mg metformin three times a day (t.i.d.). Sirtori *et al.* (J. Cardiovas. Pharm., 6:914-923, 1984), disclose that metformin in amounts of 850 mg three times a day (t.i.d) increased arterial flow in patients with peripheral vascular disease.

[00227] The present invention provides the treatment of various diseases comprising modified GLP-1 of the present invention or its fusion protein in combination with one or more therapeutic agents such as metformin. In one embodiment, the modified GLP-1 or its fusion protein in combination with metformin is used to treat diseases and conditions

associated with abnormal blood glucose level, such as diabetes. Preferably, the GLP-1/mTf fusion protein in combination with metformin is used to treat type II diabetes or obesity.

[00228] Other therapeutic agents that may be used in combination with modified GLP-1 of the present invention and its fusion proteins include but are not limited to sulfonylurea and sulfonylurea-like agents, thiazolidinediones, Peroxisome Proliferator-Activated Receptor (PPAR) gamma modulators, PPAR alpha modulators, Protein Tyrosine Phosphatase-1B inhibitors, Insulin Receptor Tyrosine Kinase activators, 11beta-hydroxysteroid dehydrogenase inhibitors, glycogen phosphorylase inhibitors, glucokinase activators, beta-3 adrenergic agonists, and glucagon receptor agonists.

DPP-IV Inhibitors

[00229] Inhibitors of DPP-IV have been shown to be promising in treating various conditions mediated by DPP-IV. For example, inhibitors of DPP-IV are an extremely promising approach in the treatment of glucose intolerance and in disorders associated with hyperglycemia, such as type II diabetes or obesity. Moreover, DPP-IV has been shown to play a part in the immune response, such as transplant rejection (Transplantation 1997, 63(10):1495-1500). Accordingly, DPP-IV may be useful in the prevention of transplant rejection. Also, inhibitors of DPP-IV may be useful in the treatment of cancer and the prevention of cancerous metastases, since the binding of endothelial DPP-IV of the lung to fibronectin of cancerous cells promotes metastasis of those cells (J. Biol. Chem. 1998, 273(37):24207-24215). DPP-IV is likewise thought to play an important part in the pathogenesis of periodontitis (Infect. Immun. 2000, 68 (2), 716-724) and to be responsible for the inactivation of GLP-2, a factor facilitating recovery of the intestine after major resection (J. Surg. Res. 1999, 87 (1), 130-133). Accordingly, DPP-IV-inhibitors also are potentially useful in recovery of the intestine.

[00230] WO 95/15309 discloses certain peptide derivatives which are inhibitors of DPP-IV and, therefore, are useful in treating a number of DPP-IV mediated processes. WO 95/13069 discloses certain cyclic amine compounds which are useful in stimulating the release of natural or endogenous growth hormone. European Patent 555,824 discloses certain benzimidazolyl compounds which prolong thrombin time and inhibit thrombin and serine-related proteases. Archives of Biochemistry and Biophysics, Vol. 323, No. 1, pgs. 148-154 (1995) discloses certain aminoacylpyrrolidine-2-nitriles which are useful as DPP-IV inhibitors. Journal of Neurochemistry, Vol. 66, pgs. 2105-2112 (1996) discloses certain

Fmoc-aminoacylpyrrolidine-2-nitriles which are useful in inhibiting prolyl oligopeptidase. Bulletin of the Chemical Society of Japan, Vol. 50, No. 7, pgs. 1827-1830 (1977) discloses the synthesis of an aminohexapeptide, viz., Z-Val-Val-ImpPro-Gly-Phe-Phe-OMe, and its related aminopeptides. In addition, the antimicrobial properties of said compounds were examined. WO 90/12005 discloses certain amino acid compounds which inhibit prolylendopeptidase activity and, therefore, are useful in treating dementia or amnesia. Derwent Abstract 95: 302548 discloses certain N-(aryl(alkyl)carbonyl) substituted heterocyclic compounds which are cholinesterase activators with enhanced peripheral selectivity useful in treating conditions due to the lowering of cholinesterase activity. Chemical Abstracts 84: 177689 discloses certain 1-acyl-pyrrolidine-2-carbonitrile compounds which are useful as intermediates for proline compounds exhibiting angiotensin converting enzyme (ACE) inhibiting activity. Chemical Abstracts 96: 116353 discloses certain 3-amino-2-mercapto-propyl-proline compounds which are Ras farnesyl-transferase inhibitors useful in treating various carcinomas or myeloid leukemias. WO 95/34538 discloses certain pyrrolidides, phosphonates, azetidines, peptides and azaprolines which inhibit DPP-IV and, therefore, are useful in treating conditions ameliorated by DPP-IV inhibition. WO 95/29190 discloses certain compounds characterized by a plurality of KPR-type repeat patterns carried by a peptide matrix enabling their multiple presentation to, and having an affinity for, the enzyme DPP-IV, which compounds exhibit the ability to inhibit the entry of HIV into cells. WO 91/16339 discloses certain tetrapeptide boronic acids which are DPP-IV inhibitors useful in treating autoimmune diseases and conditions mediated by IL-2 suppression. WO 93/08259 discloses certain polypeptide boronic acids which are DPP-IV inhibitors useful in treating autoimmune diseases and conditions mediated by IL-2 suppression. WO 95/11689 discloses certain tetrapeptide boronic acids which are DPP-IV inhibitors useful in blocking the entry of HIV into cells. East German Patent 158109 discloses certain N-protected peptidyl-hydroxamic acids and nitrobenzoyloxamides which are useful as, inter alia, DPP-IV inhibitors. WO 95/29691 discloses, inter alia, certain dipeptide proline phosphonates which are DPP-IV inhibitors useful in the treatment of immune system disorders. German Patent DD 296075 discloses certain amino acid amides which inhibit DPP-IV. Biochimica et Biophysica Acta, Vol. 1293, pgs. 147-153 discloses the preparation of certain di- and tri-peptide p-nitroanilides to study the influence of side chain modifications on their DPP-IV and PEP-catalyzed hydrolysis. Bioorganic and Medicinal Chemistry Letters, Vol. 6, No. 10, pgs. 1163-1166

(1996) discloses certain 2-cyanopyrrolidines which are inhibitors DPP-IV. J. Med. Chem., Vol. 39, pgs. 2087-2094 (1996) discloses certain prolineboronic acid-containing dipeptides which are inhibitors of DPP-IV. Diabetes, Vol. 44, pgs. 1126-1131 (September '96) is directed to a study which demonstrates that GLP-I amide is rapidly degraded when administered by subcutaneous or intravenous routes to diabetic and non-diabetic subjects.

[00231] U.S. Patent 6,727,261 provides pyrido[2,1-a]isoquinoline derivatives as novel DPP-IV inhibitors useful for the treatment and/or prophylaxis of diseases which are associated with DPP-IV, such as diabetes, particularly non-insulin dependent diabetes mellitus, and impaired glucose tolerance. These compounds are also useful in the treatment and/or prophylaxis of Bow1 disease, Colitis Ulcerosa, Morbus Crohn, obesity and/or metabolic syndrome.

[00232] U.S. Patent 6,716,843 provides alpha-amino acid sulphonyl compounds useful as inhibitors for DPP-IV.

[00233] U.S. Patent 6,645,995 discloses 2-substituted unsaturated heterocyclic compounds wherein a nitrogen atom in the heterocyclic ring is attached via an amide bond or a peptide bond to an amino acid or an amino acid derivative. These compounds are potent and selective inhibitors of DPP-IV, and are effective in treating conditions that may be regulated or normalized via inhibition of DPP-IV.

[00234] U.S. Patent 6,617,340 discloses N-(substituted glycy1)-pyrrolidines, and the use of said compounds in inhibiting dipeptidyl peptidase-IV. U.S. Patent 6124,305 discloses N-(substituted glycy1)-2-cyanopyrrolidines which inhibit DPP-IV. These compounds are effective in treating conditions mediated by DPP-IV.

[00235] Administration of the Novartis compound 1-[[[2-[(5-cyanopyridin-2-yl)amino]ethyl]amino]acetyl]-2-cyano-(S)-pyrrolidine (NVP DPP728) over a 4 week period to 93 patients with Type 2 diabetes (mean HbA_{1c} of 7.4%) reduced levels of plasma glucose, insulin, and HbA_{1c} over the 4 week study period (see Diabetes Care 2002, 25(5):869-875).

Combination Therapy Using Inhibitors of DPP-IV

[00236] In one aspect, the present invention provides the use of a transferrin fusion protein comprising a therapeutic protein, polypeptide, or peptide in combination with one or more inhibitors of DPP-IV for the treatment of various conditions. The present invention

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provides pharmaceutical compositions comprising the transferrin fusion protein and one or more inhibitors of DPP-IV. As disclosed in U.S. Application No. 10/378,094, which is herein incorporated by reference in its entirety, the transferrin to be attached to the therapeutic protein, polypeptide, or peptide may be modified. It may exhibit reduced glycosylation. The modified transferrin polypeptide may be selected from the group consisting of a single transferrin N domain, a single transferrin C domain, a transferrin N and C domain, two transferrin N domains, and two transferrin C domains. The therapeutic protein or peptide to be attached to transferrin may be in its native or modified form. Preferably, the transferrin fusion protein comprises GLP-1 as the therapeutic peptide, linked to a modified transferrin molecule, as described in U.S. Application No. 10/378,094. Moreover, the combination therapy of the present invention comprises GLP-1/mTf fusion protein, one or more inhibitors of DPP-IV, and another therapeutic molecule. Such a molecule may be Glucophage® or Glucophage® XR.

[00237] In another aspect, the present invention provides the use of a modified protein or peptide that is resistant to dipeptidyl protease cleavage or its fusion protein in combination with one or more DPP-IV inhibitors. The present invention discloses pharmaceutical compositions comprising the modified protein or peptide or its fusion protein in combination with one or more DPP-IV inhibitors. Preferably, the modified peptide is modified GLP-1 and the fusion protein is modified GLP-1/mTf protein. Further, the combination therapy of the present invention comprises modified GLP-1 or modified GLP-1/mTf protein, one or more inhibitors of DPP-IV, and another therapeutic molecule such as Glucophage® or Glucophage® XR.

[00238] DPP-IV inhibitors may be used in methods of the invention to treat any relevant disease. For instance, a GLP-1-transferrin fusion protein as herein described can be combined with a DPP-IV inhibitor, such as 1-[[[2-[(5-cyanopyridin-2-yl)amino]ethyl]amino]acetyl]-2-cyano-(S)-pyrrolidine (NVP DPP728) to treat prediabetes, diabetes, obesity, or a diabetic symptom. The therapeutic agents may be administered sequentially or concurrently.

[00239] The transferrin fusion protein may comprise the GLP-1(7-37) peptide (SEQ ID NO: 32) or the GLP-1(7-36) peptide (amino acids 1-30 of SEQ ID NO: 32). More preferably, the GLP-1(7-37) peptide or GLP-1(7-36) peptide comprises an A8 to G and K34 to A mutation. The transferrin protein may also comprise a linker between GLP-1(7-37)

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peptide or GLP-1(7-36) peptide and the transferrin molecule. Preferably, the linker is the (PEAPTD)₂ peptide.

Enhancement of the Pharmacokinetics and Pharmacodynamics of Fusion Proteins with Combination Therapy using Endopeptidase Inhibitors.

[00240] The present invention also provides combination therapy using neutral endopeptidase (NEP) inhibitors and transferrin fusion proteins. Moreover, the present invention includes combination therapy using NEP and DPP-IV inhibitors and transferrin fusion proteins. The NEP and DPP-IV inhibitors may be administered concurrently or sequentially. Furthermore, the inhibitors and the transferrin fusion proteins may be administered concurrently or sequentially. The inhibitors may be administered prior to or after administering the transferrin fusion proteins.

[00241] The transferrin fusion protein comprise the GLP-1(7-37) peptide (SEQ ID NO: 32) or the GLP-1(7-36) peptide (amino acids 1-30 of SEQ ID NO: 32). More preferably, the GLP-1(7-37) peptide or GLP-1(7-36) peptide comprises an A8 to G and K34 to A mutation. The transferrin protein may also comprise a linker between GLP-1(7-37) peptide or GLP-1(7-36) peptide and the transferrin molecule. Preferably, the linker is the (PEAPTD)₂ peptide.

[00242] Neutral endopeptidase (NEP), which is also known as enkephalinase, neprilysin, and atriopeptidase, is a membrane-bound zinc metalloendopeptidase found in many tissues including the brain, kidney, lungs, gastrointestinal tract, heart, and peripheral vasculature. NEP plays a major role in the clearance of natriuretic peptides by degrading circulating natriuretic peptides, thus preventing their effects on vasodilation, blood pressure and volume. NEP, by degrading and inactivating the natriuretic peptides, is associated with, hypertension, heart failure, and renal failure.

[00243] In addition to degrading circulating natriuretic peptides, NEP also degrades other vasodilating substances including circulating bradykinins; adrenomedullin, renal vasodilating and natriuretic-diuretic peptide; and/or urodilatin, a renal form of ANP.

