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(54) **THERAPEUTIC AGENTS COMPRISING ELASTIN-LIKE PEPTIDES**

(57) The present invention provides therapeutic agents and compositions comprising elastin-like peptides (ELPs) and therapeutic protein is a GLP-1 receptor agonist, insulin, or Factor VII/VIIa, including functional analogs. The present invention further provides encoding polynucleotides, as well as methods of making and using

the therapeutic agents. The therapeutic agents have improvements in relation to their use as therapeutics, including, *inter alia*, one or more of half-life, clearance and/or persistence in the body, solubility, and bioavailability.

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Description

PRIORITY

5 **[0001]** This application claims priority to US Provisional Application No. 61/076,221, filed June 27, 2008, which is hereby incorporated by reference in its entirety.

FIELD OF THE INVENTION

10 **[0002]** This application discloses therapeutic agents comprising elastin-like-peptides, and is related to PCT/US2007/077767 (published as WO 2008/030968 on March 13, 2008) having an International Filing Date of September 6, 2007. This application is also related to PCT/US2006/048572 (published as WO 2007/073486 on June 28, 2007) having an International Filing Date of December 20, 2006. WO 2008/030968 and WO 2007/073486 are each hereby incorporated by reference in their entirety.

DESCRIPTION OF THE TEXT FILE SUBMITTED ELECTRONICALLY

15 **[0003]** The contents of the text file submitted electronically herewith are incorporated herein by reference in their entirety: A computer readable format copy of the Sequence Listing (filename: PHAS_010_01US_SeqList_ST25.txt, date recorded: June 26, 2009, file size 50 kb).

BACKGROUND OF THE INVENTION

25 **[0004]** Therapeutic proteins or peptides in their native state or when recombinantly produced can be labile molecules exhibiting, *inter alia*, short periods of serum stability, serum half-life (i.e., circulatory half-life), or limited persistence in the body. Such molecules can also be extremely labile when formulated, such as when formulated in aqueous solutions.

[0005] In some instances, polyethylene glycol (PEG) conjugated to a proteinaceous molecule results in a longer-acting, sustained activity of the molecule. PEG attachment, however, can often substantially reduce or even destroy the protein's therapeutic activity. Therapeutic proteins and/or peptides have also been stabilized by fusion to certain proteins that are capable of extending serum half-life. For example, in some instances, therapeutic proteins fused to albumin, transferrin, and antibody fragments exhibit extended serum half-life when compared to the therapeutic protein in the unfused state. See US Patent No. 7,238,667 (particularly with respect to albumin conjugates), US Patent No. 7,176,278 (particularly with respect to transferrin conjugates), and US Patent No. 5,766,883, which are each hereby incorporated by reference in their entirety.

35 **[0006]** There remains a need in the art for more stable, longer acting, and/or effective proteinaceous molecules.

SUMMARY OF THE INVENTION

40 **[0007]** The present invention provides therapeutic agents comprising an elastin-like peptide (ELP) component and a therapeutic proteinaceous component. The ELP component contains structural peptide units, which may be repeating units, structurally related to, or derived from, sequences of the elastin protein. Such ELP components provide certain therapeutic advantages to the therapeutic agent, such as comparatively better stability, solubility, bioavailability, half-life, persistence, and/or biological action of the therapeutic proteinaceous component. Such properties may be determined, for example, with respect to the therapeutic component's unfused or unconjugated counterpart. In some embodiments, the ELP components may undergo a reversible inverse phase transition, which may impart additional practical and/or therapeutic advantages. The invention further provides polynucleotides encoding the therapeutic agents of the invention, as well as methods of treatment or prophylaxis for certain biological conditions.

45 **[0008]** In a first aspect, the invention provides a therapeutic agent comprising an elastin-like peptide (ELP) component and a therapeutic proteinaceous component, as well as pharmaceutical compositions containing the same for delivery to a subject or patient in need. The therapeutic component may be selected from active portions of the therapeutic proteins listed in Table 1, or functional analogs thereof. In certain embodiments, the therapeutic component is a GLP-1 receptor agonist, such as GLP-1, exendin-4, or a functional analog thereof. Such therapeutic components are generally effective for, among other things, increasing insulin secretion from the pancreas in a glucose-dependent manner. In other embodiments, the therapeutic component is an insulin or functional analog thereof, which is generally effective for promoting glucose uptake from the blood and storage within cells. In still other embodiments, the therapeutic component is a Factor VII/VIIa or functional analog thereof, which is generally effective for promoting coagulation by activation of Factor X or Factor IX.

55 **[0009]** The ELP and therapeutic components may be covalently coupled by various means, including chemical coupling

(e.g., conjugation) and recombinant fusion technology. In addition, the number of ELP or therapeutic components per molecule, and their respective positions within the molecule, may vary as needed. The therapeutic agent may further include one or more spacer or linker moieties, which in addition to providing the desired functional independence of the ELP and therapeutic components, may optionally provide for additional functionalities, such as a protease-sensitive feature to allow for proteolytic release or activation of the therapeutic component. The therapeutic agent may further include one or more targeting components such as, for example, a peptide or protein to target the therapeutic agent to a particular cell type, e.g., a cancer cell, or to a particular organ.

[0010] In a second aspect, the invention provides polynucleotides, such polynucleotides comprising a nucleotide sequence encoding a therapeutic agent of the invention. For example, the nucleotide sequence encodes an ELP fusion with a functional portion of at least one therapeutic protein listed in Table 1 (or functional analog thereof). In certain embodiments, the therapeutic component is a GLP-1 receptor agonist (including GLP-1 and exendin-4), insulin, Factor VII/VIIa, or functional analog thereof. Such polynucleotides may further comprise additional control element(s) operably linked to the nucleotide sequence, such as promoter elements and/or other transcription or expression-related signals. The polynucleotide may be inserted into various vectors, which may be useful for production of the therapeutic agent in host cells, including, for example, bacterial and eukaryotic host cells.

[0011] In a third aspect, the invention provides a method for treating or preventing a disease, disorder, or condition in a subject, such as in a mammalian patient, including a human patient. The method comprises administering an effective amount of the therapeutic agent of the invention (or pharmaceutical composition containing the same) to a subject or patient in need thereof. For example, the patient may be in need of an agent having a biological activity or preferred indication listed in Table 1. In certain embodiments employing a GLP-1 receptor agonist / ELP compound or employing an insulin/ELP compound, the invention provides a method for treating one or more disorders including type 1 or type 2 diabetes, hyperglycemia, and impaired glucose tolerance. In certain other embodiments employing Factor VII/VIIa / ELP compound, the invention provides a method for treating one or more disorders including hemophilia, post-surgical bleeding, anticoagulation-induced bleeding, thrombocytopenia, factor VII deficiency, factor XI deficiency, and intracranial hemorrhage.

[0012] Various other aspects, features and embodiments of the invention will be more fully apparent from the following disclosure and appended claims.

BRIEF DESCRIPTION OF THE FIGURES

[0013]

Figure 1 depicts plasmid pET24d-ELP1-90, encoding an ELP component with a 10 unit VPGXG (SEQ ID NO: 3) repeat motif, where guest position X is V, G, and A in the ratio of 5:3:2. This motif is repeated eight times with a final C-terminal 10-unit repeat where X is V, G, A, and W in the ratio 4:3:2:1. This ELP component is represented generally as $[(VPGXG)_{10}]_9$.

Figure 2A depicts plasmid pET24d-Ex-4 ELP1-90 encoding an ELP component with VPGXG (SEQ ID NO: 3) repeat motif (as in Fig. 1) cloned in frame with an N-terminal exendin-4 component. **Figure 2B** depicts the nucleotide and amino acid sequence of the exendin-4/ELP fusion (SEQ ID NOS: 23 and 24). Primer sequences are indicated (SEQ ID NOS:35-40).

Figure 3A depicts the nucleotide and amino acid sequence of an exendin-4 construct having an N-terminal Tev (Tobacco Etch Virus cysteine protease) cleavage site (SEQ ID NOS: 25 and 26). Primer sequences are indicated (SEQ ID NOS:38, 41, 42). **Figure 3B** also depicts the nucleotide and amino acid sequence of an exendin-4 construct having an N-terminal Tev cleavage site, but with an additional sequence N-terminal to the Tev cleavage site to provide a better target for the protease (SEQ ID NOS: 27 and 28). Primer sequences are indicated (SEQ ID NOS:38, 43,44).

Figure 4A depicts the nucleotide and amino acid sequence of an exendin-4/ELP fusion as in Figures 1-3, but with a DsbA leader sequence to direct secretion into the periplasmic space (SEQ ID NOS: 29 and 30). Primer sequences are indicated (SEQ ID NOS:38, 45, 46). **Figure 4B** depicts plasmid pET24d-DsbA-Ex-4 ELP1-90 encoding the fusion of Fig. 4A.

Figure 5A depicts pPB0868, which encodes GLP-1(A8G,7-37)ELP1-90. **Figure 5B** depicts the nucleotide and amino acid sequence of the encoded fusion protein (SEQ ID NOS: 53 and 54, respectively).

Figure 6A depicts pPB1022, which encodes GLP-1(A8G,7-37)ELP1-120. **Figure 6B** depicts the nucleotide and

amino acid sequence of the encoded fusion protein (SEQ ID NOS: 55 and 56, respectively).

Figure 7A depicts pPB0788, which encodes Factor VII-ELP1-90. **Figure 7B** depicts the nucleotide and amino acid sequence of the encoded fusion protein (SEQ ID NOS: 57 and 58, respectively).