[00244] NEP is also involved in the degradation of endothelin isoform ET-1, a vasoconstrictor, and may be involved in the formation of ET-1 (Brunner-La Rocca *et al.*, Cardiovascular Research 51 (2001) 510-520). NEP also degrades angiotensin II, a potent

vasoconstrictor, endomorphins, and a number of peptides involved in metabolism such as bradykinin, GLP-1 (Hupe-Sodmann, K., McGregor, G P., Bridenbaugh, R *et al* Regulatory Peptides 1995; 58: 149-56, Hupe-Sodmann, K, Goeke, R., Goeke, B *et al* Peptides 1997; 18: 625-32), PYY (Medeiros MD, Turner AJ., Endocrinology. 1994 May;134(5):2088-94) and glucagon (Trebien R *et al* Am J Physiol Endocrinol Metab. 2004 Sep;287(3):E431-8).

[00245] A number of NEP inhibitors such as phosphoramidon or NEP/ACE inhibitors (including omapatrilat disclosed in U.S. Pat. No. 5,508,272, gempatrilat disclosed in U.S. Pat. No. 5,552,397, sampatrilat and MDL100240 disclosed in U.S. Pat. No. 5,430,145) have been reported in the literature as useful for the monotherapeutic treatment of, for example, hypertension and heart failure. Nathisuwan *et al.*, "A Review of Vasopeptidase Inhibitors: A New Modality in the Treatment of Hypertension and Chronic Heart Failure," Pharmacotherapy, Vol. 22(1), pp. 27-42 (2002). Candoxatril and ecadotril are the two highly specific inhibitors of NEP presently undergoing trials as future drugs for heart failure. Both compounds are prodrugs which are metabolized in the body to active congeners. Candoxatril is activated in the liver to candoxatrilat, while ecadotril is converted to its active congener, S-thiorphan.

[00246] Some examples of ACE/NEP inhibitors disclosed in U.S. Pat. Nos. 5,508,272, 5,362,727, 5,366,973, 5,225,401, 4,722,810, 5,223,516, 5,552,397, 4,749,688, 5,504,080, 5,612,359, and 5,525,723, and European Patent Applications 0481,522, 0534363A2, 534,396, and 534,492.

[00247] The present invention provides combination therapy comprising transferrin fusion proteins and DPP-IV inhibitors and or ACE/NEP inhibitors for the treatment of various diseases or conditions. Such diseases or conditions include but are not limited to diabetes, preferably type II diabetes, congestive heart failure, obesity, hypertension, and irritable bowel syndrome.

Transgenic Animals

[00248] The production of transgenic non-human animals that express a modified polypeptide or peptide that is protected from DPP activity is contemplated in one embodiment of the present invention. In some embodiments, transgenic non-human

animals expressing fusion proteins comprising a modified polypeptide or peptide and having increased stability is contemplated.

[00249] The successful production of transgenic, non-human animals has been described in a number of patents and publications, such as, for example U.S. Patent 6,291,740 (issued September 18, 2001); U.S. Patent 6,281,408 (issued August 28, 2001); and U.S. Patent 6,271,436 (issued August 7, 2001) the contents of which are hereby incorporated by reference in their entirety.

[00250] The ability to alter the genetic make-up of animals, such as domesticated mammals including cows, pigs, goats, horses, cattle, and sheep, allows a number of commercial applications. These applications include the production of animals which express large quantities of exogenous proteins in an easily harvested form (*e.g.*, expression into the milk or blood), the production of animals with increased weight gain, feed efficiency, carcass composition, milk production or content, disease resistance and resistance to infection by specific microorganisms and the production of animals having enhanced growth rates or reproductive performance. Animals which contain exogenous DNA sequences in their genome are referred to as transgenic animals.

[00251] The most widely used method for the production of transgenic animals is the microinjection of DNA into the pronuclei of fertilized embryos (Wall *et al.*, *J. Cell. Biochem.* 49:113 [1992]). Other methods for the production of transgenic animals include the infection of embryos with retroviruses or with retroviral vectors. Infection of both pre- and post-implantation mouse embryos with either wild-type or recombinant retroviruses has been reported (Janenich, *Proc. Natl. Acad. Sci. USA* 73:1260 [1976]; Janenich *et al.*, *Cell* 24:519 [1981]; Stuhlmann *et al.*, *Proc. Natl. Acad. Sci. USA* 81:7151 [1984]; Jahner *et al.*, *Proc. Natl. Acad. Sci. USA* 82:6927 [1985]; Van der Putten *et al.*, *Proc. Natl. Acad. Sci. USA* 82:6148-6152 [1985]; Stewart *et al.*, *EMBO J.* 6:383-388 [1987]).

[00252] An alternative means for infecting embryos with retroviruses is the injection of virus or virus-producing cells into the blastocoele of mouse embryos (Jahner, D. *et al.*, *Nature* 298:623 [1982]). The introduction of transgenes into the germline of mice has been reported using intrauterine retroviral infection of the midgestation mouse embryo (Jahner *et al.*, *supra* [1982]). Infection of bovine and ovine embryos with retroviruses or retroviral vectors to create transgenic animals has been reported. These protocols involve the microinjection of retroviral particles or growth arrested (*i.e.*, mitomycin C-treated) cells which

shed retroviral particles into the perivitelline space of fertilized eggs or early embryos (PCT International Application WO 90/08832 [1990]; and Haskell and Bowen, *Mol. Reprod. Dev.*, 40:386 [1995]. PCT International Application WO 90/08832 describes the injection of wild-type feline leukemia virus B into the perivitelline space of sheep embryos at the 2 to 8 cell stage. Fetuses derived from injected embryos were shown to contain multiple sites of integration.

[00253] U.S. Patent 6,291,740 (issued September 18, 2001) describes the production of transgenic animals by the introduction of exogenous DNA into pre-maturation oocytes and mature, unfertilized oocytes (*i.e.*, pre-fertilization oocytes) using retroviral vectors which transduce dividing cells (*e.g.*, vectors derived from murine leukemia virus [MLV]). This patent also describes methods and compositions for cytomegalovirus promoter-driven, as well as mouse mammary tumor LTR expression of various recombinant proteins.

[00254] U.S. Patent 6,281,408 (issued August 28, 2001) describes methods for producing transgenic animals using embryonic stem cells. Briefly, the embryonic stem cells are used in a mixed cell co-culture with a morula to generate transgenic animals. Foreign genetic material is introduced into the embryonic stem cells prior to co-culturing by, for example, electroporation, microinjection or retroviral delivery. ES cells transfected in this manner are selected for integrations of the gene via a selection marker such as neomycin.

[00255] U.S. Patent 6,271,436 (issued August 7, 2001) describes the production of transgenic animals using methods including isolation of primordial germ cells, culturing these cells to produce primordial germ cell-derived cell lines, transforming both the primordial germ cells and the cultured cell lines, and using these transformed cells and cell lines to generate transgenic animals. The efficiency at which transgenic animals are generated is greatly increased, thereby allowing the use of homologous recombination in producing transgenic non-rodent animal species.

Gene Therapy

[00256] The use of modified polypeptide or peptide constructs of the present invention for gene therapy is contemplated in one embodiment of this invention. The polypeptide or peptide has been modified to protect it from DPP activity by the addition of one or more additional amino acids at its N-terminus. For example, the nucleic acid construct encoding

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GLP-1 comprising an additional His residue at its N-terminus is provided for gene therapy. Also, the nucleic acid construct encoding modified GLP-1/transferrin fusion protein is provided for gene therapy. The modified GLP-1 constructs of the present invention are protected from DPP activity and are more stable; thus, they are ideally suited to gene therapy treatments.

[00257] Briefly, gene therapy via injection of an adenovirus vector containing a gene encoding a soluble fusion protein consisting of cytotoxic lymphocyte antigen 4 (CTLA4) and the Fc portion of human immunoglobulin G1 was recently shown in Ijima *et al.* (June 10, 2001) Human Gene Therapy (United States) 12/9:1063-77. In this application of gene therapy, a murine model of type II collagen-induced arthritis was successfully treated via intraarticular injection of the vector.

[00258] Gene therapy is also described in a number of U.S. patents including U.S. Pat. 6,225,290 (issued May 1, 2001); U.S. Pat. 6,187,305 (issued February 13, 2001); and U.S. Pat. 6,140,111 (issued October 31, 2000).

[00259] U.S. Patent 6,225,290 provides methods and constructs whereby intestinal epithelial cells of a mammalian subject are genetically altered to operatively incorporate a gene which expresses a protein which has a desired therapeutic effect. Intestinal cell transformation is accomplished by administration of a formulation composed primarily of naked DNA, and the DNA may be administered orally. Oral or other intragastrintestinal routes of administration provide a simple method of administration, while the use of naked nucleic acid avoids the complications associated with use of viral vectors to accomplish gene therapy. The expressed protein is secreted directly into the gastrointestinal tract and/or blood stream to obtain therapeutic blood levels of the protein thereby treating the patient in need of the protein. The transformed intestinal epithelial cells provide short or long term therapeutic cures for diseases associated with a deficiency in a particular protein or which are amenable to treatment by overexpression of a protein.

[00260] U.S. Pat. 6,187,305 provides methods of gene or DNA targeting in cells of vertebrate, particularly mammalian, origin. Briefly, DNA is introduced into primary or secondary cells of vertebrate origin through homologous recombination or targeting of the DNA, which is introduced into genomic DNA of the primary or secondary cells at a preselected site.

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[00261] U.S. Pat. 6,140,111 (issued October 31, 2000) describes retroviral gene therapy vectors. The disclosed retroviral vectors include an insertion site for genes of interest and are capable of expressing high levels of the protein derived from the genes of interest in a wide variety of transfected cell types. Also disclosed are retroviral vectors lacking a selectable marker, thus rendering them suitable for human gene therapy in the treatment of a variety of disease states without the co-expression of a marker product, such as an antibiotic. These retroviral vectors are especially suited for use in certain packaging cell lines. The ability of retroviral vectors to insert into the genome of mammalian cells has made them particularly promising candidates for use in the genetic therapy of genetic diseases in humans and animals. Genetic therapy typically involves (1) adding new genetic material to patient cells *in vivo*, or (2) removing patient cells from the body, adding new genetic material to the cells and reintroducing them into the body, *i.e.*, *in vitro* gene therapy. Discussions of how to perform gene therapy in a variety of cells using retroviral vectors can be found, for example, in U.S. Pat. Nos. 4,868,116, issued Sep. 19, 1989, and 4,980,286, issued Dec. 25, 1990 (epithelial cells), WO 89/07136 published Aug. 10, 1989 (hepatocyte cells), EP 378,576 published Jul. 25, 1990 (fibroblast cells), and WO 89/05345 published Jun. 15, 1989 and WO/90/06997, published Jun. 28, 1990 (endothelial cells), the disclosures of which are incorporated herein by reference.

[00262] Without further description, it is believed that one of ordinary skill in the art can, using the preceding description and the following illustrative examples, make and utilize the claimed invention. The following working examples therefore, specifically point out preferred embodiments of the present invention, and are not to be construed as limiting in any way the remainder of the disclosure. All articles, publications, patents and documents referred to throughout this application are hereby incorporated by reference in their entirety.

EXAMPLES

Example 1: Modified GLP-1 Having Dipeptidyl-Peptidase IV Protection

[00263] This Example describes modified GLP-1 peptides protected from DPP-IV activity. The following peptides were synthesized using standard solid phase Fmoc chemistry and

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purified by reverse phase HPLC using a C18 column and quantitated by absorbance at 220nm. The purified peptides were analyzed by mass spectrometry (MALDI-TOF):

GLP-1

NH₂-His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-COOH (amino acids 1-30 of SEQ ID NO: 32)

GLP-1 (A8G)

NH₂-His-Gly-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-COOH (SEQ ID NO: 90)

H-GLP-1

NH₂-His-His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-COOH (SEQ ID NO: 91)

H-GLP-1 (A8G)

NH₂-His-His-Gly-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-COOH (SEQ ID NO: 92)

HH-GLP-1

NH₂-His-His-His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-COOH (SEQ ID NO: 93)

G-GLP-1

NH₂-Gly-His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-COOH (SEQ ID NO: 94)

H-Exendin-4

NH₂-His-His-Gly-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Leu-Ser-Lys-Gln-Met-Glu-Glu-Glu-Ala-Val-Arg-Leu-Phe-Ile-Glu-Trp-Leu-Lys-Asn-Gly-Gly-Pro-Ser-Ser-Gly-Ala-Pro-Pro-Pro-Ser-COOH (SEQ ID NO: 95)

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Dipeptidylpeptidase-IV Treatment

[00264] Equimolar concentrations of each peptide (6 μ M) were treated with 2 μ g of recombinant human DPP-IV (1 μ g/ μ L, R&D Systems, Minneapolis, MN) in 25 mM Tris-Cl (pH 8.0). Control reactions excluding DPP-IV were set up in parallel for each peptide. The digests were incubated at room temperature for 2 hours, at which time the reactions were diluted 10-fold in Krebs-Ringer buffer (Biosource International, Camarillo, CA) supplemented with 1mM 3-Isobutyl-1-methylxanthine (IBMX, Calbiochem, San Diego, CA). The peptides were then analyzed to determine residual GLP-1 receptor activating activity, as described below.