Figure 8A depicts the nucleotide and amino acid sequence of an insulin (B, C, and A chains) having the ELP component cloned in frame (SEQ ID NOS: 31 and 32). Primer sequences are indicated (SEQ ID NOS: 47 and 48). **Figure 8B** depicts plasmid pET24d Insulin-ELP1-90 expressing the insulin/ELP fusion of Fig. 8A.

Figure 9 is a Western blot for FVII-ELP1-90 from transient transfection of Freestyle HEK293, detected with mouse anti-human FVII monoclonal antibody. Lanes are: (1) culture media; (2) FVII ELP1-90 after purification by phase transition; and FVII control.

Figure 10 is an SDS-PAGE showing recombinant production of an Exendin-4/ELP4-60 fusion. Lanes are: (M) Protein markers; (1) Exendin-4 ELP4-60 from total lysate; (2) Exendin-4 ELP4-60 from insoluble lysate; (3) Exendin-4 ELP4-60 from soluble lysate; (4) Exendin-4 ELP4-60 from 1st transition (equal volume); (5) Exendin-4 ELP4-60 from 2nd transition (concentrated); (6) Exendin-4 ELP4-60 from 3rd transition (concentrated).

Figure 11 shows the activation of Factor X by FactorVIIa-ELP1-90, and by Factor VIIa as a comparison. As shown, FactorVIIa-ELP retains full activity.

Figure 12 shows that Factor VIIa-ELP1-90 has a long PK when administered by i.v. in rats. FactorVIIa has a $T_{1/2}$ of about 690 min. as compared to about 45-60 min. for Factor VIIa.

Figure 13 shows the high *in vitro* activity of GLP1-ELP and Exendin-4-ELP, when compared to the activity of Exendin peptide.

Figure 14 shows that GLP1-ELP has a $T_{1/2}$ of about 12.9 hours when administered by i.v. to rats, and a $T_{1/2}$ of about 8.6 hours when administered subcutaneously (SQ).

Figure 15 shows that GLP-1 ELP has a long half-life in rabbits of about 20 hours when administered i.v., and about 24 hours when administered sub-cutaneously.

Figure 16 shows sustained glycemic control in diabetic mice with GLP-1-ELP.

DETAILED DESCRIPTION OF THE INVENTION

[0014] The present invention provides therapeutic agents comprising an elastin-like peptide (ELP) ("the ELP component") and a therapeutic component. The therapeutic component may be selected from Table 1 (e.g., selected from a Therapeutic Protein, or functional portion or functional analog thereof, listed in Table 1). In certain embodiments, the therapeutic component is a GLP-1 receptor agonist, such as GLP-1 or exendin-4, or may be insulin, Factor VII/VIIa, or functional analog thereof. The ELP component contains structural units related to, or derived from, sequences of the elastin protein, which provides certain therapeutic advantages, such as comparatively better persistence, stability, solubility, bioavailability, half-life, and/or biological action of the therapeutic component. Such properties may be determined with respect to, for example, an unfused or unconjugated counterpart of the therapeutic component. The invention further provides polynucleotides encoding the therapeutic agents of the invention, as well as methods of treatment or prophylaxis for certain biological conditions, including the preferred indications listed in Table 1, and including diabetes (e.g., Type I and Type II), hyperglycemia, bleeding, hemophilia, and hemorrhage, among others.

[0015] For ease of reference in the ensuing discussion, set out below are definitions of some terms appearing in the discussion.

[0016] As used herein, the term "therapeutic agent" or "therapeutic component" refers to an agent or component capable of inducing a biological effect *in vivo* and/or *in vitro*. The biological effect may be useful for treating and/or preventing a condition, disorder, or disease in a subject or patient.

[0017] As used herein, the term "coupled" means that the specified components are either directly covalently bonded to one another (e.g., via chemical conjugation or recombinant fusion technology), or indirectly covalently joined to one another (e.g., via chemical conjugation or recombinant fusion technology) through an intervening moiety or moieties, such as a bridge, spacer, or linker.

[0018] As used herein, "half-life" (which generally refers to *in vivo* half-life or circulatory half-life) is the period of time

that is required for a 50% diminution of bioactivity of the active agent to occur. Such term is to be contrasted with "persistence," which is the overall temporal duration of the active agent in the body, and "rate of clearance" as being a dynamically changing variable that may or may not be correlative with the numerical values of half-life and persistence.

[0019] The term "functional analog" refers to a protein that is an active analog (e.g., either chemical or protein analog), derivative, fragment, truncation isoform or the like of a native protein. For example, the functional analog may be a functional analog of a therapeutic protein listed in Table 1, or may be a functional analog of a GLP-1 receptor agonist (e.g., GLP-1, exendin), insulin, or Factor VII/VIIa. A polypeptide is active when it retains some or all of the biological activity of the corresponding native polypeptide, as determined *in vivo* or in one or more indicative *in vitro* assays. Exemplary activity assays for certain therapeutic proteins, which are determinative of activity, are listed Table 1. Further, such biological activities and assays for GLP-1 receptor agonists, insulin, and Factor VII/VIIa, which are determinative of whether a given molecule is a "functional analog," are described in detail elsewhere herein.

[0020] As used herein, the term "native," as used in reference to an amino acid sequence, indicates that the amino acid sequence is found in a naturally-occurring protein.

[0021] As used herein, the term "spacer" refers to any moiety, peptide or other chemical entity, that may be interposed between the ELP component and the therapeutic component. For example, the spacer may be a divalent group that is covalently bonded at one terminus to the ELP component, and covalently bonded at the other terminus to the therapeutic component. The therapeutic agents may therefore be open to the inclusion of additional chemical structure that does not preclude the efficacy of the agent for its intended purpose. The spacer may, for example, be a protease-sensitive spacer moiety that is provided to control the pharmacokinetics of the agent, or the spacer may be a protease-resistant moiety.

[0022] The therapeutic component and the ELP component may be coupled with one another in any suitable covalent manner, including chemical coupling and recombinant technology, such that the therapeutic agent is efficacious for its intended purpose, and such that the presence of the ELP component enhances the therapeutic component in some functional, therapeutic or physiological aspect. For example, the ELP-coupled therapeutic component may be enhanced in, e.g., its bioavailability, bio-unavailability, therapeutically effective dose, biological action, formulation compatibility, resistance to proteolysis or other degradative modality, solubility, half-life or other measure of persistence in the body subsequent to administration, rate of clearance from the body subsequent to administration, etc. Such enhancement may be determined, for example, in relation to a corresponding unconjugated or unfused counterpart therapeutic (e.g., determined relative to native GLP-1, exendin, insulin, or Factor VII/VIIa, or a therapeutic protein listed in Table 1).

[0023] In some embodiments, the therapeutic agent of the invention circulates or exists in the body in a soluble form, and escapes filtration by the kidney thereby persisting in the body in an active form. In some embodiments, the therapeutic agents of the invention have a molecular weight of less than the generally recognized cut-off for filtration through the kidney, such as less than about 60 kD, or in some embodiments less than about 55, 50, 45, 40, 30, or 20 kDa, and persist in the body by at least 2-fold, 3-fold, 4-fold, 5-fold, 10-fold, 20-fold, or 100-fold longer than an uncoupled (e.g., unfused or unconjugated) therapeutic counterpart.

[0024] The number of ELP and/or therapeutic components per molecule, and their respective positions within the molecule, may vary among embodiments of the invention. For example, in embodiments where the agent is a recombinant fusion, at least one ELP component may be placed at one or both of the N-terminus and the C-terminus. Where the ELP component is at both the N-terminus and C-terminus of the fusion, the ELP components will flank the therapeutic component. Alternatively, the therapeutic component may be positioned at either or both of the N-terminus and C-terminus. Where the therapeutic component is at both the N-terminus and C-terminus, the therapeutic component will flank the ELP component. In a further embodiment, different therapeutic components are positioned at the N-terminus and C-terminus of the molecule. As discussed in detail herein, in certain embodiments, such therapeutic component(s) may be released by proteolysis of a spacer moiety separating the ELP and therapeutic components. In certain embodiments, the therapeutic component may be inactive in the fused state, and becoming active upon proteolytic release from the ELP component(s). Alternatively, the therapeutic component remains active in the fused state, making proteolytic processing of the therapeutic agent unnecessary for biological activity.

[0025] When prepared as recombinant fusions, the therapeutic agent can be prepared by known recombinant expression techniques. For example, to recombinantly produce the therapeutic agent, a nucleic acid sequence encoding the chimeric gene is operatively linked to a suitable promoter sequence such that the nucleic acid sequence encoding such fusion protein will be transcribed and/or translated into the desired fusion protein in the host cells. Preferred promoters are those useful for expression in *E. coli*, such as the T7 promoter. Any commonly used expression system may be used, including eukaryotic or prokaryotic systems. Specific examples include yeast (e.g., *Saccharomyces spp.*, *Pichia spp.*), baculovirus, mammalian, and bacterial systems, such as *E. coli*, and *Caulobacter*.

[0026] The various aspects and embodiments of the invention are described in greater detail in the following sections.

Elastin-Like Peptide (ELP) Component

[0027] The therapeutic agent of the invention may comprise one or more ELP components. The ELP components comprise or consist of structural peptide units or sequences that are related to, or derived from, the elastin protein. Such sequences are useful for improving the properties of therapeutic proteins, such as those listed in Table 1, as well as GLP-1 receptor agonists (e.g., GLP-1 or exendin-4), insulin, and Factor VII/VIIa in one or more of bioavailability, therapeutically effective dose, biological action, formulation compatibility, resistance to proteolysis, solubility, half-life or other measure of persistence in the body subsequent to administration, and/or rate of clearance from the body.

[0028] The ELP component is constructed from structural units of from three to about twenty amino acids, or in some embodiments, from four to ten amino acids, such as five or six amino acids. The length of the individual structural units, in a particular ELP component, may vary or may be uniform. In certain embodiments, the ELP component is constructed of a polytetra-, polypenta-, polyhexa-, polyhepta-, polyocta-, and polynona-peptide motif of repeating structural units. Exemplary structural units include units defined by SEQ ID NOS: 1-12 (below), which may be employed as repeating structural units, including tandem-repeating units, or may be employed in some combination, to create an ELP effective for improving the properties of the therapeutic component. Thus, the ELP component may comprise or consist essentially of structural unit(s) selected from SEQ ID NOS: 1-12, as defined below.

[0029] The ELP component, comprising such structural units, may be of varying sizes. For example, the ELP component may comprise or consist essentially of from about 10 to about 500 structural units, or in certain embodiments about 15 to about 150 structural units, or in certain embodiments from about 20 to about 100 structural units, or from about 50 to about 90 structural units, including one or a combination of units defined by SEQ ID NOS: 1-12. Thus, the ELP component may have a length of from about 50 to about 2000 amino acid residues, or from about 100 to about 600 amino acid residues, or from about 200 to about 500 amino acid residues, or from about 200 to about 400 amino acid residues.

[0030] In some embodiments, the ELP component, or in some cases the therapeutic agent, has a size of less than about 65 kDa, or less than about 60 kDa, or less than about 55 kDa, or less than about 50 kDa, or less than about 40 kDa, or less than about 30 or 25 kDa. Three major blood proteins, Human Serum Albumin (HSA), Transferrin (Tf) and IgG, or the Fc portion of IgGs in their glycosylated form, have been exploited to extend the half-lives of proteins and peptides for improved therapeutic use. These molecules are 585, 679 and 480 amino acids in length giving molecular weights of about 66, 77, and ~75 kDa (including glycosylations), respectively. They are each globular and relatively compact. The half life of these molecules is determined by a number of factors, including charge distribution, rescue of molecules by the neonatal Fc receptor (FcRn) (HSA and Fc) or cycling of Tf through the Tf receptor (TfR), and their size which prevents filtering through the kidney glomerulus. HSA is slightly below the generally regarded cut-off for filtration through the kidney (~70 kDa) but its charge distribution helps prevent this. It would be anticipated that, in order to achieve half-life extension of the same order as that achieved with HSA, Tf and Fc, a protein of at least this molecular weight range would be required or desirable, i.e. having over 550 amino acids and being over 65 kDa. However, an ELP with a small number of amino acids relative to HSA, Tf and Fc (e.g., in the range of about 300 to 400) and around 30 to 40 kDa may have a half life that matches and/or exceeds that of HSA, Tf, and Fc.

[0031] In some embodiments, the ELP component in the untransitioned state may have an extended, relatively unstructured and non-globular form, and thus such molecules may have a large expanded structure in comparison to HSA, Tf and Fc, so as to escape kidney filtration. In such embodiments, the therapeutic agents of the invention have a molecular weight of less than the generally recognized cut-off for filtration through the kidney, such as less than about 60 kD, or in some embodiments less than about 55, 50, 45, 40, 30, or 25 kDa, and persist in the body by at least 2-fold, 3-fold, 4-fold, 5-fold, 10-fold, 20-fold, or 100-fold longer than an uncoupled (e.g., unfused or unconjugated) therapeutic counterpart.

[0032] In certain embodiments, the ELP component undergoes a reversible inverse phase transition. That is, the ELP components are structurally disordered and highly soluble in water below a transition temperature (Tt), but exhibit a sharp (2-3°C range) disorder-to-order phase transition when the temperature is raised above the Tt, leading to desolvation and aggregation of the ELP components. For example, the ELP forms insoluble polymers, when reaching sufficient size, which can be readily removed and isolated from solution by centrifugation. Such phase transition is reversible, and isolated insoluble ELPs can be completely resolubilized in buffer solution when the temperature is returned below the Tt of the ELPs. Thus, the therapeutic agents of the invention can, in some embodiments, be separated from other contaminating proteins to high purity using inverse transition cycling procedures, e.g., utilizing the temperature-dependent solubility of the therapeutic agent, or salt addition to the medium. Successive inverse phase transition cycles can be used to obtain a high degree of purity. In addition to temperature and ionic strength, other environmental variables useful for modulating the inverse transition of the therapeutic agents include pH, the addition of inorganic and organic solutes and solvents, side-chain ionization or chemical modification, and pressure.

[0033] In certain embodiments, the ELP component does not undergo a reversible inverse phase transition, or does not undergo such a transition at a biologically relevant Tt, and thus the improvements in the biological and/or physiological properties of the molecule (as described elsewhere herein), may be entirely or substantially independent of any phase transition properties. Nevertheless, such phase transition properties may impart additional practical advantages, for

example, in relation to the recovery and purification of such molecules.

[0034] In certain embodiments, the ELP component(s) may be formed of structural units, including but not limited to:

(a) the tetrapeptide Val-Pro-Gly-Gly, or VPGG (SEQ ID NO: 1);

(b) the tetrapeptide Ile-Pro-Gly-Gly, or IPGG (SEQ ID NO: 2);

(c) the pentapeptide Val-Pro-Gly-X-Gly (SEQ ID NO: 3), or

VPGXG, where X is any natural or non-natural amino acid residue, and where X optionally varies among polymeric or oligomeric repeats;

(d) the pentapeptide Ala-Val-Gly-Val-Pro, or AVGVP (SEQ ID NO: 4);

(e) the pentapeptide Ile-Pro-Gly-X-Gly, or IPGXG (SEQ ID NO: 5), where X is any natural or non-natural amino acid residue, and where X optionally varies among polymeric or oligomeric repeats;

(e) the pentapeptide Ile-Pro-Gly-Val-Gly, or IPGVG (SEQ ID NO: 6);

(f) the pentapeptide Leu-Pro-Gly-X-Gly, or LPGXG (SEQ ID NO: 7), where X is any natural or non-natural amino acid residue, and where X optionally varies among polymeric or oligomeric repeats;

(g) the pentapeptide Leu-Pro-Gly-Val-Gly, or LPGVG (SEQ ID NO: 8);

(h) the hexapeptide Val-Ala-Pro-Gly-Val-Gly, or VAPGVG (SEQ ID NO: 9);

(I) the octapeptide Gly-Val-Gly-Val-Pro-Gly-Val-Gly, or GVGVPGVG (SEQ ID NO: 10);

(J) the nonapeptide Val-Pro-Gly-Phe-Gly-Val-Gly-Ala-Gly, or VPGFVGAG (SEQ ID NO: 11); and

(K) the nonapeptides Val-Pro-Gly-Val-Gly-Val-Pro-Gly-Gly, or VPGVGVPGG (SEQ ID NO: 12).

Such structural units defined by SEQ ID NOS:1-12 may form structural repeat units, or may be used in combination to form an ELP component in accordance with the invention. In some embodiments, the ELP component is formed entirely (or almost entirely) of one or a combination of (e.g., 2, 3 or 4) structural units selected from SEQ ID NOS: 1-12. In other embodiments, at least 75%, or at least 80%, or at least 90% of the ELP component is formed from one or a combination of structural units selected from SEQ ID NOS: 1-12, and which may be present as repeating units.

[0035] In certain embodiments, the ELP component(s) contain repeat units, including tandem repeating units, of the pentapeptide Val-Pro-Gly-X-Gly (SEQ ID NO:3), where X is as defined above, and where the percentage of Val-Pro-Gly-X-Gly (SEQ ID NO:3) pentapeptide units taken with respect to the entire ELP component (which may comprise structural units other than VPGXG (SEQ ID NO:3)) is greater than about 75%, or greater than about 85%, or greater than about 95% of the ELP component. The ELP component may contain motifs having a 5 to 15-unit repeat (e.g. about 10-unit repeat) of the pentapeptide of SEQ ID NO: 3, with the guest residue X varying among at least 2 or at least 3 of the units. The guest residues may be independently selected, such as from the amino acids V, I, L, A, G, and W (and may be selected so as to retain a desired inverse phase transition property). The repeat motif itself may be repeated, for example, from about 5 to about 12 times, such as about 8 to 10 times, to create an exemplary ELP component. The ELP component as described in this paragraph may of course be constructed from any one of the structural units defined by SEQ ID NOS: 1-12, or a combination thereof.

[0036] In some embodiments, the ELP component may include a β -turn structure. Exemplary peptide sequences suitable for creating a β -turn structure are described in International Patent Application PCT/US96/05186, which is hereby incorporated by reference in its entirety. For example, the fourth residue (X) in the elastin pentapeptide sequence, VPGXG (SEQ ID NO:3), can be altered without eliminating the formation of a β -turn. Alternatively, the ELP component may lack a β -turn, or otherwise have a different conformation and/or folding character.

[0037] In certain embodiments, the ELP components include polymeric or oligomeric repeats of the pentapeptide VPGXG (SEQ ID NO: 3), where the guest residue X is any amino acid. X may be a naturally occurring or non-naturally occurring amino acid. In some embodiments, X is selected from alanine, arginine, asparagine, aspartic acid, cysteine, glutamic acid, glutamine, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, serine, threonine, tryptophan, tyrosine and valine. In some embodiments, X is a natural amino acid other than proline or cysteine.