Cyclic AMP Stimulation Assay

[00265] Four 96-well tissue culture plates were seeded with CHO-GLP1R cells (Montrose-Rafizadeh, *et al.* 1997 *J. Biol. Chem.* 272, 21201-21206) at a density of 2×10^4 cells/well in RPMI/10% FBS medium one day prior to treatment. The next day the cells appeared uniformly distributed with an approximate confluency of 60-80 percent. One day after seeding the culture plates the cells were washed twice with Krebs-Ringer buffer (KRB) followed by incubation in KRB for 1hr at 37°C to lower the intracellular levels of cAMP. This was followed by incubation for 10 minutes in KRB/IBMX to inhibit intracellular enzymes that break down cAMP. Dilutions of each test compound were prepared in KRB/IBMX and triplicate wells of CHO-GLP1R cells were treated with 50 μ l of test compound per well for exactly 20 minutes at 37°C. The treatment was halted by washing the cultures twice with ice-cold phosphate-buffered saline. Lysates were prepared by the addition of 0.1ml lysis buffer 1B (Amersham Biosciences cAMP Biotrak EIA kit) for 10 minutes at room temperature. The entire volume of each cell extract was then assayed to determine the cAMP concentration using the cAMP Biotrak Enzyme Immunoassay System (Amersham Biosciences Corporation, Piscataway, NJ, product code RPN225) according to kit instructions. Peptides of the invention were found to be more resistant to DPP-IV than the unmodified forms.

Active GLP-1 specific ELISA

[00266] Alternatively, DPP-IV degradation of GLP-1 and GLP-1 derivatives of the invention was assayed using an ELISA system (Glucagon-Like Peptide-1 [Active] ELISA

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kit [Linco Research, Inc., St. Charles, MO]) that is specific for intact, active GLP-1 and does not recognize GLP-1 in which the N-terminal two amino acids have been removed due to the action of DPP-IV, *i.e.* GLP-1(9-36 or 9-37). Equimolar concentrations of GLP-1 and H-GLP-1 (1200pM) were treated with recombinant human DPP-IV (200ng/ μ L, R&D Systems, Minneapolis, MN) in 25 mM Tris-Cl (pH 8.0) and the reaction stopped by dilution in the assay buffer supplied with the kit, which contains protease inhibitors.

[00267] The kit comprises a 96-well microtitre plate coated with anti-GLP-1 monoclonal antibody. The plate was washed (25mM Borate-buffered Saline x4 in a plate washer, ThermoLabsystems Ultrawash Plus), then incubated with peptide samples (300pM and 10-fold serial dilutions down the plate) for 3 hours at room temperature. After washing as described above, the plate was incubated with Alkaline-Phosphatase-conjugated anti-GLP antibody (supplied as a ready-to-use component of the kit) for 2 hours at room temperature. After washing, 4-Methylumbelliferyl Phosphate (MUP) substrate (1:200 dilution in 50mM Borate pH 9.5) was applied to all wells, and incubated in the dark at room temperature for 30 minutes. The plate was read at 355 excitation and 460nm emission wavelengths on a SpectraMax Gemini EM fluorescence plate reader. As H-GLP-1 bound less readily to the monoclonal antibody than GLP-1 itself, the concentration of active H-GLP-1 remaining after DPP-IV treatment was determined using an H-GLP-1 standard curve. Figure 8 shows that H-GLP-1 is substantially more resistant to the action of DPP-IV than GLP-1.

Example 2: Modified GLP-1 Fusion Protein

[00268] This Example describes a fusion protein comprising a modified GLP-1 protected from DPP-IV activity fused to a modified transferrin molecule.

[00269] In order to construct a sequence encoding the transferrin secretion leader followed by GLP-1 and the N-terminal part of transferrin, the following overlapping primers were designed:

P0236-

TTCCCATACAAACTTAAGAGTCCAATTAGCTTCATCGCCA (SEQ ID NO: 96)

P0237-

GGTTTAGCTTGTTTTTTTATTGGCGATGAAGCTAATTGGACTCTTAAGTTTGTAT
GGGAA (SEQ ID NO: 97)

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P0244-

ATAAAAAACAAGCTAACCTAATTCTAACAAAGCAAAGATGAGGCTCGCCGTG
GGAGCCC (SEQ ID NO: 98)

P0245-

CAGGACGGCGCAGACCAGCAGGGCTCCACGGCGAGCCTCATCTTTGCTTGTTA
GAATTA (SEQ ID NO: 99)

P0248-

TGCTGGTCTGCGCCGTCCTGGGGCTGTGTCTGGCGCATGCTGAAGGTACTTTTA
CTTCTGATGTTTCTTC (SEQ ID NO: 100)

P0249-

AATTCTTTAGCAGCTTGACCTTCCAAATAAGAAGAAACATCAGAAGTAAAAGT
ACCTTCAGCATGCGCCAGACACAGCCC (SEQ ID NO: 101)

P0250-

TTATTTGGAAGGTCAAGCTGCTAAAGAATTTATTGCTTGGTTGGTTAAAGGTAG
GGTACCTGATAAAACT (SEQ ID NO: 102)

P0251-

AGTTTTATCAGGTACCCTACCTTTAACCAACCAAGCAATA (SEQ ID NO: 103)

The positions of these primers are shown below.

```

                                AflIII
                                -+----
                                >>.....P0236.....>>
721 ccaatgttac gtcccgttat attggagttc ttcccataca aacttaagag tccaattagc
    gtttacaatg cagggcaata taacctcaag aagggtatgt ttgaattctc aggttaatcg
    <<.....P0237.....<<

>>P0236.>> >>.....P0244.....>>
781 ttcatcgcca ataaaaaac aagctaaaoc taattctaac aagcaaagat gaggctcgcc
    aagttagcggg tatttttttg ttcgatttgg attaagattg ttcgtttcta ctccgagcgg
    <<.....P0237.....<< <<.....P0245.....<<
                                >>...nL.....>
                                m r l a

>>P0244.>> >>.....P0248.....>>
841 gtgggagccc tctctggctg cgcgcctctg ggcctgtgtc tggcgcagtc tgaaggtact
    caccctcggg acgaccagac gcggcaggac cccgacacag acccgttacc acttccatga
    <<.....P0245.....<< <<.....P0249.....<<
                                >>...nL.....>
                                v g a l l v c a v l g l c l a
                                >>...GLP-1.....>
                                h a e g t

>>.....P0248.....>> >>.....P0250.....>>
901 tttacttctg atgtttcttc ttatttggaa ggtcaagctg ctaaagaatt tattgcttgg
    aatgaagac tacaagaag aataaacctt ccagttcgac gatttcttaa ataacgaac
    <<.....P0249.....<< <<.....P0251.....<<
                                >>...GLP-1.....>
                                f t s d v s s y l e g q a a k e f i a w

                                KpnI
                                -----+
                                >>.....P0250.....>>
961 ttggttaaag gtaggtacc tgataaaact gtgagatggt gtgcagtgtc ggagcatgag
    aaccaatttc catcccatgg actattttga cactctacca cacgtcacag cctcgtactc
    <<.....P0251.....<<
                                >>...GLP-1.....>
                                l v k g r
                                >>.....mTf.....>
                                v p d k t v r w c a v s e h e

```

(SEQ ID NO: 104 is the coding strand; SEQ ID NO: 105 is the encoded protein.)

The primers (8µL of 20pmol conc.) were combined and heated to 65°C for 5 min. and then the annealing reaction was allowed to cool slowly to room temperature.

[00270] After adding T4 DNA ligase to the annealing reaction and incubating for a further 2hr at room temperature, 1 µL of the reaction was removed and used in a PCR reaction to amplify the completed insert with the outer primers P0236 and P0251. The PCR conditions were as follows:

- 5 min at 94°C
- 25 cycles of : 30sec at 94°C
- 30sec at 50°C
- 1 min at 72°C

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7 min at 72°C

hold at 4°C

[00271] The resulting PCR product was digested *Afl*III and *Kpn*I and ligated into pREX0094 (Figure 1) which had previously been digested with *Afl*III and *Kpn*I. The ligation was used to transform *E. coli*. The DNA from the resultant clones was sequenced and a clone correct the length of the *Afl*III/*Kpn*I insert was selected and designated pREX0198 (Figure 2). Next, pREX0198 was digested with *Not*I and *Pvu*I and inserted into pSAC35 (Figure 3) to create pREX0240 (Figure 4).

[00272] To create a plasmid encoding the natural transferrin secretion leader followed by H-GLP-1(7-36) fused to modified transferrin (mTf), overlapping primers P0424 and P0425 were designed to add the extra N-terminal histidine to the sequence encoded by pREX0198.

P0424 5' to 3'

CTGTGTCTGGCGCATCATGCTGAAG (SEQ ID NO: 106)

P0425 5' to 3'

CTTCAGCATGATGCGCCAGACACAG (SEQ ID NO: 107)

[00273] pREX0198 was used as the template for the initial PCR reactions using the two overlapping mutagenic primers and two outer primers in separate reactions, *i.e.* P0424 plus P0012 and P0425 plus P0025. The products of these reactions were then used as templates in a second round of PCR with just the outer primers, *i.e.* P0012 plus P0025, in order to join them together. The reaction conditions for both rounds of PCR were 1 x 94°C for 1 min, 20 x 94°C for 30 seconds, 50°C for 30 seconds, 72°C for 1 minute and 1 x 72°C for 7 minutes to finish.

[00274] The PCR product from the final reaction was digested with *Afl*III and *Kpn*I and ligated into *Afl*III/*Kpn*I digested pREX0052 (Figure 5) to create pREX0367 (Figure 6). The construct was DNA sequenced to confirm the insertion of the codon for the extra histidine.

[00275] pREX0367 was then digested with *Not*I and *Pvu*I (the latter to destroy the ampicillin resistance gene) and ligated into pSAC35 previously digested with *Not*I to create pREX0368 (Figure 7).

[00276] pREX0368 was transformed into the host *Saccharomyces cerevisiae* strain by electroporation and transformed colonies selected on the basis of leucine prototrophy on buffered minimal medium plates. After selection of single colonies, yeast transformants were stocked in 40% Trehalose and stored at -70°C. Expression was determined by growth in liquid minimal medium buffered to pH6.5 and analysis of supernatant by SDS-PAGE, western blot and ELISA.

[00277] The plasmids encoding GLP-1/mTf (pREX0100) and H-GLP-1/mTf were constructed as described in U.S. Application 10/378,094, filed March 4, 2003, which is herein incorporated by reference in its entirety. To produce the GLP-1/mTf fusion protein, the amino acid sequence of GLP-1(7-36) and GLP-1(7-37) may be used.

haegtftsdvssylegqaakefiawlvkgr (amino acids 1-30 of SEQ ID NO: 32)

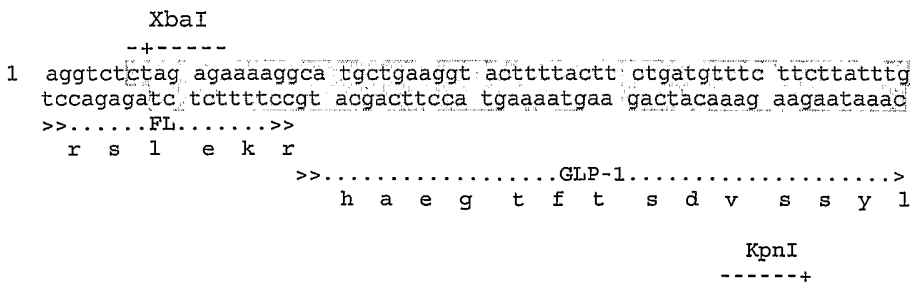
haegtftsdvssylegqaakefiawlvkgrg (SEQ ID NO: 32)

[00278] For example, the peptide sequence of GLP-1(7-36) may be back translated into DNA and codon optimized for yeast:

```
catgctgaaggtacttttacttctgatgtttcttcttatttgggaaggtcaagctgctaaagaa
h a e g t f t s d v s s y l e g q a a k e

tttattgcttggttggttaaaggtaga (SEQ ID NO: 117)
f i a w l v k g r (amino acids 1-30 of SEQ ID NO: 32)
```

[00279] The primers were specifically designed to form 5' *Xba*I and 3' *Kpn*I sticky ends after annealing and to enable direct ligation into *Xba*I/*Kpn*I cut pREX0052, just 5' of the end of the leader sequence and at the N-terminus of mTf. Alternatively, other sticky ends may be engineered for ligations into other vectors.