[0038] The guest residue X (e.g., with respect to SEQ ID NO: 3, or other ELP structural unit) may be a non-classical (non-genetically encoded) amino acid. Examples of non-classical amino acids include: D- isomers of the common amino acids, 2,4-diaminobutyric acid, α -amino isobutyric acid, A- aminobutyric acid, Abu, 2-amino butyric acid, γ -Abu, ϵ -Ahx, 6-amino hexanoic acid, Aib, 2-amino isobutyric acid, 3-amino propionic acid, ornithine, norleucine, norvaline, hydroxy-proline, sarcosine, citrulline, homocitrulline, cysteic acid, t-butylglycine, t-butylalanine, phenylglycine, cyclohexylalanine, β -alanine, fluoro-amino acids, designer amino acids such as β -methyl amino acids, C α -methyl amino acids, N α -methyl amino acids, and amino acid analogs in general.

[0039] Selection of X is independent in each ELP structural unit (e.g., for each structural unit defined herein having a guest residue X). For example, X may be independently selected for each structural unit as an amino acid having a positively charged side chain, an amino acid having a negatively charged side chain, or an amino acid having a neutral side chain, including in some embodiments, a hydrophobic side chain.

[0040] In still other embodiments, the ELP component(s) may include polymeric or oligomeric repeats of the pentapeptides VPGXG (SEQ ID NO:3), IPGXG (SEQ ID NO:5) or LPGXG (SEQ ID NO:7), or a combination thereof, where X is as defined above.

[0041] In each embodiment, the structural units, or in some cases polymeric or oligomeric repeats, of the ELP sequences may be separated by one or more amino acid residues that do not eliminate the overall effect of the molecule, that is, in imparting certain improvements to the therapeutic component as described. In certain embodiments, such one or more amino acids also do not eliminate or substantially affect the phase transition properties of the ELP component (relative to the deletion of such one or more amino acids).

[0042] In each repeat, X is independently selected. The structure of the resulting ELP components may be described using the notation ELP_k [X_iY_j-n], where k designates a particular ELP repeat unit, the bracketed capital letters are single letter amino acid codes and their corresponding subscripts designate the relative ratio of each guest residue X in the structural units (where applicable), and n describes the total length of the ELP in number of the structural repeats. For example, ELP1 [V₅A₂G₃-10] designates an ELP component containing 10 repeating units of the pentapeptide VPGXG (SEQ ID NO:3), where X is valine, alanine, and glycine at a relative ratio of 5:2:3; ELP1 [K₁V₂F₁-4] designates an ELP component containing 4 repeating units of the pentapeptide VPGXG (SEQ ID NO:3), where X is lysine, valine, and phenylalanine at a relative ratio of 1:2:1; ELP1 [K₁V₇F₁-9] designates a polypeptide containing 9 repeating units of the pentapeptide VPGXG (SEQ ID NO:3), where X is lysine, valine, and phenylalanine at a relative ratio of 1:7:1; ELP1 [V-5] designates a polypeptide containing 5 repeating units of the pentapeptide VPGXG (SEQ ID NO:3), where X is exclusively valine; ELP1 [V-20] designates a polypeptide containing 20 repeating units of the pentapeptide VPGXG (SEQ ID NO:3), where X is exclusively valine; ELP2 [5] designates a polypeptide containing 5 repeating units of the pentapeptide AVGVV (SEQ ID NO:4); ELP3 [V-5] designates a polypeptide containing 5 repeating units of the pentapeptide IPGXG (SEQ ID NO:5), where X is exclusively valine; ELP4 [V-5] designates a polypeptide containing 5 repeating units of the pentapeptide LPGXG (SEQ ID NO:7), where X is exclusively valine. Such ELP components as described in this paragraph may be used in connection with the present invention to increase the therapeutic properties of the therapeutic component.

[0043] Further, the T_t is a function of the hydrophobicity of the guest residue. Thus, by varying the identity of the guest residue(s) and their mole fraction(s), ELPs can be synthesized that exhibit an inverse transition over a 0-100°C range. Thus, the T_t at a given ELP length may be decreased by incorporating a larger fraction of hydrophobic guest residues in the ELP sequence. Examples of suitable hydrophobic guest residues include valine, leucine, isoleucine, phenylalanine, tryptophan and methionine. Tyrosine, which is moderately hydrophobic, may also be used. Conversely, the T_t may be increased by incorporating residues, such as those selected from the group consisting of: glutamic acid, cysteine, lysine, aspartate, alanine, asparagine, serine, threonine, glycine, arginine, and glutamine; preferably selected from alanine, serine, threonine and glutamic acid.

[0044] The ELP component in some embodiments is selected or designed to provide a T_t ranging from about 10 to about 80°C, such as from about 35 to about 60°C, or from about 38 to about 45°C. In some embodiments, the T_t is greater than about 40°C or greater than about 42 °C, or greater than about 45 °C, or greater than about 50 °C. The transition temperature, in some embodiments, is above the body temperature of the subject or patient (e.g., >37°C) thereby remaining soluble *in vivo*, or in other embodiments, the T_t is below the body temperature (e.g., < 37°C) to provide alternative advantages, such as *in vivo* formation of a drug depot for sustained release of the therapeutic agent.

[0045] The T_t of the ELP component can be modified by varying ELP chain length, as the T_t generally increases with decreasing MW. For polypeptides having a molecular weight > 100,000, the hydrophobicity scale developed by Urry et al. (PCT/US96/05186, which is hereby incorporated by reference in its entirety) is preferred for predicting the approximate T_t of a specific ELP sequence. However, in some embodiments, ELP component length can be kept relatively small, while maintaining a target T_t, by incorporating a larger fraction of hydrophobic guest residues (e.g., amino acid residues having hydrophobic side chains) in the ELP sequence. For polypeptides having a molecular weight <100,000, the T_t may be predicted or determined by the following quadratic function: $T_t = M_0 + M_1X + M_2X^2$ where X is the MW of the fusion protein, and $M_0 = 116.21$; $M_1 = -1.7499$; $M_2 = 0.010349$.

[0046] While the T_t of the ELP component, and therefore of the ELP component coupled to a therapeutic component,

is affected by the identity and hydrophobicity of the guest residue, X, additional properties of the molecule may also be affected. Such properties include, but are not limited to solubility, bioavailability, persistence, and half-life of the molecule.

[0047] As described in PCT/US2007/077767 (published as WO 2008/030968), which is hereby incorporated by reference in its entirety, the ELP-coupled therapeutic component can retain the therapeutic component's biological activity.

Additionally, ELPs themselves can exhibit long half-lives. Therefore, ELP components in accordance with the present invention substantially increase (e.g. by greater than 10%, 20%, 30%, 50%, 100%, 200%, 500% or more, in specific embodiments) the half-life of the therapeutic component when conjugated thereto. Such half-life (or in some embodiments persistence or rate of clearance) is determined in comparison to the half-life of the free (unconjugated or unfused) form of the therapeutic component. Furthermore, ELPs may target high blood content organs, when administered *in vivo*, and thus, can partition in the body, to provide a predetermined desired corporeal distribution among various organs or regions of the body, or a desired selectivity or targeting of a therapeutic agent. In sum, the therapeutic agents contemplated by the invention are administered or generated *in vivo* as active compositions having extended half-lives (e.g., circulatory half-life), among other potential benefits described herein.

[0048] The invention thus provides various agents for therapeutic (*in vivo*) application, where the therapeutic component is biologically active. Such therapeutic components include those listed in Table 1 (e.g., full length or functional portions or functional analogs thereof), as well as GLP-1 receptor agonists such as GLP-1 or exendin-4, insulin, or Factor VII/VIIIa, and functional analogs thereof. The structure and activity of such therapeutic components are described in detail below. In some forms of the therapeutic agent, the coupling of the therapeutic component to the ELP component is effected by direct covalent bonding or indirect (through appropriate spacer groups) bonding (as described elsewhere herein). Further, the therapeutic component(s) and the ELP component(s) can be structurally arranged in any suitable manner involving such direct or indirect covalent bonding, relative to one another.

Glucagon-Like Peptide (GLP)-1 Receptor Agonists

[0049] In certain embodiments of the invention, the therapeutic agent comprises an ELP component fused or conjugated to a GLP-1 receptor agonist, such as GLP-1, exendin-4, or functional analogs thereof.

[0050] Human GLP-1 is a 37 amino acid residue peptide originating from proglucagon which is synthesized in the L-cells in the distal ileum, in the pancreas, and in the brain. Processing of proglucagon to give GLP-1 (7-36)amide, GLP-1 (7-37) and GLP-2 occurs mainly in the L-cells. A simple system is used to describe fragments and analogs of this peptide. For example, Gly⁸-GLP-1 (7-37) designates a fragment of GLP-1 formally derived from GLP-1 by deleting the amino acid residues Nos. 1 to 6 and substituting the naturally occurring amino acid residue in position 8 (Ala) by Gly. Similarly, Lys³⁴ (N^ε-tetradecanoyl)-GLP-1(7-37) designates GLP-1 (7-37) wherein the ε-amino group of the Lys residue in position 34 has been tetradecanoylated. Where reference in this text is made to C-terminally extended GLP-1 analogues, the amino acid residue in position 38 is Arg unless otherwise indicated, the optional amino acid residue in position 39 is also Arg unless otherwise indicated and the optional amino acid residue in position 40 is Asp unless otherwise indicated. Also, if a C-terminally extended analogue extends to position 41, 42, 43, 44 or 45, the amino acid sequence of this extension is as in the corresponding sequence in human proglucagon unless otherwise indicated.