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61  gaaggtcaag ctgctaaaga atttattgct tggttggtta aaggtagggT acctgata
    cttccagttc gacgatttct taaataacga accaaccaat ttccatcca tggactat
>.....GLP-1.....>>
    e g q a a k e f i a w l v k g r
                                     >>..mTf..>>
                                     v p d

```

SEQ ID NOS: 118 and 119

[00280] After annealing and ligation, the clones were sequenced to confirm correct insertion. This vector was designated pREX0094. The cassette was cut out of pREX0094 with *NotI* and sub-cloned into *NotI* cut yeast vector, pSAC35, to make pREX0100.

[00281] This plasmid was then electroporated into the host *Saccharomyces* yeast strains and transformants selected for leucine prototrophy on minimal media plates. Expression was determined by growth in liquid minimal media and analysis of supernatant by SDS-PAGE, western blot, and ELISA.

[00282] GLP-1/mTf and H-GLP-1/mTf were expressed and purified from fermentation cultures, grown under standard conditions by cation exchange and anion exchange chromatography.

Dipeptidylpeptidase-IV Treatment

[00283] Equimolar concentrations of GLP-1/mTf and H-GLP-1-mTF (2 μ M) were treated with recombinant human DPP-IV (1 μ g/ μ L, R&D Systems) in a solution of 25mM Tris-Cl (pH 8.0). Control reactions excluding DPP-IV were set-up in parallel for each fusion protein. The digests were incubated at room temperature for 2 hours, at which time the reactions were diluted 20-fold in Krebs-Ringer buffer (Biosource International) supplemented with 1 mM IBMX (Calbiochem).

Cyclic AMP Stimulation Assay

[00284] Tissue culture plates (24-well) were seeded with CHO-GLP1R cells at a density of 1×10^5 cells per/well in RPMI/10% FBS medium one day prior to treatment. The next day the cells appeared uniformly distributed with an approximate confluency of 60-80 percent. One day after seeding the culture plates the cells were washed twice with Krebs-Ringer buffer (KRB) followed by incubation in KRB for 1hr at 37°C to lower the intracellular levels of cAMP. This was followed by incubation for 10 minutes in KRB/IBMX to inhibit intracellular enzymes that break down cAMP. Dilutions of each test compound were

prepared in KRB/IBMX and triplicate wells of CHO-GLP1R cells were treated with 0.15ml of test compound per well for exactly 50 minutes at 37°C. The treatment was halted by washing the cultures two times with ice-cold phosphate-buffered saline. Lysates were prepared by the addition of 0.2ml lysis buffer 1B (Amersham Biosciences cAMP Biotrak EIA kit) for 10 minutes at room temperature, then 100µl of each cell extract was then assayed to determine the cAMP concentration using the cAMP Biotrak Enzyme Immunoassay System (Amersham Biosciences) according to kit instructions.

[00285] H-GLP-1/mTf was found to be more resistant to DPP-IV than GLP-1/mTf

Example 3: Modified GLP-1/mTf for the Treatment of Diabetes

[00286] In this Example, modified GLP-1/mTf of the present invention is used as a therapeutic agent to treat diabetes. Modified GLP-1/mTf is administered to Zucker rats, a standard animal model for type II diabetes. Zucker rats have abnormally high blood glucose levels. It has been shown that treatment of these animals with GLP-1 induces insulin secretion and reduces blood glucose.

[00287] Zucker rats are fasted overnight and then treated with H-GLP-1 or H-GLP-1 fused to transferrin (H-GLP-1/mTf). Thirty minutes after subcutaneous injection of H-GLP-1 or H-GLP-1/mTf, the animals are subjected to a Glucose Tolerance Test (GTT). For this test, fasted animals are fed glucose solution (1.5mg/g body weight), and the blood glucose is measured at appropriate time intervals. Soon after the glucose administration, the blood glucose level of the untreated animals rises and slowly drops towards the base line while the animals which are injected with H-GLP-1 or H-GLP-1/mTf show faster normalization of blood glucose level due to the insulinotropic effect of the GLP-1.

[00288] In a further experiment, modified H-GLP-1 or H-GLP-1/mTf is used to normalize the high fasting glucose of the Zucker rats without glucose administration. While the blood glucose levels remain high in the untreated animals, a significant drop is seen in the H-GLP-1 or modified H-GLP-1/mTf treated animals.

Example 4: Modified Glucagon Having Dipeptidyl-Peptidase IV Protection

[00289] This Example describes modified glucagon molecules protected from DPP-IV activity.

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[00290] The following peptides are synthesized using standard solid phase Fmoc chemistry and purified by reverse phase HPLC using a C18 column and quantitated by absorbance at 220nm. The purified peptides are analyzed by mass spectrometry (MALDI-TOF):

Glucagon

NH₂-His-Ser-Gln-Gly-Thr-Phe-Thr-Ser-Asp-Tyr-Ser-Lys-Tyr-Leu-Asp-Ser-Arg-Arg-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-Met-Asn-Thr-COOH (SEQ ID NO: 35)

H-Glucagon

NH₂-His-His-Ser-Gln-Gly-Thr-Phe-Thr-Ser-Asp-Tyr-Ser-Lys-Val-Leu-Asp-Ser-Arg-Arg-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-Met-Asn-Thr-COOH (SEQ ID NO: 108)

[00291] The peptides are pre-treated with DPP-IV as described above and then assayed for the ability to activate the glucagon receptor using a recombinant cell line expressing a cloned glucagon receptor.

Example 5: Modified GIP Having Dipeptidyl-Peptidase IV Protection

[00292] This Example provides modified GIP molecules protected from DPP-IV activity.

[00293] The following peptides are synthesized using standard solid phase Fmoc chemistry and purified by reverse phase HPLC using a C18 column and quantitated by absorbance at 220nm. The purified peptides are analysed by mass spectrometry (MALDI-TOF):

GIP

NH₂-Tyr-Ala-Glu-Gly-Thr-Phe-Ile-Ser-Asp-Tyr-Ser-Ile-Ala-Met-Asp-Lys-Ile-His-Gln-Gln-Asp-Phe-Val-Asn-Trp-Leu-Leu-Ala-Gln-Lys-Gly-Lys-Lys-Asn-Asp-Trp-Lys-His-Asn-Ile-Thr-Gln-COOH (SEQ ID NO: 31)

Y-GIP

NH₂-Tyr-Tyr-Ala-Glu-Gly-Thr-Phe-Ile-Ser-Asp-Tyr-Ser-Ile-Ala-Met-Asp-Lys-Ile-His-Gln-Gln-Asp-Phe-Val-Asn-Trp-Leu-Leu-Ala-Gln-Lys-Gly-Lys-Lys-Asn-Asp-Trp-Lys-His-Asn-Ile-Thr-Gln-COOH (SEQ ID NO: 109)

[00294] The peptides are pre-treated with DPP-IV as described above and then assayed for the ability to activate the GIP receptor using a recombinant cell line expressing a cloned GIP receptor.

[00295] It should be understood that the foregoing discussion and examples merely present a detailed description of certain preferred embodiments. It therefore should be apparent to those of ordinary skill in the art that various modifications and equivalents can be made without departing from the spirit and scope of the invention. All journal articles, other references, patents, and patent applications that are identified in this patent application are incorporated by reference in their entirety.