[0051] The parent peptide of GLP-1, proglucagon (PG), has several cleavage sites that produce various peptide products dependent on the tissue of origin including glucagon (PG[32-62]) and GLP-1[7-36]NH₂ (PG[72-107]) in the pancreas, and GLP-1[7-37] (PG[78-108]) and GLP-1 [7-36]NH₂ (PG [78-107]) in the L cells of the intestine where GLP-1 [7-36]NH₂ (78-107 PG) is the major product. The GLP-1 component in accordance with the invention may be any biologically active product or derivative of proglucagon, or functional analog thereof, including: GLP-1 (1-35), GLP-1(1-36), GLP-1 (1-36)amide, GLP-1 (1-37), GLP-1 (1-38), GLP-1 (1-39), GLP-1 (1-40), GLP-1 (1-41), GLP-1 (7-35), GLP-1 (7-36), GLP-1 (7-36)amide, GLP-1 (7-37), GLP-1 (7-38), GLP-1 (7-39), GLP-1 (7-40) and GLP-1 (7-41), or a analog of the foregoing. Generally, the GLP-1 component in some embodiments may be expressed as GLP-1 (A-B), where A is an integer from 1 to 7 and B is an integer from 38 to 45, optionally with one or more amino acid substitutions as defined below.

[0052] As an overview, after processing in the intestinal L-cells, GLP-1 is released into the circulation, most notably in response to a meal. The plasma concentration of GLP-1 rises from a fasting level of approximately 15 pmol/L to a peak postprandial level of 40 pmol/L. For a given rise in plasma glucose concentration, the increase in plasma insulin is approximately threefold greater when glucose is administered orally compared with intravenously (Kreymann et al., 1987, Lancet 2(8571): 1300-4). This alimentary enhancement of insulin release, known as the incretin effect, is primarily humoral and GLP-1 is now thought to be the most potent physiological incretin in humans. GLP-1 mediates insulin production via binding to the GLP-1 receptor, known to be expressed in pancreatic β cells. In addition to the insulinotropic effect, GLP-1 suppresses glucagon secretion, delays gastric emptying (Wettergen et al., 1993, Dig Dis Sci 38: 665-73) and may enhance peripheral glucose disposal (D'Alessio et al., 1994, J. Clin Invest 93: 2293-6).

[0053] A combination of actions gives GLP-1 unique therapeutic advantages over other agents currently used to treat non-insulin-dependent diabetes mellitus (NIDDM). First, a single subcutaneous dose of GLP-1 can completely normalize post prandial glucose levels in patients with NIDDM (Gutniak et al., 1994, Diabetes Care 17: 1039-44). This effect may

be mediated both by increased insulin release and by a reduction in glucagon secretion. Second, intravenous infusion of GLP-1 can delay postprandial gastric emptying in patients with NIDDM (Williams et al., 1996, J. Clin Endo Metab 81: 327-32). Third, unlike sulphonylureas, the insulinotropic action of GLP-1 is dependent on plasma glucose concentration (Holz et al., 1993, Nature 361:362-5). Thus, the loss of GLP-1-mediated insulin release at low plasma glucose concentration protects against severe hypoglycemia.

[0054] When given to healthy subjects, GLP-1 potently influences glycemic levels as well as insulin and glucagon concentrations (Orskov, 1992, Diabetologia 35:701-11), effects which are glucose dependent (Weir et al., 1989, Diabetes 38: 338-342). Moreover, it is also effective in patients with diabetes (Gutniak, M., 1992, N. Engl J Med 226: 1316-22), normalizing blood glucose levels in type 2 diabetic subjects and improving glycemic control in type 1 patients (Nauck et al., 1993, Diabetologia 36: 741-4, Creutzfeldt et al., 1996, Diabetes Care 19:580-6).

[0055] GLP-1 is, however, metabolically unstable, having a plasma half-life ($t_{1/2}$) of only 1-2 minutes *in vivo*. Moreover, exogenously administered GLP-1 is also rapidly degraded (Deacon et al., 1995, Diabetes 44: 1126-31). This metabolic instability has limited the therapeutic potential of native GLP-1.

[0056] GLP-1[7-36]NH₂ has the following amino acid sequence: HAEGTFTSDVSSYLEGQAAKEFIAWLVKGR (SEQ ID NO: 13), which may be employed as the GLP-1 component in accordance with the invention. Alternatively, the GLP-1 component may contain glycine (G) at the second position, giving, for example, the sequence HEGTFTSDVSSYLEGQAAKEFIAWLVKGR (SEQ ID NO: 17). The GLP-1 component may be a biologically active fragment of GLP-1, for example, as disclosed in US 2007/0041951, which is hereby incorporated by reference in its entirety. Other fragments and modified sequences of GLP-1 are known in the art (U.S. Pat. No. 5,614,492; U.S. Pat. No. 5,545,618; European Patent Application, Publication No. EP 0658568 A1; WO 93/25579, which are hereby incorporated by reference in their entireties). Such fragments and modified sequences may be used in connection with the present invention, as well as those described below.

[0057] Certain structural and functional analogs of GLP-1 have been isolated from the venom of the Gila monster lizards (*Heloderma suspectum* and *Heloderma horridum*) and have shown clinical utility. Such molecules find use in accordance with the present invention. In particular, exendin-4 is a 39 amino acid residue peptide isolated from the venom of *Heloderma suspectum* and shares approximately 52% homology with human GLP-1. Exendin-4 is a potent GLP-1 receptor agonist that stimulates insulin release, thereby lowering blood glucose levels. Exendin-4 has the following amino acid sequence: HEGTFTSDLSKQMEEEAVRLFELKNGGPSSGAPPPS (SEQ ID NO: 14). A synthetic version of exendin-4 known as exenatide (marketed as Byetta®) has been approved for the treatment of Type-2 Diabetes. Although exenatide is structurally analogous to native GLP-1, it has a longer half-life after injection.

[0058] While exenatide has the ability to lower blood glucose levels on its own, it can also be combined with other medications such as metformin, a thiazolidinedione, a sulphonylureas, and/or insulin to improve glucose control. Exenatide is administered by injection subcutaneously twice per day using a pre-filled pen device. Typical human responses to exenatide include improvements in the initial rapid release of endogenous insulin, an increase in β -cell growth and replication, suppression of pancreatic glucagon release, delayed gastric emptying, and reduced appetite - all of which function to lower blood glucose. Unlike sulphonylureas and meglitinides, exenatide increases insulin synthesis and secretion in the presence of glucose only, thus lessening the risk of hypoglycemia. Despite the therapeutic utility of exenatide, it has certain undesirable traits, including the requirement of twice daily injections, gastrointestinal side effects, and similar to native GLP-1, a relatively short half-life (i.e. approximately 2 hr).

[0059] Various functional analogs of GLP-1 and exendin-4 are known, and which find use in accordance with the invention. These include liraglutide (Novo Nordisk, WO98/008871), R1583/taspoglutide (Roche, WO00/034331), CJC-1131 (ConjuChem, WO00/069911), ZP-10/AVE0010 (Zealand Pharma, Sanofi-Aventis, WO01/004156), and LY548806 (Eli Lilly, WO03/018516).

[0060] Liraglutide, also known as NN2211, is a GLP-1 receptor agonist analog that has been designed for once-daily injection (Harder et al., 2004, Diabetes Care 27: 1915-21). Liraglutide has been tested in patients with type-2 diabetes in a number of studies and has been shown to be effective over a variety of durations. In one study, treatment with liraglutide improved glycemic control, improved β -cell function, and reduced endogenous glucose release in patients with type-2 diabetes after one week of treatment (Degen et al., 2004, Diabetes 53: 1187-94). In a similar study, eight weeks of 0.6-mg liraglutide therapy significantly improved glycemic control without increasing weight in subjects with type 2 diabetes compared with those on placebo (Harder et al., 2004, Diabetes Care 27: 1915-21).

[0061] Thus, in certain embodiments, the GLP-1 receptor agonist in accordance with the invention is as described in WO98/008871, which is hereby incorporated by reference in its entirety. The GLP-1 receptor agonist may have at least one lipophilic substituent, in addition to one, two, or more amino acid substitutions with respect to native GLP-1. For example, the lipophilic substituent may be an acyl group selected from CH₃(CH₂)_nCO-, wherein n is an integer from 4 to 38, such as an integer from 4 to 24. The lipophilic substituent may be an acyl group of a straight-chain or branched alkyl or fatty acid (for example, as described in WO98/008871, which description is hereby incorporated by reference).