CLAIMS

1. A method of treating a disease or condition in a patient comprising administering an effective amount of a transferrin (Tf) fusion protein and at least one second agent.
2. The method of claim 1, wherein the second agent is selected from the group consisting of a DPP-IV inhibitor and a neutral endopeptidase (NEP) inhibitor.
3. The method of claim 1, wherein the Tf fusion protein is susceptible to DPP-IV clearance.
4. The method of claim 1, wherein the Tf fusion protein is sensitive, resistant or partially resistant to DPP-IV clearance.
5. The method of claim 1, wherein the Tf fusion protein comprises one or more GLP-1 peptides fused to a Tf molecule.
6. The method of claim 5, wherein the Tf fusion protein further comprises a linker.
7. The method of claim 6, wherein the linker is PEAPTD or (PEAPTD)₂.
8. The method of claim 5, wherein the GLP-1 peptide is at the N-terminus of the fusion protein.
9. The method of claim 5, wherein the GLP-1 peptide is GLP-1(7-37) having SEQ ID NO: 32 or GLP-1(7-36) (amino acids 1-30 of SEQ ID NO: 32).
10. The method of claim 9, wherein the GLP-1 peptide has been modified by mutating A8 to G.
11. The method of claim 9, wherein the GLP-1 peptide has been modified by mutating K34 to A.
12. The method of claim 9, wherein the GLP-1 peptide has been modified by

mutating A8 to G and K34 to A.

13. The method of claim 12, wherein the GLP-1 peptide is linked to the Tf molecule via the linker (PEAPTD)₂.

14. The method of claim 5, wherein the Tf molecule has been modified to exhibit reduced glycosylation as compared to a fully glycosylated Tf molecule.

15. The method of claim 5, wherein the Tf molecule has been modified to exhibit no glycosylation.

16. The method of claim 5, wherein the Tf molecule has been modified to comprise at least one mutation that prevents glycosylation.

17. The method of claim 5, wherein the Tf molecule has modified to have reduced affinity for a transferrin receptor (TfR) as compared to a wild-type Tf molecule.

18. The method of claim 5, wherein the Tf molecule has been modified to have no binding for a TfR.

19. The method of claim 5, wherein the Tf molecule has been modified to have reduced affinity for iron as compared to a wild-type Tf molecule.

20. The method of claim 5, wherein the Tf molecule has been modified to have no binding to iron.

21. The method of claim 5, wherein the Tf molecule is lactoferrin or melanotransferrin.

22. The method of claim 1, wherein the disease is a metabolic disease.

23. The method of claim 22, wherein the metabolic disease is prediabetes, diabetes, or obesity.

24. The method of claim 23, wherein the diabetes is type II diabetes.

25. The method of claim 1, wherein the disease is congestive heart failure.

26. The method of claim 1, wherein the disease is irritable bowel syndrome or dyspepsia.

27. The method of claim 1, wherein there are two second agents.

28. The method of claim 27, wherein the two second agents are a DPP-IV inhibitor and an NEP inhibitor.

29. The method of claim 29, wherein the two second agents are administered concurrently.

30. The method of claim 1 or 29 wherein the transferrin fusion protein and the one or more second agents are administered sequentially.

31. The method of claim 1 or 29, wherein the transferrin fusion protein and the one or more second agents are administered concurrently.

32. A composition comprising a transferrin (Tf) fusion protein and at least one second agent.

33. The composition of claim 32, wherein the second agent is selected from the group consisting of a DPP-IV inhibitor and a neutral endopeptidase (NEP) inhibitor.

34. The composition of claim 32, wherein the Tf fusion protein is susceptible to DPP-IV clearance.

35. The composition of claim 32, wherein the Tf fusion protein is sensitive, resistant or partially resistant to DPP-IV clearance.

36. The composition of claim 32, wherein the Tf fusion protein comprises one or more GLP-1 peptides fused to a Tf molecule.

37. The composition of claim 36, wherein the Tf fusion protein further comprises a linker.

38. The composition of claim 37, wherein the linker is PEAPTD or (PEAPTD)₂.
39. The composition of claim 36, wherein the GLP-1 peptide is at the N-terminus of the fusion protein.
40. The composition of claim 36, wherein the GLP-1 peptide is GLP-1(7-37) having SEQ ID NO: 32 or GLP-1(7-36) (amino acids 1-30 of SEQ ID NO: 32).
41. The composition of claim 40, wherein the GLP-1 peptide has been modified by mutating A8 to G.
42. The composition of claim 40, wherein the GLP-1 peptide has been modified by mutating K34 to A.
43. The composition of claim 40, wherein the GLP-1 peptide has been modified by mutating A8 to G and K34 to A.
44. The composition of claim 43, wherein the GLP-1 peptide is linked to the Tf molecule via the linker (PEAPTD)₂.
45. The composition of claim 32, wherein the Tf molecule has been modified to exhibit reduced glycosylation as compared to a fully glycosylated Tf molecule.
46. The composition of claim 32, wherein the Tf molecule has been modified to exhibit no glycosylation.
47. The composition of claim 32, wherein the Tf molecule has been modified to comprise at least one mutation that prevents glycosylation.
48. The composition of claim 32, wherein the Tf molecule has modified to have reduced affinity for a transferrin receptor (TfR) as compared to a wild-type Tf molecule.
49. The composition of claim 32, wherein the Tf molecule has been modified to have no binding for a TfR.
50. The composition of claim 32, wherein the Tf molecule has been modified to have reduced affinity for iron as compared to a wild-type Tf molecule.

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51. The composition of claim 32, wherein the Tf molecule has been modified to have no binding to iron.

52. The composition of claim 32, wherein the Tf molecule is lactoferrin or melanotransferrin.

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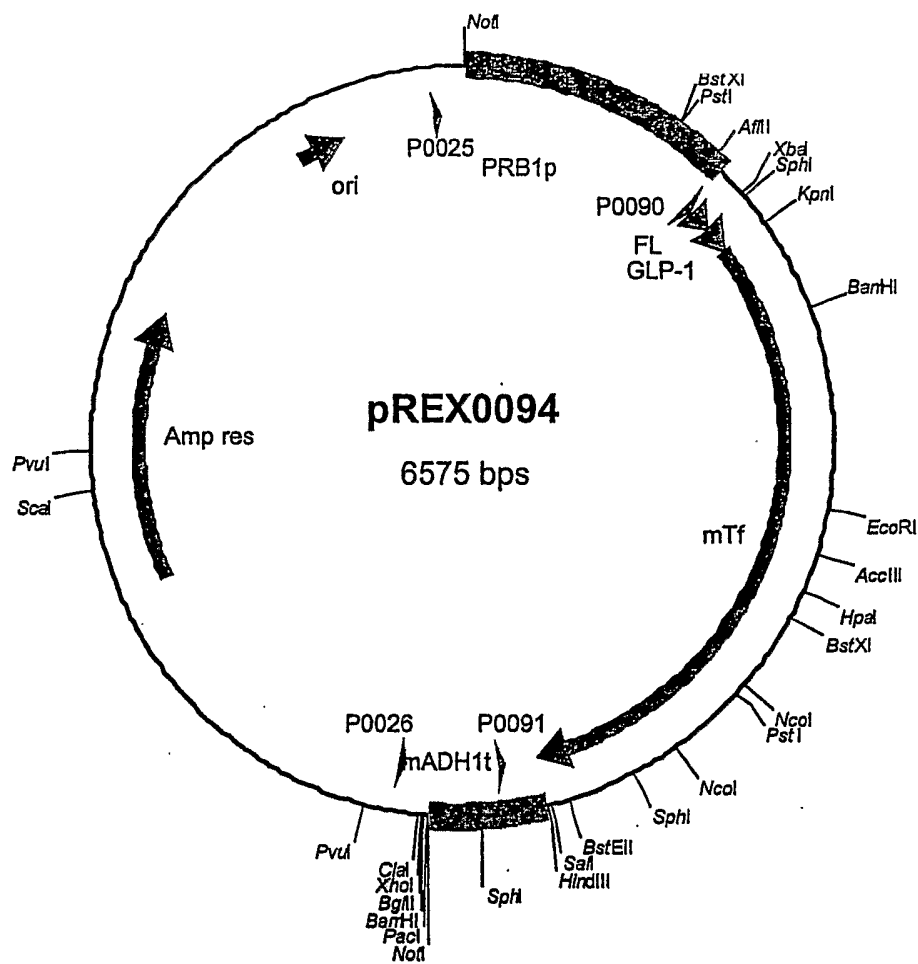


Figure 1

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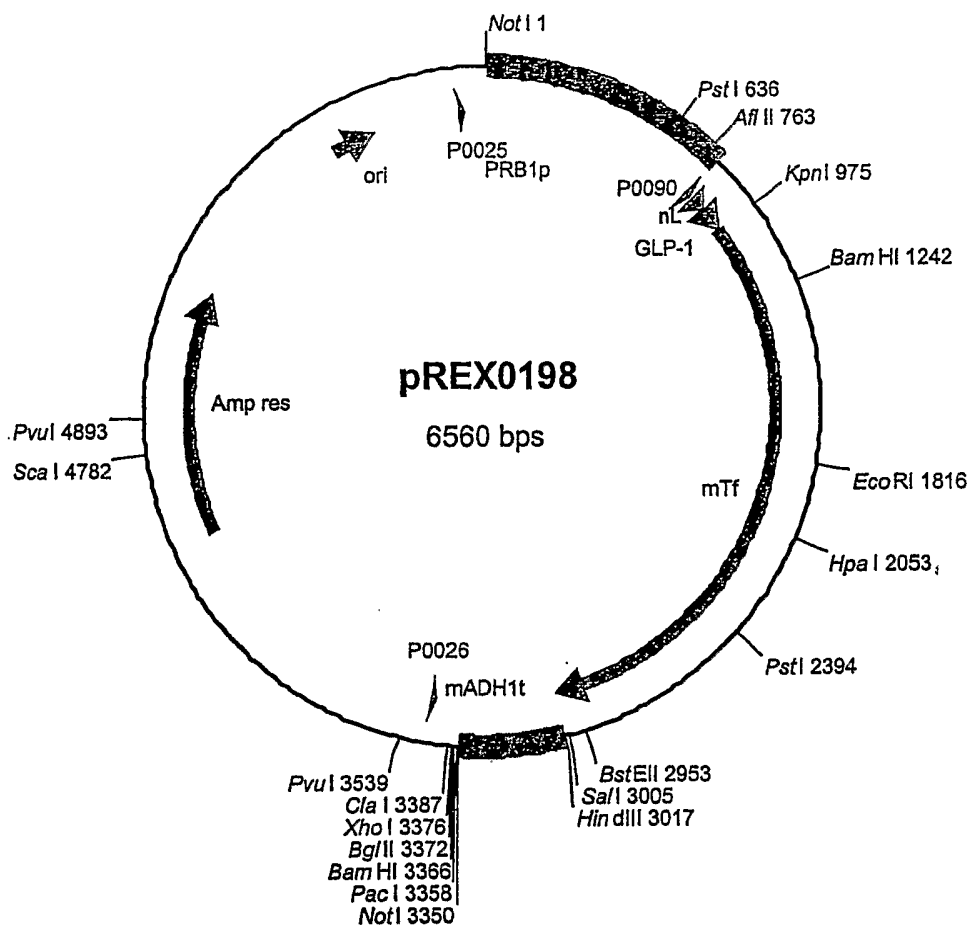


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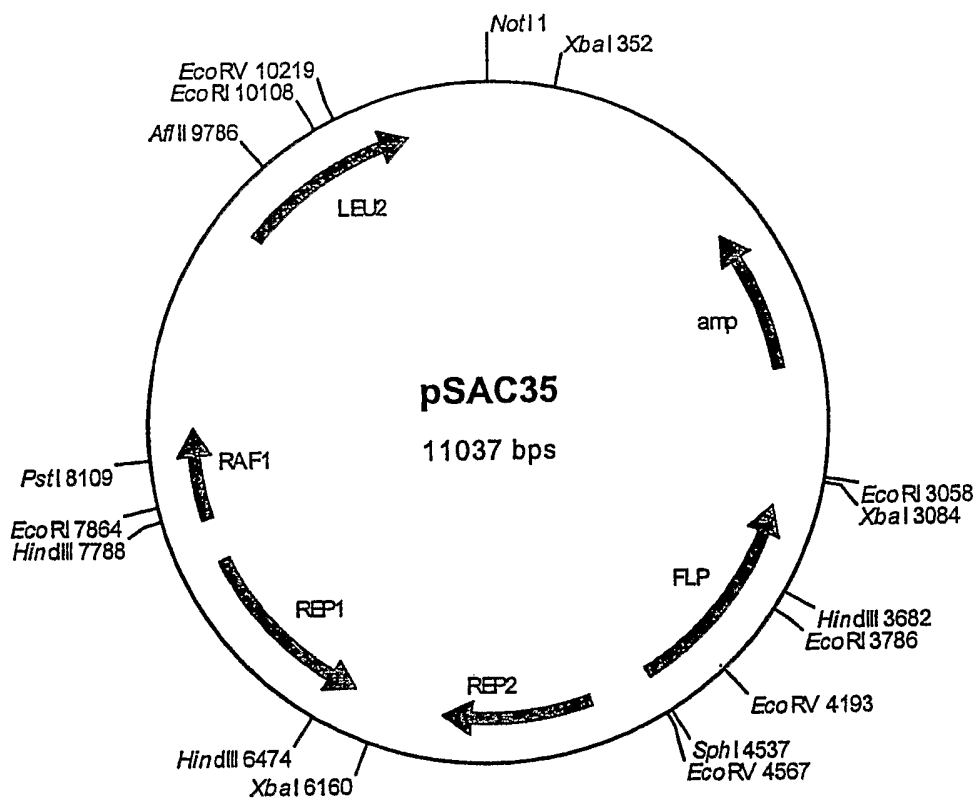


Figure 3

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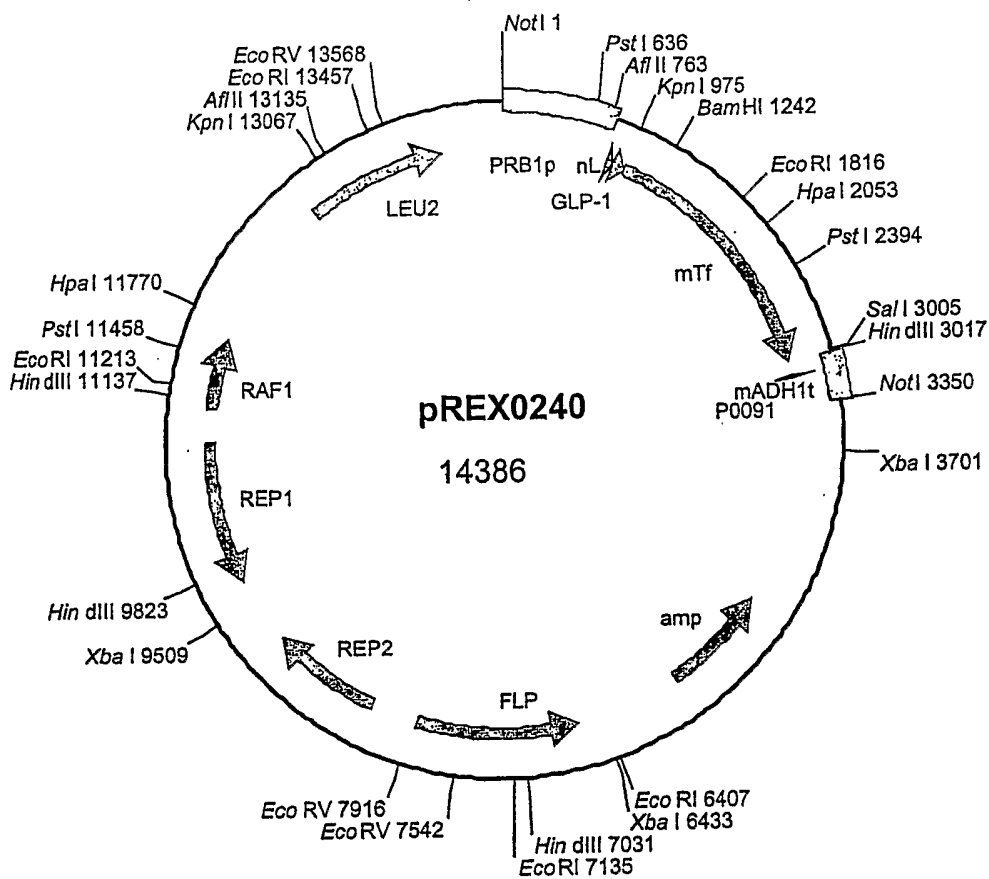


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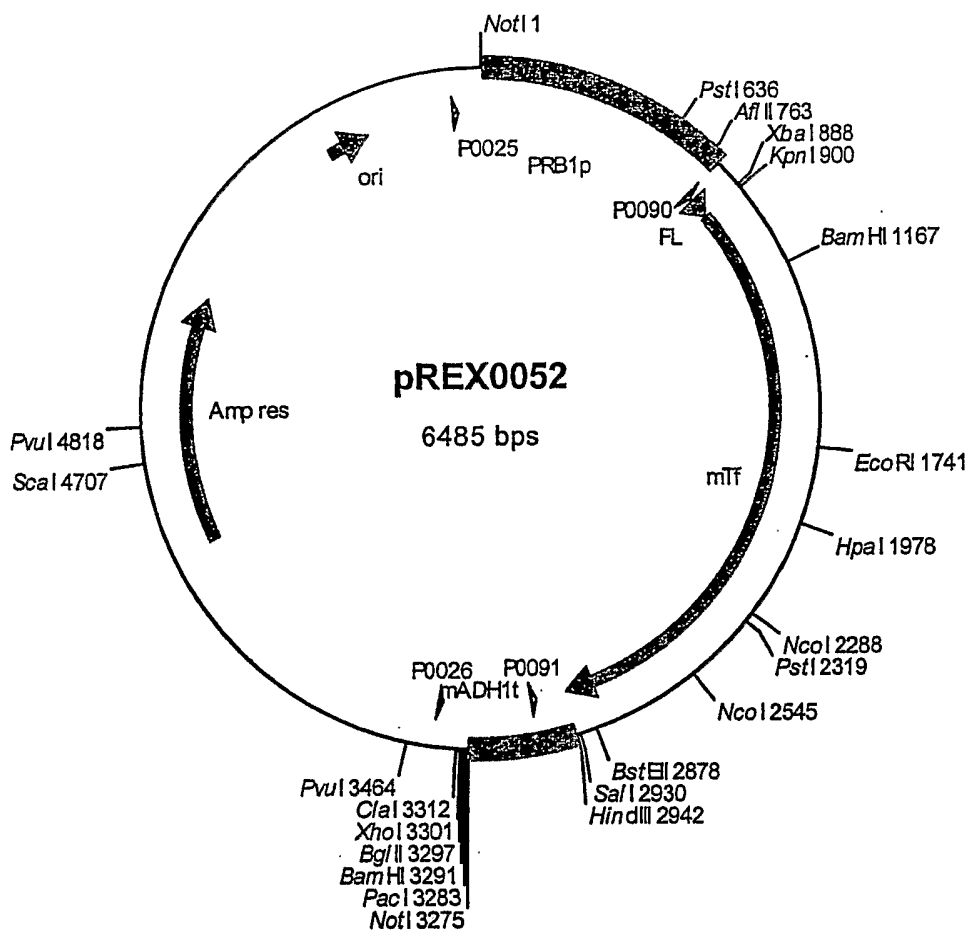


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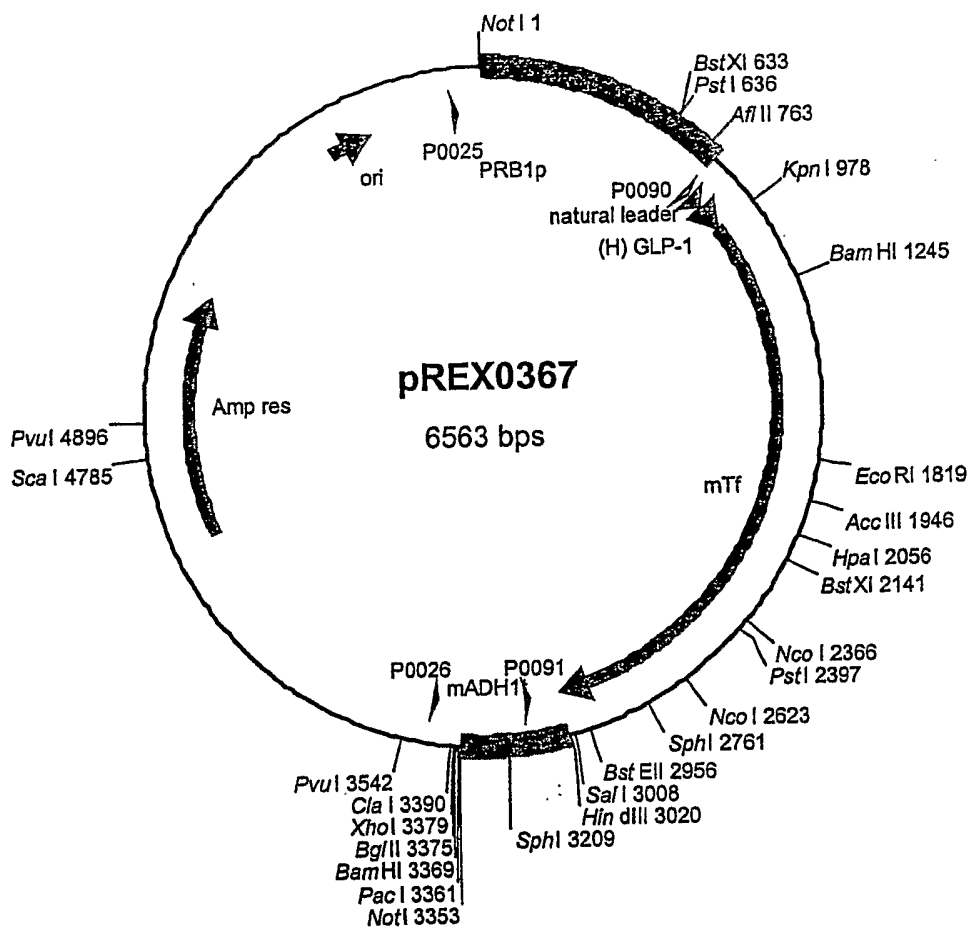


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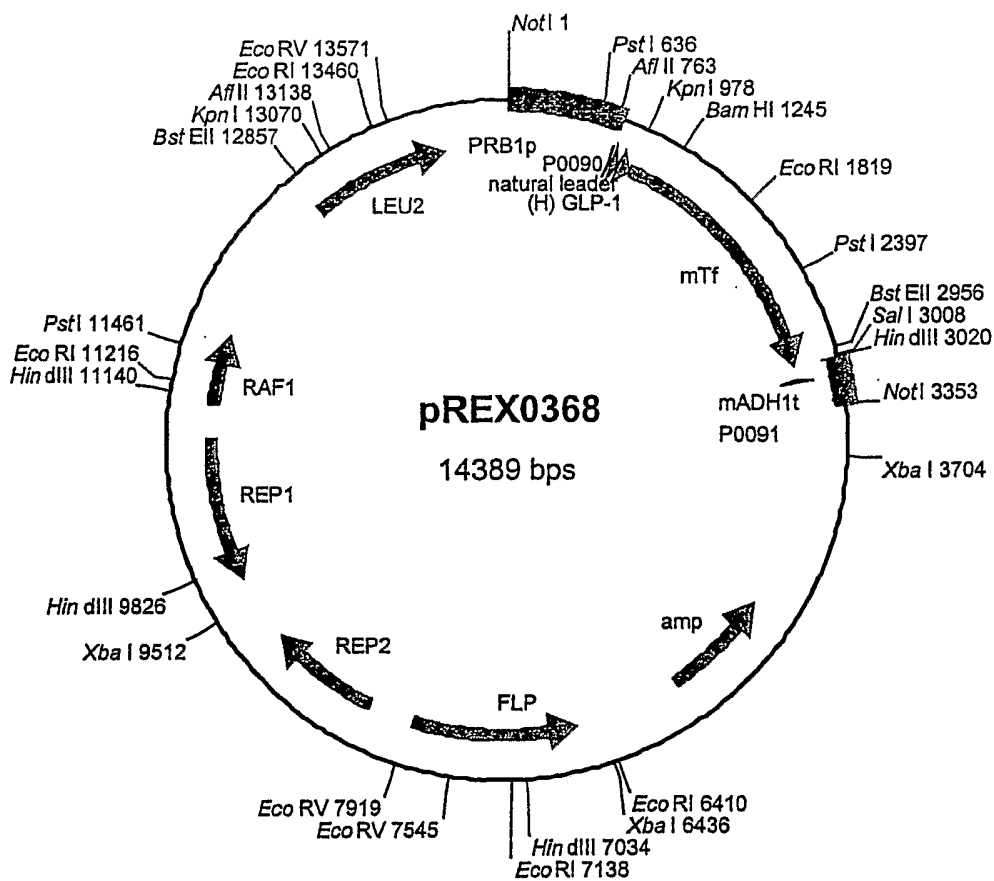


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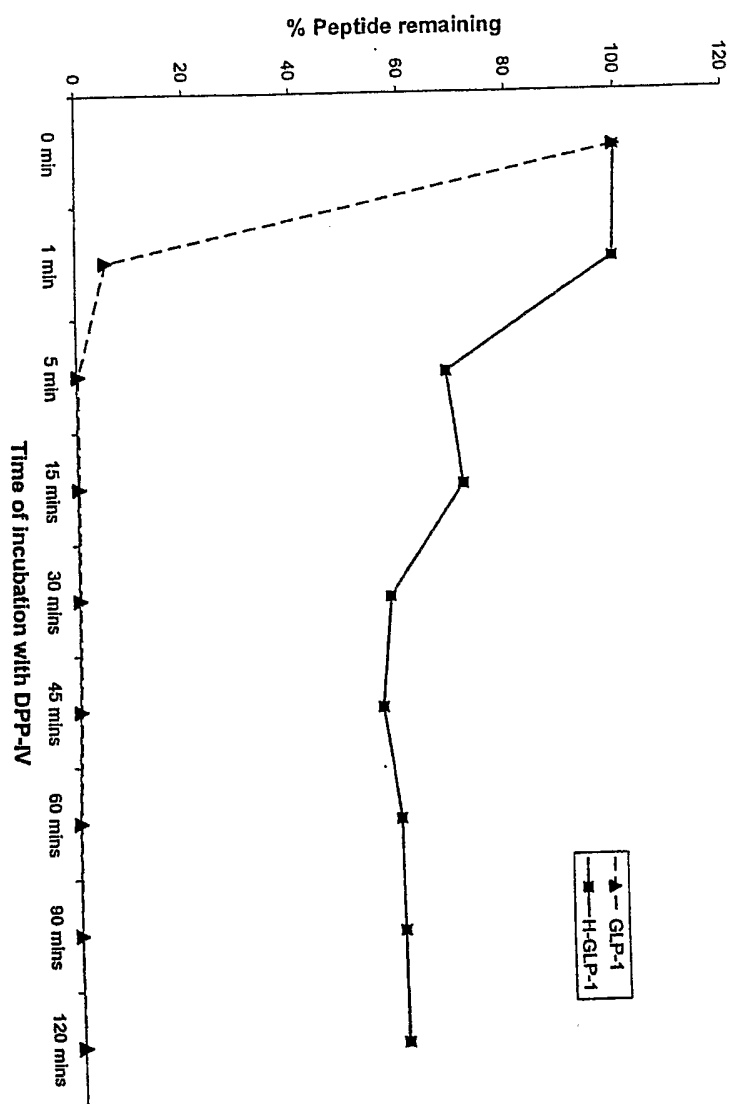


Figure 8

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Prior, Christopher P.