[0062] In certain embodiments, the GLP-1 component is Arg²⁶-GLP-1 (7-37), Arg³⁴-GLP-1 (7-37), Lys³⁶-GLP-1 (7-37), Arg^{26,34}Lys³⁶-GLP-I (7-37), Arg^{26,34}Lys³⁸-GLP-I (7-38), Arg^{28,34} Lys³⁹-GLP-1 (7-39), Arg^{26,34}Lys⁴⁰-GLP-1(7-40),

Arg²⁶Lys³⁶-GLP-1(7-37), Arg³⁴Lys³⁶-GLP-1(7-37), Arg²⁶Lys³⁹-GLP-1(7-39), Arg³⁴Lys⁴⁰-GLP-1(7-40), Arg^{26,34}Lys^{36,39}-GLP-1(7-39), Arg^{26,34}Lys^{36,40}-GLP-1(7-40), Gly⁸Arg²⁶-GLP-1(7-37); Gly⁸Arg³⁴-GLP-1(7-37); Gly⁸Lys³⁸-GLP-1(7-37); Gly⁸Arg^{26,34}Lys³⁶-GLP-1(7-37), Gly⁸Arg^{26,34}Lys³⁹-GLP-1(7-39), Gly⁸Arg^{26,34}Lys⁴⁰-GLP-1(7-40), Gly⁶Arg²⁶Lys³⁶-GLP-1(7-37), Gly⁸Arg³⁴Lys³⁶-GLP-1(7-37), Gly⁸Arg²⁶Lys³⁹-GLP-1(7-39); Gly⁸Arg³⁴Lys⁴⁰-GLP-1(7-40), Gly⁸Arg^{28,34}Lys^{36,39}-GLP-1(7-39) and Gly⁸Arg^{26,34}Lys^{35,40}-GLP-1(7-40), each optionally having a lipophilic substituent. For example, the GLP-1 receptor agonist may have the sequence/structure Arg³⁴Lys²⁶-(N-ε-(γ-Glu(N-α-hexadecanoyl)))-GLP-1(7-37).

[0063] Taspoglutide, also known as R1583 or BIM 51077, is a GLP-1 receptor agonist that has been shown to improve glycemic control and lower body weight in subjects with type 2 diabetes mellitus treated with metformin (Abstract No. A-1604, June 7, 2008, 68th American Diabetes Association Meeting, San Francisco, CA).

[0064] Thus, in certain embodiments, the GLP-1 receptor agonist is as described in WO00/034331, which is hereby incorporated by reference in its entirety. In certain exemplary embodiments, the GLP-1 receptor agonist has the sequence [Aib^{8,35}]hGLP-1(7-36)NH₂ (e.g. taspoglutide), wherein Aib is alpha-aminoisobutyric acid.

[0065] CJC-1131 is a GLP-1 analog that consists of a DPP-IV-resistant form of GLP-1 joined to a reactive chemical linker group that allows GLP-1 to form a covalent and irreversible bond with serum albumin following subcutaneous injection (Kim et al., 2003, Diabetes 52: 751-9). In a 12-week, randomized, double-blind, placebo-controlled multicenter study, CJC-1131 and metformin treatment was effective in reducing fasting blood glucose levels in type 2 diabetes patients (Ratner et al., Abstract No. 10-OR, June 10-14th, 2005, 65th American Diabetes Association Meeting, San Francisco, CA).

[0066] Thus, in certain embodiments, the GLP-1 receptor agonist is as described in WO00/069911, which is hereby incorporated by reference in its entirety. In some embodiments, the GLP-1 receptor agonist is modified with a reactive group which reacts with amino groups, hydroxyl groups or thiol groups on blood components to form a stable covalent bond. In certain embodiments, the GLP-1 receptor agonist is modified with a reactive group selected from the group consisting of succinimidyl and maleimido groups. In certain exemplary embodiments, the GLP-1 receptor agonist has the sequence/structure: D-Ala⁸Lys³⁷-(2-(2-(2-maleimidopropionamido(ethoxy)ethoxy)acetamide))-GLP-1(7-37) (e.g. CJC-1131).

[0067] AVE0010, also known as ZP-10, is a GLP-1 receptor agonist that may be employed in connection with the invention. In a recent double-blind study, patients treated with once daily dosing of AVE0010 demonstrated significant reductions in HbA1c levels (Ratner et al., Abstract No. 433-P, 68th American Diabetes Association Meeting, San Francisco, CA.). At the conclusion of the study, the percentages of patients with HbA1c <7% ranged from 47-69% for once daily dosing compared to 32% for placebo. In addition, AVE0010 treated patients showed dose-dependent reductions in weight and post-prandial plasma glucose.

[0068] Thus, in certain embodiments, the GLP-1 receptor agonist is as described in WO01/004156, which is hereby incorporated by reference in its entirety. For example, the GLP-1 receptor agonist may have the sequence: HGEFTFTSDLSKQMEEEAVRLFIEWLKNGGPSSGAPPSKKKKKK-NH₂ (SEQ ID NO: 18) (e.g. AVE0010).

[0069] LY548806 is a GLP-1 derivative designed to be resistant to proteolysis by dipeptidase-peptidyl IV (DPP-IV) (Jackson et al., Abstract No. 562, June 10-14th, 2005, 65th American Diabetes Association Meeting, San Francisco, CA). In an animal model of hyperglycemia, LY548806 has been shown to produce a significant lowering of blood glucose levels during the hyperglycemic phase (Saha et al., 2006, J. Pharm. Exp. Ther. 316: 1159-64). Moreover, LY548806 was shown to produce a significant increase in insulin levels consistent with its known mechanism of action, namely stimulation of insulin release in the presence of hyperglycemia.

[0070] Thus, in certain embodiments, the GLP-1 receptor agonist is as described in WO03/018516, which is hereby incorporated by reference in its entirety. In some embodiments, the therapeutic agents of the present invention comprise GLP-1 analogs wherein the backbone for such analogs or fragments contains an amino acid other than alanine at position 8 (position 8 analogs). The backbone may also include L-histidine, D-histidine, or modified forms of histidine such as desamino-histidine, 2-amino-histidine, β-hydroxy-histidine, homohistidine, α-fluoromethyl-histidine, or α-methyl-histidine at position 7. In some embodiments, these position 8 analogs may contain one or more additional changes at positions 12, 16, 18, 19, 20, 22, 25, 27, 30, 33, and 37 compared to the corresponding amino acid of native GLP-1. In other embodiments, these position 8 analogs may contain one or more additional changes at positions 16, 18, 22, 25 and 33 compared to the corresponding amino acid of native GLP-1. In certain exemplary embodiments, the GLP-1 receptor agonist has the sequence: HVEGFTFTSDVSSYLEEQAAKEFIWLKGRG-OH (SEQ ID NO: 19) (e.g. LY548806).

[0071] Thus, the present invention provides therapeutic agents comprising an elastin-like peptide (ELP) and a GLP-1 receptor agonist. For example, in certain embodiments, the GLP-1 receptor agonist is GLP-1 (SEQ ID NO:13, 17, or 59) or a functional analog thereof. In other embodiments, the GLP-1 receptor agonist is exendin-4 (SEQ ID NO:14) or a functional analog thereof. Such functional analogs of GLP-1 or exendin-4 include functional fragments truncated at the C-terminus by from 1 to 10 amino acids, including by 1, 2, 3, or up to about 5 amino acids (with respect to SEQ ID NOS: 13, 14, 17, or 59). Such functional analogs may contain from 1 to 10 amino acid insertions, deletions, and/or substitutions (collectively) with respect to the native sequence (e.g., SEQ ID NOS 13, 14, and 59), and in each case

retaining the activity of the peptide. For example, the functional analog of GLP-1 or exendin-4 may have from 1 to about 3, 4, or 5 insertions, deletions and/or substitutions (collectively) with respect to SEQ ID NOS: 13, 59 and 14, and in each case retaining the activity of the peptide. Such activity may be confirmed or assayed using any available assay, including those described herein. In these or other embodiments, the GLP-1 receptor agonist component has at least about 50%, 75%, 80%, 85%, 90%, or 95% identity with the native sequence (SEQ ID NOS: 13, 59, and 14). The determination of sequence identity between two sequences (e.g., between a native sequence and a functional analog) can be accomplished using any alignment tool, including Tatusova et al., Blast 2 sequences - a new tool for comparing protein and nucleotide sequences, FEMS Microbiol Lett. 174:247-250 (1999). Such functional analogs may further comprise additional chemical modifications, such as those described in this section and/or others known in the art.

[0072] In certain embodiments, the GLP1-ELP fusion has a sequence exemplified herein as SEQ ID NOS: 54 and 56. When processed, the mature form of such fusion protein will begin with the His⁷ of GLP.

[0073] In another aspect, the present invention provides methods for the treatment or prevention of type 2 diabetes, impaired glucose tolerance, type 1 diabetes, hyperglycemia, obesity, binge eating, bulimia, hypertension, syndrome X, dyslipidemia, cognitive disorders, atherosclerosis, non-fatty liver disease, myocardial infarction, coronary heart disease and other cardiovascular disorders. The method comprises administering the therapeutic agent comprising the elastin-like peptide (ELP) and the GLP-1 receptor agonist (as described above) to a patient in need of such treatment. In these or other embodiments, the present invention provides methods for decreasing food intake, decreasing β -cell apoptosis, increasing β -cell function and β -cell mass, and/or for restoring glucose sensitivity to β -cells. Generally, the patient may be a human or non-human animal patient (e.g., dog, cat, cow, or horse). Preferably, the patient is human.

[0074] The treatment with a ELP/GLP-1 receptor agonist compound according to the present invention may also be combined with one or more pharmacologically active substances, e.g. selected from antidiabetic agents, antiobesity agents, appetite regulating agents, antihypertensive agents, agents for the treatment and/or prevention of complications resulting from or associated with diabetes and agents for the treatment and/or prevention of complications and disorders resulting from or associated with obesity. In the present context, the expression "antidiabetic agent" includes compounds for the treatment and/or prophylaxis of insulin resistance and diseases wherein insulin resistance is the pathophysiological mechanism.