Ballance, David J.

<120> Combination Therapy Using Transferrin Fusion Proteins Comprising
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<213> Homo sapiens

<400> 54

Asn Pro Leu Leu Ile Leu Thr Phe Val Ala Ala Ala Leu Ala Ala Pro
1 5 10 15

Phe Asp Asp Asp Asp Lys Ile Val Gly Gly Tyr Asn Cys Glu Glu Asn
20 25 30

Ser Val Pro Tyr Gln Val Ser Leu Asn Ser Gly Tyr His Phe Cys Gly
35 40 45

Gly Ser Leu Ile Asn Glu Gln Trp Val Val Ser Ala Gly His Cys Tyr
50 55 60

Lys Ser Arg Ile Gln Val Arg Leu Gly Glu His Asn Ile Glu Val Leu
65 70 75 80

Glu Gly Asn Glu Gln Phe Ile Asn Ala Ala Lys Ile Ile Arg His Pro
85 90 95

Gln Tyr Asp Arg Lys Thr Leu Asn Asn Asp Ile Met Leu Ile Lys Leu
100 105 110

Ser Ser Arg Ala Val Ile Asn Ala Arg Val Ser Thr Ile Ser Leu Pro
115 120 125

Thr Ala Pro Pro Ala Thr Gly Thr Lys Cys Leu Ile Ser Gly Trp Gly
130 135 140

Asn Thr Ala Ser Ser Gly Ala Asp Tyr Pro Asp Glu Leu Gln Cys Leu
145 150 155 160

Asp Ala Pro Val Leu Ser Gln Ala Lys Cys Glu Ala Ser Tyr Pro Gly
165 170 175

Lys Ile Thr Ser Asn Met Phe Cys Val Gly Phe Leu Glu Gly Gly Lys
180 185 190

Asp Ser Cys Gln Gly Asp Ser Gly Gly Pro Val Val Cys Asn Gly Gln
 195 200 205

Leu Gln Gly Val Val Ser Trp Gly Asp Gly Cys Ala Gln Lys Asn Lys
 210 215 220

Pro Gly Val Tyr Thr Lys Val Tyr Asn Tyr Val Lys Trp Ile Lys Asn
 225 230 235 240

Thr Ile Ala Ala Asn Ser
 245

<210> 55
 <211> 184
 <212> PRT
 <213> Homo sapiens

<400> 55

Gly Pro Val Pro Thr Pro Pro Asp Asn Ile Gln Val Gln Glu Asn Phe
 1 5 10 15

Asn Ile Ser Arg Ile Tyr Gly Lys Trp Tyr Asn Leu Ala Ile Gly Ser
 20 25 30

Thr Cys Pro Trp Leu Lys Lys Ile Met Asp Arg Met Thr Val Ser Thr
 35 40 45

Leu Val Leu Gly Glu Gly Ala Thr Glu Ala Glu Ile Ser Met Thr Ser
 50 55 60

Thr Arg Trp Arg Lys Gly Val Cys Glu Glu Thr Ser Gly Ala Tyr Glu
 65 70 75 80

Lys Thr Asp Thr Asp Gly Lys Phe Leu Tyr His Lys Ser Lys Trp Asn
 85 90 95

Ile Thr Met Glu Ser Tyr Val Val His Thr Asn Tyr Asp Glu Tyr Ala
 100 105 110

Ile Phe Leu Thr Lys Lys Phe Ser Arg His His Gly Pro Thr Ile Thr
 115 120 125

Ala Lys Leu Tyr Gly Arg Ala Pro Gln Leu Arg Glu Thr Leu Leu Gln
 130 135 140

Asp Phe Arg Val Val Ala Gln Gly Val Gly Ile Pro Glu Asp Ser Ile
 145 150 155 160

Phe Thr Met Ala Asp Arg Gly Glu Cys Val Pro Gly Glu Gln Glu Pro
 165 170 175

Glu Pro Ile Leu Ile Pro Arg Val
 180

<210> 56
 <211> 77
 <212> PRT
 <213> Homo sapiens

<400> 56

Val Pro Leu Ser Arg Thr Val Arg Cys Thr Cys Ile Ser Ile Ser Asn
 1 5 10 15

Gln Pro Val Asn Pro Arg Ser Leu Glu Lys Leu Glu Ile Ile Pro Ala
 20 25 30

Ser Gln Phe Cys Pro Arg Val Glu Ile Ile Ala Thr Met Lys Lys Lys
 35 40 45

Gly Glu Lys Arg Cys Leu Asn Pro Glu Ser Lys Ala Ile Lys Asn Leu
 50 55 60

Leu Lys Ala Val Ser Lys Glu Arg Ser Lys Arg Ser Pro
 65 70 75

<210> 57
 <211> 74
 <212> PRT
 <213> Homo sapiens

<400> 57

Gly Pro Ala Ser Val Pro Thr Thr Cys Cys Phe Asn Leu Ala Asn Arg
 1 5 10 15

Lys Ile Pro Leu Gln Arg Leu Glu Ser Tyr Arg Arg Ile Thr Ser Gly
 20 25 30

Lys Cys Pro Gln Lys Ala Val Ile Phe Leu Thr Lys Leu Ala Lys Asp
 35 40 45

Ile Cys Ala Asp Pro Lys Lys Lys Tyr Val Gln Asp Ser Met Lys Tyr
 50 55 60

Leu Asp Gln Lys Ser Pro Thr Pro Lys Pro
 65 70

<210> 58
 <211> 74
 <212> PRT
 <213> Homo sapiens

<400> 58

Gln Pro Asp Ala Ile Asn Ala Pro Val Thr Cys Cys Tyr Asn Phe Thr
 1 5 10 15

Asn Arg Lys Ile Ser Val Gln Arg Leu Ala Ser Tyr Arg Arg Ile Thr
 20 25 30

Ser Ser Lys Cys Pro Lys Glu Ala Val Ile Phe Lys Thr Ile Val Ala
 35 40 45

Lys Glu Ile Cys Ala Asp Pro Lys Gln Lys Trp Val Gln Asp Ser Met
 50 55 60

Asp His Leu Asp Lys Gln Thr Gln Thr Pro
 65 70

<210> 59
 <211> 76
 <212> PRT
 <213> Homo sapiens

<400> 59

Gln Pro Asp Ser Val Ser Ile Pro Ile Thr Cys Cys Phe Asn Val Ile
 1 5 10 15

Asn Arg Lys Ile Pro Ile Gln Arg Leu Glu Ser Tyr Thr Arg Ile Thr
 20 25 30

Asn Ile Gln Cys Pro Lys Glu Ala Val Ile Phe Lys Thr Lys Arg Gly
 35 40 45

Lys Glu Val Cys Ala Asp Pro Lys Glu Arg Trp Val Arg Asp Ser Met
 50 55 60

Lys His Leu Asp Gln Ile Phe Gln Asn Leu Lys Pro

65

70

75

<210> 60
 <211> 76
 <212> PRT
 <213> Homo sapiens

<400> 60

Gln Pro Val Gly Ile Asn Thr Ser Thr Thr Cys Cys Tyr Arg Phe Ile
 1 5 10 15

Asn Lys Lys Ile Pro Lys Gln Arg Leu Glu Ser Tyr Arg Arg Thr Thr
 20 25 30

Ser Ser His Cys Pro Arg Glu Ala Val Ile Phe Lys Thr Lys Leu Asp
 35 40 45

Lys Glu Ile Cys Ala Asp Pro Thr Gln Lys Trp Val Gln Asp Phe Met
 50 55 60

Lys His Leu Asp Lys Lys Thr Gln Thr Pro Lys Leu
 65 70 75

<210> 61
 <211> 77
 <212> PRT
 <213> Homo sapiens

<400> 61

Gly Pro Val Ser Ala Val Leu Thr Glu Leu Arg Cys Thr Cys Leu Arg
 1 5 10 15

Val Thr Leu Arg Val Asn Pro Lys Thr Ile Gly Lys Leu Gln Val Phe
 20 25 30

Pro Ala Gly Pro Gln Cys Ser Lys Val Glu Val Val Ala Ser Leu Lys
 35 40 45

Asn Gly Lys Gln Val Cys Leu Asp Pro Glu Ala Pro Phe Leu Lys Lys
 50 55 60

Val Ile Gln Lys Ile Leu Asp Ser Gly Asn Lys Lys Asn
 65 70 75

<210> 62
 <211> 68

<212> PRT

<213> Homo sapiens

<400> 62

Lys Pro Val Ser Leu Ser Tyr Arg Cys Pro Cys Arg Phe Phe Glu Ser
 1 5 10 15

His Val Ala Arg Ala Asn Val Lys His Leu Lys Ile Leu Asn Thr Pro
 20 25 30

Asn Cys Ala Leu Gln Ile Val Ala Arg Leu Lys Asn Asn Asn Arg Gln
 35 40 45

Val Cys Ile Asp Pro Lys Leu Lys Trp Ile Gln Glu Tyr Leu Glu Lys
 50 55 60

Ala Leu Asn Lys
 65

<210> 63

<211> 72

<212> PRT

<213> Homo sapiens

<400> 63

Lys Pro Val Ser Leu Ser Tyr Arg Cys Pro Cys Arg Phe Phe Glu Ser
 1 5 10 15

His Val Ala Arg Ala Asn Val Lys His Leu Lys Ile Leu Asn Thr Pro
 20 25 30

Asn Cys Ala Leu Gln Ile Val Ala Arg Leu Lys Asn Asn Asn Arg Gln
 35 40 45

Val Cys Ile Asp Pro Lys Leu Lys Trp Ile Gln Glu Tyr Leu Glu Lys
 50 55 60

Ala Leu Asn Lys Arg Phe Lys Met
 65 70

<210> 64

<211> 69

<212> PRT

<213> Homo sapiens

<400> 64

Gly Pro Tyr Gly Ala Asn Met Glu Asp Ser Val Cys Cys Arg Asp Tyr
 1 5 10 15

Val Arg Tyr Arg Leu Pro Leu Arg Val Val Lys His Phe Tyr Trp Thr
 20 25 30

Ser Asp Ser Cys Pro Arg Pro Gly Val Val Leu Leu Thr Phe Arg Asp
 35 40 45

Lys Glu Ile Cys Ala Asp Pro Arg Val Pro Trp Val Lys Met Ile Leu
 50 55 60

Asn Lys Leu Ser Gln
 65

<210> 65
 <211> 7
 <212> PRT
 <213> Homo sapiens

<400> 65

Tyr Pro Phe Val Glu Pro Ile
 1 5

<210> 66
 <211> 95
 <212> PRT
 <213> Homo sapiens

<400> 66

Ala Pro Gly Pro Arg Gly Ile Ile Ile Asn Leu Glu Asn Gly Glu Leu
 1 5 10 15

Cys Met Asn Ser Ala Gln Cys Lys Ser Asn Cys Cys Gln His Ser Ser
 20 25 30

Ala Leu Gly Leu Ala Arg Cys Thr Ser Met Ala Ser Glu Asn Ser Glu
 35 40 45

Cys Ser Val Lys Thr Leu Tyr Gly Ile Tyr Tyr Lys Cys Pro Cys Glu
 50 55 60

Arg Gly Leu Thr Cys Glu Gly Asp Lys Thr Ile Val Gly Ser Ile Thr
 65 70 75 80

Asn Thr Asn Phe Gly Ile Cys His Asp Ala Gly Arg Ser Lys Gln

85

90

95

<210> 67
<211> 8
<212> PRT
<213> Homo sapiens

<400> 67

His Ser Asp Ala Val Phe Thr Asp
1 5

<210> 68
<211> 6
<212> PRT
<213> Homo sapiens

<400> 68

His Ser Asp Gly Ile Phe
1 5