[0075] The ability of a GLP-1 or exendin-4 analog, or an GLP-1 receptor agonist/ELP compound, to bind the GLP-1 receptor may be determined by standard methods, for example, by receptor-binding activity screening procedures which involve providing appropriate cells that express the GLP-1 receptor on their surface, for example, insulinoma cell lines such as RINmSF cells or INS-1 cells. In addition to measuring specific binding of tracer to membrane using radioimmunoassay methods, cAMP activity or glucose dependent insulin production can also be measured. In one method, a polynucleotide encoding the GLP-1 receptor is employed to transfect cells to thereby express the GLP-1 receptor protein. Thus, these methods may be employed for testing or confirming whether a suspected GLP receptor agonist is active. An exemplary assay is described in greater detail herein.

[0076] In addition, known methods can be used to measure or predict the level biological activity of a GLP-1 receptor agonist or GLP-1 receptor agonist/ELP *in vivo* (See e.g. Siegel, et al., 1999, Regul Pept 79(2-3): 93-102). In particular, GLP-1 receptor agonist or GLP-1 receptor agonist/ELP compounds can be assessed for their ability to induce the production of insulin *in vivo* using a variety of known assays for measuring GLP-1 activity. For example, an ELP/GLP-1 receptor agonist compound can be introduced into a cell, such as an immortalized β -cell, and the resulting cell can be contacted with glucose. If the co-produces insulin in response to the glucose, then the modified GLP-1 is generally considered biologically active *in vivo* (Fehmann et al., 1992, Endocrinology 130: 159-166). An exemplary assay is described in greater detail herein.

[0077] The ability of an GLP-1 receptor agonist/ELP compound to enhance β -cell proliferation, inhibit β -cell apoptosis, and regulate islet growth may also be measured using known assays. Pancreatic β -cell proliferation may be assessed by ³H-thymidine or BrdU incorporation assays (See e.g. Buteau et al., 2003, Diabetes 52: 124-32), wherein pancrea β -cells such as INS(832/13) cells are contacted with an ELP/ GLP-1 receptor agon compound and analyzed for increases in ³H-thymidine or BrdU incorporation. The antiapoptotic activity of an ELP/GLP-1 receptor agonist compound can be measured cultured insulin-secreting cells and/or in animal models where diabetes occurs as consequence of an excessive rate of beta-cell apoptosis (See e.g. Bulotta et al., 2004, C Biochem Biophys 40(3 suppl): 65-78).

[0078] In addition to GLP-1, other peptides of this family, such as those derived from processing of the pro-glucagon gene, such as GLP-2, GIP, and oxyntomodulin, could be conjugated or fused to the ELP component (as described herein) to enhance the therapeutic potential.

Insulin

[0079] In other embodiments, the present invention provides a therapeutic agent comprising an ELP component coupled to insulin (e.g., via fusion or conjugation). Insulin injections, e.g. of human insulin, can be used to treat diabetes. The insulin-making cells in the body are called β -cells, and they are found in the pancreas gland. These cells cluster together

to form the "islets of Langerhans", named for the German medical student who described them.

[0080] The synthesis of insulin begins at the translation of the insulin gene, which resides on chromosome 11. During translation, two introns are spliced out of the mRNA product, which encodes a protein of 110 amino acids in length. This primary translation product is called preproinsulin and is inactive. It contains a signal peptide of 24 amino acids in length, which is required for the protein to cross the cell membrane.

[0081] Once the preproinsulin reaches the endoplasmic reticulum, a protease cleaves off the signal peptide to create proinsulin. Proinsulin consists of three domains: an amino-terminal B chain, a carboxyl-terminal A chain, and a connecting peptide in the middle known as the C-peptide. Insulin is composed of two chains of amino acids named chain A (21 amino acids - GIVEQCCASVCSLYQLENYCN) (SEQ ID NO: 15) and chain B (30 amino acids FVNQHLCGSHLVEA-LYLVCGERGFFYTPKA) (SEQ ID NO: 16) that are linked together by two disulfide bridges. There is a 3rd disulfide bridge within the A chain that links the 6th and 11th residues of the A chain together. In most species, the length and amino acid compositions of chains A and B are similar, and the positions of the three disulfide bonds are highly conserved. For this reason, pig insulin can replace deficient human insulin levels in diabetes patients. Today, porcine insulin has largely been replaced by the mass production of human proinsulin by bacteria (recombinant insulin).

[0082] Insulin molecules have a tendency to form dimers in solution, and in the presence of zinc ions, insulin dimers associate into hexamers. Whereas monomers of insulin readily diffuse through the blood and have a rapid effect, hexamers diffuse slowly and have a delayed onset of action. In the design of recombinant insulin, the structure of insulin can be modified in a way that reduces the tendency of the insulin molecule to form dimers and hexamers but that does not interrupt binding to the insulin receptor. In this way, a range of preparations are made, varying from short acting to long acting.

[0083] Within the endoplasmic reticulum, proinsulin is exposed to several specific peptidases that remove the C-peptide and generate the mature and active form of insulin. In the Golgi apparatus, insulin and free C-peptide are packaged into secretory granules, which accumulate in the cytoplasm of the β -cells. Exocytosis of the granules is triggered by the entry of glucose into the beta cells. The secretion of insulin has a broad impact on metabolism.

[0084] There are two phases of insulin release in response to a rise in glucose. The first is an immediate release of insulin. This is attributable to the release of preformed insulin, which is stored in secretory granules. After a short delay, there is a second, more prolonged release of newly synthesized insulin.

[0085] Once released, insulin is active for only a brief time before it is degraded by enzymes. Insulinase found in the liver and kidneys breaks down insulin circulating in the plasma, and as a result, insulin has a half-life of only about 6 minutes. This short duration of action results in rapid changes in the circulating levels of insulin.

[0086] Insulin analogs have been developed with improved therapeutic properties (Owens et al., 2001, Lancet 358: 739-46; Vajo et al., 2001, Endocr Rev 22: 706-17), and such analogs may be employed in connection with the present invention. Various strategies, including elongation of the COOH-terminal end of the insulin B-chain and engineering of fatty acid-acylated insulins with substantial affinity for albumin are used to generate longer-acting insulin analogs. However, *in vivo* treatments with available longer-acting insulin compounds still result in a high frequency of hypo- and hyperglycemic excursions and modest reduction in HbA_{1c}. Accordingly, development of a truly long-acting and stable human insulin analog still remains an important task.

[0087] Functional analogs of insulin that may be employed in accordance with the invention include rapid acting analogs such as lispro, aspart and glulisine, which are absorbed rapidly (< 30 minutes) after subcutaneous injection, peak at one hour, and have a relatively short duration of action (3 to 4 hours). In addition, two long acting insulin analogs have been developed: glargine and detemir, and which may be employed in connection with the invention. The long acting insulin analogs have an onset of action of approximately two hours and reach a plateau of biological action at 4 to 6 hours, and may last up to 24 hours.

[0088] Thus, in one embodiment, the insulin component may contain the A and/or B chain of lispro (also known as Humalog, Eli Lilly). Insulin lispro differs from human insulin by the substitution of proline with lysine at position 28 and the substitution of lysine with proline at position 29 of the insulin B chain. Although these modifications do not alter receptor binding, they help to block the formation of insulin dimers and hexamers, allowing for larger amounts of active monomeric insulin to be available for postprandial injections.

[0089] In another embodiment, the insulin may contain an A and/or B chain of aspart (also known as Novolog, Novo Nordisk). Insulin aspart is designed with the single replacement of the amino acid proline by aspartic acid at position 28 of the human insulin B chain. This modification helps block the formation of insulin hexamers, creating a faster acting insulin.

[0090] In yet another embodiment, the insulin may contain an A and/or B chain of glulisine (also known as Apidra, Sanofi-Aventis). Insulin glulisine is a short acting analog created by substitution of asparagine at position 3 by lysine and lysine at position 29 by glutamine of human insulin B chain. Insulin glulisine has more rapid onset of action and shorter duration of action compared to regular human insulin.

[0091] In another embodiment, the insulin may contain an A and/or B chain of glargine (also known as Lantus, Sanofi-Aventis). Insulin glargine differs from human insulin in that the amino acid asparagine at position 21 of the A chain is

replaced by glycine and two arginines are added to the C-terminus of the B-chain. Compared with bedtime neutral protamine Hagedorn (NPH) insulin (an intermediate acting insulin), insulin glargine is associated with less nocturnal hypoglycemia in patients with type 2 diabetes.

[0092] In yet another embodiment, the insulin may contain an A and/or B chain from detemir (also known as Levemir, Novo Nordisk). Insulin detemir is a soluble (at neutral pH) long-acting insulin analog, in which the amino acid threonine at B30 is removed and a 14-carbon, myristoyl fatty acid is acetylated to the epsilon-amino group of LysB29. After subcutaneous injection, detemir dissociates, thereby exposing the free fatty acid which enables reversible binding to albumin molecules. So at steady state, the concentration of free unbound insulin is greatly reduced resulting in stable plasma glucose levels.

[0093] In some embodiments, the insulin may be a single-chain insulin analog (SIA) (e.g. as described in 6,630,438 and WO08/019368, which are hereby incorporated by reference in their entirety). Single-chain insulin analogs encompass a group of structurally-related proteins wherein the A and B chains are covalently linked by a polypeptide linker. The polypeptide linker connects the C-terminus of the B chain to the N-terminus of the A chain. The linker may be of any length so long as the linker provides the structural conformation necessary for the SIA to have a glucose uptake and insulin receptor binding effect. In some embodiments, the linker is about 5-18 amino acids in length. In other embodiments, the linker is about 9-15 amino acids in length. In certain embodiments, the linker is about 12 amino acids long. In certain exemplary embodiments, the linker has the sequence KDDNPNLRLVR (SEQ ID NO.: 20) or GAGSSRRAPQT (SEQ ID NO.: 21). However, it should be understood that many variations of this sequence are possible such as in the length (both addition and deletion) and substitutions of amino acids without substantially compromising the effectiveness of the produced SIA in glucose uptake and insulin receptor binding activities. For example, several different amino acid residues may be added or removed from either end without substantially decreasing the activity of the produced SIA.