<210> 69
<211> 8
<212> PRT
<213> Homo sapiens

<400> 69

His Ser Gln Gly Thr Phe Thr Ser
1 5

<210> 70
<211> 8
<212> PRT
<213> Homo sapiens

<400> 70

Phe Pro Thr Ile Pro Leu Ser Arg
1 5

<210> 71
<211> 8
<212> PRT
<213> Homo sapiens

<400> 71

His Ser Asp Gly Thr Phe Thr Ser
1 5

<210> 72

-24-

<211> 31
 <212> PRT
 <213> artificial

<220>
 <223> substituted GLP-1

<220>
 <221> MISC_FEATURE
 <222> (1)..(31)
 <223> X at position 2 = V or S; X at position 3 = Gln, D-Gln or Asp; X
 at position 10 = T or G; X at position 12 = K or A.

<400> 72

His Xaa Xaa Gly Thr Phe Thr Ser Asp Xaa Ser Xaa Tyr Leu Glu Gly
 1 5 10 15

Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly Arg Gly
 20 25 30

<210> 73
 <211> 31
 <212> PRT
 <213> artificial

<220>
 <223> substituted GLP-1

<220>
 <221> MISC_FEATURE
 <222> (1)..(31)
 <223> X at position 20 = G, S, C, T, N, Q, Y, A, V, I, L, M, F, A,
 d-Lys or K; X at position 28 = G, S, C, T, N, Q, Y, A, V, I, L,
 M, F, A, d-Lys or K; X at position 30 = G, S, C, T, N, Q, Y, A,
 V, I, L, M, F, K, R, d-Arg or -OH.

<220>
 <221> MISC_FEATURE
 <222> (1)..(31)
 <223> X at position 29 = G or -OH; X at position 31 = G or -OH.

<400> 73

His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu Gly
 1 5 10 15

Gln Ala Ala Xaa Glu Phe Ile Ala Trp Leu Val Xaa Xaa Xaa Xaa
 20 25 30

<210> 74
 <211> 31
 <212> PRT

<213> artificial

<220>

<223> substituted GLP-1

<220>

<221> MISC_FEATURE

<222> (1)..(31)

<223> X at position 25 = oxidation-resistant amino acid; X at position 29 = G or -OH; X at position 30 = R or -OH; X at position 31 = G or -OH.

<400> 74

His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu Gly
1 5 10 15

Gln Ala Ala Lys Glu Phe Ile Ala Xaa Leu Val Xaa Xaa Xaa Xaa
20 25 30

<210> 75

<211> 31

<212> PRT

<213> artificial

<220>

<223> substituted GLP-1

<220>

<221> MISC_FEATURE

<222> (1)..(31)

<223> X at position 10 = Y or V; X at position 12 = K or S; X at position 15 = D or E; X at position 16 = S or G; X at position 17 = R or Q; X at position 18 = R or A; X at position 20 = Q or K.

<220>

<221> MISC_FEATURE

<222> (1)..(31)

<223> X at position 29 = G or -OH; X at position 30 = R or -OH; X at position 31 = G or -OH.

<400> 75

His Ala Glu Gly Thr Phe Thr Ser Asp Xaa Ser Xaa Tyr Leu Xaa Xaa
1 5 10 15

Xaa Xaa Ala Xaa Glu Phe Ile Ala Trp Leu Val Xaa Xaa Xaa Xaa
20 25 30

<210> 76

<211> 31

<212> PRT

<213> artificial

<220>
 <223> substituted GLP-1

<220>
 <221> MISC_FEATURE
 <222> (1)..(31)
 <223> X at position 2 = G or S or C or A; X at position 3 = D or G or S or C or T or N or Q or Y or A or V or I or L or M or F or E; X at position 4 = S or C or T or N or Q or Y or A or V or I or L or M or F or G; X at position 9 = E or D.

<220>
 <221> MISC_FEATURE
 <222> (1)..(31)
 <223> X at position 29 = G or -OH; X at position 30 = R or -OH; X at position 31 = G or -OH.

<400> 76

His Xaa Xaa Xaa Thr Phe Thr Ser Xaa Val Ser Ser Tyr Leu Glu Gly
 1 5 10 15

Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Xaa Xaa Xaa Xaa
 20 25 30

<210> 77
 <211> 31
 <212> PRT
 <213> artificial

<220>
 <223> substituted GLP-1

<220>
 <221> MISC_FEATURE
 <222> (1)..(31)
 <223> X at position 1 = G or S or C or T or N or Q or Y or A or V or I or L or M or F or d-His or alkylated His or acylated His; X at position 29 = G or -OH; X at position 30 = R or -OH; X at position 31 = G or -OH.

<400> 77

Xaa Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu Gly
 1 5 10 15

Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Xaa Xaa Xaa Xaa
 20 25 30

<210> 78
 <211> 31
 <212> PRT

<213> artificial

<220>

<223> substituted GLP-1

<220>

<221> MISC_FEATURE

<222> (1)..(31)

<223> X at position 1 = l- or d-His, desamino-His, 2-amino-His, beta-hydroxy-His, homohistidine, alpha-fluoromethyl-His, and alpha-methyl-His; X at position 2 = A, G, V, T, I or alpha-methyl-Ala; X at positions 15 and 21 = E, Q, A, T, S or G.

<220>

<221> MISC_FEATURE

<222> (1)..(31)

<223> X at position 31 = Gly or NH2.

<400> 78

Xaa Xaa Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Xaa Gly
1 5 10 15

Gln Ala Ala Lys Xaa Phe Ile Ala Trp Leu Val Lys Gly Arg Xaa
20 25 30

<210> 79

<211> 39

<212> PRT

<213> Homo sapiens

<400> 79

His Ser Asp Gly Thr Phe Thr Ser Asp Leu Ser Lys Gln Met Glu Glu
1 5 10 15

Glu Ala Val Arg Leu Phe Ile Glu Trp Leu Lys Asn Gly Gly Pro Ser
20 25 30

Ser Gly Ala Pro Pro Pro Ser
35

<210> 80

<211> 39

<212> PRT

<213> Homo sapiens

<400> 80

His Gly Glu Gly Thr Phe Thr Ser Asp Leu Ser Lys Gln Met Glu Glu
1 5 10 15

Glu Ala Val Arg Leu Phe Ile Glu Trp Leu Lys Asn Gly Gly Pro Ser
 20 25 30

Ser Gly Ala Pro Pro Pro Ser
 35

<210> 81
 <211> 31
 <212> PRT
 <213> artificial

<220>
 <223> modified exendin-4

<400> 81

His Gly Glu Gly Thr Phe Thr Ser Asp Leu Ser Lys Gln Met Glu Glu
 1 5 10 15

Ala Val Arg Leu Phe Ile Glu Trp Leu Lys Asn Gly Gly Pro Tyr
 20 25 30

<210> 82
 <211> 4
 <212> PRT
 <213> artificial

<220>
 <223> modified GLP-1 N-terminus

<400> 82

His His Ala Glu
 1

<210> 83
 <211> 4
 <212> PRT
 <213> artificial

<220>
 <223> modified GLP-1 N-terminus

<400> 83

His His Gly Glu
 1

<210> 84
 <211> 4
 <212> PRT
 <213> artificial

<220>
<223> modified GLP-1 N-terminus

<400> 84

His His Ser Glu
1

<210> 85
<211> 4
<212> PRT
<213> artificial

<220>
<223> modified GLP-1 N-terminus

<400> 85

Gly His Ala Glu
1

<210> 86
<211> 4
<212> PRT
<213> artificial

<220>
<223> modified GLP-1 N-terminus

<400> 86

Gly His Gly Glu
1

<210> 87
<211> 4
<212> PRT
<213> artificial

<220>
<223> modified GLP-1 N-terminus

<400> 87

Gly His Ser Glu
1

<210> 88
<211> 32
<212> PRT
<213> artificial

<220>
<223> modified GLP-1

<400> 88

His His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu
 1 5 10 15

Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly Arg Gly
 20 25 30

<210> 89

<211> 32

<212> PRT

<213> artificial

<220>

<223> modified GLP-1

<220>

<221> MISC_FEATURE

<222> (1)..(31)

<223> X = an amino acid other than l- or d-Lys

<400> 89

His His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu
 1 5 10 15

Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Xaa Gly Arg Gly
 20 25 30

<210> 90

<211> 30

<212> PRT

<213> artificial

<220>

<223> A8G modified GLP-1

<400> 90

His Gly Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu Gly
 1 5 10 15

Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly Arg
 20 25 30

<210> 91

<211> 31

<212> PRT

<213> artificial

<220>

<223> modified GLP-1 with additional N-terminal His

<400> 91

His His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu
 1 5 10 15

Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly Arg
 20 25 30

<210> 92

<211> 31

<212> PRT

<213> artificial

<220>

<223> modified GLP-1 with additional N-terminal His and A8G

<400> 92

His His Gly Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu
 1 5 10 15

Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly Arg
 20 25 30

<210> 93

<211> 32

<212> PRT

<213> artificial

<220>

<223> modified GLP-1 with 2 additional N-terminal His residues

<400> 93

His His His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu
 1 5 10 15

Glu Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly Arg
 20 25 30

<210> 94

<211> 31

<212> PRT

<213> artificial

<220>

<223> modified GLP-1 with additional N-terminal Gly

<400> 94

Gly His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu
 1 5 10 15

Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly Arg
 20 25 30

<210> 95
 <211> 40
 <212> PRT
 <213> artificial

<220>
 <223> modified exendin-4 with additional N-terminal His

<400> 95

His His Gly Glu Gly Thr Phe Thr Ser Asp Leu Ser Lys Gln Met Glu
 1 5 10 15

Glu Glu Ala Val Arg Leu Phe Ile Glu Trp Leu Lys Asn Gly Gly Pro
 20 25 30

Ser Ser Gly Ala Pro Pro Pro Ser
 35 40

<210> 96
 <211> 40
 <212> DNA
 <213> artificial

<220>
 <223> primer for constructing modified GLP-1-modified Tf fusion protein

<400> 96
 ttcccatataca aacttaagag tccaattagc ttcacgcca 40

<210> 97
 <211> 60
 <212> DNA
 <213> artificial

<220>
 <223> primer for constructing modified GLP-1-modified Tf fusion protein

<400> 97
 ggtttagcctt gtttttttat tggcgatgaa gctaattgga ctcttaagtt tgtatgggaa 60

<210> 98
 <211> 60
 <212> DNA
 <213> artificial

<220>
 <223> primer for constructing modified GLP-1-modified Tf fusion protein

<400> 98
ataaaaaaac aagctaaacc taattctaac aagcaaagat gaggctcgcc gtgggagccc 60

<210> 99
<211> 60
<212> DNA
<213> artificial

<220>
<223> primer for constructing modified GLP-1-modified Tf fusion protein

<400> 99
caggacggcg cagaccagca gggctccac ggcgagcctc atctttgctt gttagaatta 60

<210> 100
<211> 70
<212> DNA
<213> artificial

<220>
<223> primer for constructing modified GLP-1-modified Tf fusion protein

<400> 100
tgctggtctg cgccgtcctg gggctgtgtc tggcgcacatg tgaaggtact tttacttctg 60
atgtttcttc 70

<210> 101
<211> 80
<212> DNA
<213> artificial

<220>
<223> primer for constructing modified GLP-1-modified Tf fusion protein

<400> 101
aattctttag cagcttgacc ttcaaataa gaagaaacat cagaagtaaa agtaccttca 60
gcatgcgcca gacacagccc 80

<210> 102
<211> 70
<212> DNA
<213> artificial

<220>
<223> primer for constructing modified GLP-1-modified Tf fusion protein

<400> 102
ttatttggaa ggtcaagctg ctaaagaatt tattgcttgg ttggttaaag gtagggatacc 60
tgataaaact 70

<210> 103
 <211> 40
 <212> DNA
 <213> artificial

<220>
 <223> primer for constructing modified GLP-1-modified Tf fusion protein

<400> 103
 agttttatca ggtaccctac ctttaaccaa ccaagcaata 40

<210> 104
 <211> 300
 <212> DNA
 <213> artificial

<220>
 <223> sequence encoding modified GLP-1-modified Tf fusion protein

<220>
 <221> CDS
 <222> (109)..(300)

<400> 104
 ccaatgttac gtccogttat attggagttc ttccataca aacttaagag tccaattagc 60

ttcatcgcca ataaaaaaaaac aagctaaacc taattctaac aagcaaag atg agg ctc 117
 Met Arg Leu
 1

gcc gtg gga gcc ctg ctg gtc tgc gcc gtc ctg ggg ctg tgt ctg gcg 165
 Ala Val Gly Ala Leu Leu Val Cys Ala Val Leu Gly Leu Cys Leu Ala
 5 10 15

cat gct gaa ggt act ttt act tct gat gtt tct tct tat ttg gaa ggt 213
 His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr Leu Glu Gly
 20 25 30 35

caa gct gct aaa gaa ttt att gct tgg ttg gtt aaa ggt agg gta cct 261
 Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly Arg Val Pro
 40 45 50

gat aaa act gtg aga tgg tgt gca gtg tcg gag cat gag 300
 Asp Lys Thr Val Arg Trp Cys Ala Val Ser Glu His Glu
 55 60

<210> 105
 <211> 64
 <212> PRT
 <213> artificial

<220>
 <223> sequence encoding modified GLP-1-modified Tf fusion protein

<400> 105

Met Arg Leu Ala Val Gly Ala Leu Leu Val Cys Ala Val Leu Gly Leu
 1 5 10 15

Cys Leu Ala His Ala Glu Gly Thr Phe Thr Ser Asp Val Ser Ser Tyr
 20 25 30

Leu Glu Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu Val Lys Gly
 35 40 45

Arg Val Pro Asp Lys Thr Val Arg Trp Cys Ala Val Ser Glu His Glu
 50 55 60

<210> 106

<211> 25

<212> DNA

<213> artificial

<220>

<223> primer for constructing plasmid with natural Tf secretion signal

<400> 106

ctgtgtctgg cgcacatgc tgaag 25

<210> 107

<211> 25

<212> DNA

<213> artificial

<220>

<223> primer for constructing plasmid with natural Tf secretion signal

<400> 107

cttcagcatg atgcgccaga cacag 25

<210> 108

<211> 30

<212> PRT

<213> artificial

<220>

<223> modified glucagon with additional N-terminal His

<400> 108

His His Ser Gln Gly Thr Phe Thr Ser Asp Tyr Ser Lys Val Leu Asp
 1 5 10 15

Ser Arg Arg Ala Gln Asp Phe Val Gln Trp Leu Met Asn Thr
 20 25 30

<210> 109

<211> 43
<212> PRT
<213> artificial

<220>
<223> modified GIP with additional N-terminal Tyr

<400> 109

Tyr Tyr Ala Glu Gly Thr Phe Ile Ser Asp Tyr Ser Ile Ala Met Asp
1 5 10 15

Lys Ile His Gln Gln Asp Phe Val Asn Trp Leu Leu Ala Gln Lys Gly
20 25 30

Lys Lys Asn Asp Trp Lys His Asn Ile Thr Gln
35 40

<210> 110
<211> 5
<212> PRT
<213> artificial

<220>
<223> dipeptidyl peptidase serine protease motif

<400> 110

Gly Trp Ser Tyr Gly
1 5

<210> 111
<211> 6
<212> PRT
<213> artificial

<220>
<223> transferrin secretion signal sequence

<400> 111

Arg Ser Leu Asp Lys Arg
1 5

<210> 112
<211> 6
<212> PRT
<213> artificial

<220>
<223> transferrin secretion signal sequence

<400> 112

Arg Ser Leu Asp Arg Arg
1 5

<210> 113
<211> 6
<212> PRT
<213> artificial

<220>
<223> transferrin secretion signal sequence

<400> 113

Arg Ser Leu Glu Lys Arg
1 5

<210> 114
<211> 6
<212> PRT
<213> artificial

<220>
<223> transferrin secretion signal sequence

<400> 114

Arg Ser Leu Glu Arg Arg
1 5

<210> 115
<211> 4
<212> PRT
<213> artificial

<220>
<223> DPP-resistant N-terminus

<400> 115

His His Ala Glu
1

<210> 116
<211> 4
<212> PRT
<213> artificial

<220>
<223> DPP-resistant N-terminus

<400> 116

Gly His Ala Glu
1

<210> 117
 <211> 90
 <212> DNA
 <213> artificial

<220>
 <223> DNA sequence encoding GLP-1(7-36)

<400> 117
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 gaatttattg cttggttggt taaaggtaga 90

<210> 118
 <211> 118
 <212> DNA
 <213> artificial

<220>
 <223> site of pREX0052 with GLP-1 insert

<220>
 <221> CDS
 <222> (1)..(117)

<400> 118
 agg tct cta gag aaa agg cat gct gaa ggt act ttt act tct gat gtt 48
 Arg Ser Leu Glu Lys Arg His Ala Glu Gly Thr Phe Thr Ser Asp Val
 1 5 10 15
 tct tct tat ttg gaa ggt caa gct gct aaa gaa ttt att gct tgg ttg 96
 Ser Ser Tyr Leu Glu Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu
 20 25 30
 gtt aaa ggt agg gta cct gat a 118
 Val Lys Gly Arg Val Pro Asp
 35

<210> 119
 <211> 39
 <212> PRT
 <213> artificial

<220>
 <223> site of pREX0052 with GLP-1 insert

<400> 119
 Arg Ser Leu Glu Lys Arg His Ala Glu Gly Thr Phe Thr Ser Asp Val
 1 5 10 15
 Ser Ser Tyr Leu Glu Gly Gln Ala Ala Lys Glu Phe Ile Ala Trp Leu
 20 25 30

Val Lys Gly Arg Val Pro Asp
35