[0094] An exemplary single-chain insulin analog currently in clinical development is abulin (Duttaroy et al., 2005, Diabetes 54: 251-8). Abulin can be produced in yeast or in mammalian cells. It consists of the B and A chain of human insulin (100% identity to native human insulin) linked together by a dodecapeptide linker and fused to the NH₂ terminals of the native human serum albumin. For expression and purification of abulin, Duttaroy et al. constructed a synthetic gene construct encoding a single-chain insulin containing the B- and A- chain of mature human insulin linked together by a dodecapeptide linker using four overlapping primers and PCR amplification. The resulting PCR product was ligated in-frame between the signal peptide of human serum albumin (HSA) and the NH₂ terminus of mature HSA, contained within a pSAC35 vector for expression in yeast. In accordance with the present invention, the HSA component of abulin may be replaced with an ELP component as described herein.

[0095] Thus, in one aspect, the present invention provides therapeutic agents comprising an elastin-like peptide (ELP) and an insulin or functional analog thereof. For example, in certain embodiments, the insulin is a mammalian insulin, such as human insulin or porcine insulin. In accordance with the invention, the ELP component may be coupled (e.g., via recombinant fusion or chemical conjugation) to the insulin A chain, or B chain, or both. The insulin may comprise each of chains A, B, and C (SEQ ID NOS: 51 and 52), or may contain a processed form, containing only chains A and B. In some embodiments, chains A and B are connected by a short linking peptide, to create a single chain insulin. The insulin may be a functional analog of human insulin, including functional fragments truncated at the N-terminus and/or C-terminus (of either or both of chains A and B) by from 1 to 10 amino acids, including by 1, 2, 3, or about 5 amino acids. Functional analogs may contain from 1 to 10 amino acid insertions, deletions, and/or substitutions (collectively) with respect to the native sequence (e.g., SEQ ID NOS 15 and 16), and in each case retaining the activity of the peptide. For example, functional analogs may have 1, 2, 3, 4, or 5 amino acid insertions, deletions, and/or substitutions (collectively) with respect to the native sequence (which may contain chains A and B, or chains A, B, and C). Such activity may be confirmed or assayed using any available assay, including those described herein. In these or other embodiments, the insulin component has at least about 75%, 80%, 85%, 90%, 95%, or 98% identity with each of the native sequences for chains A and B (SEQ ID NOS: 15 and 16). The determination of sequence identity between two sequences (e.g., between a native sequence and a functional analog) can be accomplished using any alignment tool, including Tatusova et al., Blast 2 sequences - a new tool for comparing protein and nucleotide sequences, FEMS Microbiol Lett. 174:247-250 (1999). The insulin component may contain additional chemical modifications known in the art.

[0096] In another aspect, the present invention provides methods for the treatment or prevention of diabetes, including type I and II diabetes. The method comprises administering an effective amount of the therapeutic agent comprising an elastin-like peptide

[0097] (ELP) component and an insulin (or functional analog thereof) component to a patient in need thereof. Generally, the patient may be a human or non-human animal (e.g., dog, cat, cow, or horse) patient. Preferably, the patient is human.

[0098] To characterize the *in vitro* binding properties of an insulin analog or an ELP-containing insulin analog, competition binding assays may be performed in various cell lines that express the insulin receptor (Jehle et al., 1996, Diabetologia 39: 421-432). For example, competition binding assays using CHO cells overexpressing the human insulin receptor may be employed. Insulin can also bind to the IGF-1 receptor with a lower affinity than the insulin receptor. To determine the binding affinity of an ELP-containing insulin analog, a competition binding assay can be performed using

¹²⁵I-labeled IGF-1 in L6 cells.

[0099] The activities of insulin include stimulation of peripheral glucose disposal and inhibition of hepatic glucose production. The ability of an ELP-containing insulin analog to mediate these biological activities can be assayed *in vitro* using known methodologies. For example, the effect of an ELP-containing analog on glucose uptake in 3T3-L1 adipocytes can be measured and compared with that of insulin. Pretreatment of the cells with a biologically active analog will generally produce a dose-dependent increase in 2-deoxyglucose uptake. The ability of an ELP-containing insulin analog to regulate glucose production may be measured in any number of cells types, for example, H4Ile hepatoma cells. In this assay, pretreatment with a biologically active analog will generally result in a dose-dependent inhibition of the amount of glucose released.

Factor VII(VIIa)

[0100] In certain embodiments, the invention provides therapeutic agents comprising an ELP component coupled (e.g., via fusion or conjugation) to a Factor VII/VIIa. Coagulation is the biological process of blood clot formation involving many different serine proteases as well as their essential cofactors and inhibitors. It is initiated by exposure of Factor VII (FVII) and Factor VIIa (FVIIa) to its membrane bound cofactor, tissue factor (TF), resulting in production of Factor Xa (FXa) and more FVIIa. The process is propagated upon production of Factor IXa (FIXa) and additional FXa that, upon binding with their respective cofactors FVIIIa and FVa, form platelet bound complexes, ultimately resulting in the formation of thrombin and a fibrin clot. Thrombin also serves to further amplify coagulation by activation of cofactors such as FV and FVII and zymogens such as Factor XI. Moreover, thrombin activates platelets leading to platelet aggregation, which is necessary for the formation of a hemostatic plug.

[0101] Factor VII circulates in the blood in a zymogen form, and is converted to its active form, Factor VIIa, by either factor IXa, factor Xa, factor XIIa, or thrombin by minor proteolysis. Factor VIIa is a two-chain, 50 kilodalton (kDa) plasma serine protease. The active form of the enzyme comprises a heavy chain (254 amino acid residues) containing a catalytic domain and a light chain (152 residues) containing 2 epidermal growth factor (EGF)-like domains. The mature factor VII/VIIa that circulates in plasma is composed of 406 amino acid residues (SEQ ID NO: 33). The light and heavy chains are held together by a disulfide bond.

[0102] As noted above, Factor VIIa is generated by proteolysis of a single peptide bond from its single chain zymogen, Factor VII, which is present at approximately 0.5 μ g/ml in plasma. The conversion of zymogen Factor VII into the activated two-chain molecule occurs by cleavage of an internal peptide bond. In human Factor VII, the cleavage site is at Arg 152-Ile153 (Hagen et al., 1986, PNAS USA 83: 2412-6).

[0103] "Factor, VII/VIIa" as used in this application means a product consisting of either the unactivated form (factor VII) or the activated form (factor VIIa) or mixtures thereof. "Factor VII/VIIa" within the above definition includes proteins that have an amino acid sequence of native human factor VII/VIIa. It also includes proteins with a slightly modified amino acid sequence, for instance, a modified N-terminal end including N-terminal amino acid deletions or additions so long as those proteins substantially retain the activity of factor VIIa. "Factor VII" within the above definition also includes natural allelic variations that may exist and occur from one individual to another. Also, degree and location of glycosylation or other post-translation modifications may vary depending on the chosen host cells and the nature of the host cellular environment.

[0104] In the presence of calcium ions, Factor VIIa binds with high affinity to TF. TF is a 263 amino acid residue glycoprotein composed of a 219 residue extracellular domain, a single transmembrane domain, and a short cytoplasmic domain (Morrissey et al., 1987, Cell 50: 129-35). The TF extracellular domain is composed of two fibronectin type III domains of about 105 amino acids each. The binding of FVIIa is mediated entirely by the TF extracellular domain (Muller et al., 1994, Biochem. 33:10864-70). Residues in the area of amino acids 16-26 and 129-147 contribute to the binding of FVIIa as well as the coagulant function of the molecule. Residues Lys20, Trp45, Asp58, Tyr94, and Phe140 make a large contribution (1 kcal/mol) to the free energy (ΔG) of binding to FVIIa.

[0105] TF is expressed constitutively on cells separated from plasma by the vascular endothelium. Its expression on endothelial cells and monocytes is induced by exposure to inflammatory cytokines or bacterial lipopolysaccharides (Drake et al., 1989, J. Cell Biol. 109: 389). Upon tissue injury, the exposed extracellular domain of TF forms a high affinity, calcium dependent complex with FVII. Once bound to TF, FVII can be activated by peptide bond cleavage to yield serine protease FVIIa. The enzyme that catalyzes this step *in vivo* has not been elucidated, but *in vitro* FXa, thrombin, TF:FVIIa and FIXa can catalyze this cleavage. FVIIa has only weak activity upon its physiological substrates FX and FIX whereas the TF:FVIIa complex rapidly activates FX and FIX.

[0106] The TF:FVIIa complex constitutes the primary initiator of the extrinsic pathway of blood coagulation. The complex initiates the extrinsic pathway by activation of FX to Factor Xa (FXa), FIX to Factor IXa (FIXa), and additional FVII to FVIIa. The action of TF:FVIIa leads ultimately to the conversion of prothrombin to thrombin, which carries out many biological functions. Among the most important activities of thrombin is the conversion of fibrinogen to fibrin, which polymerizes to form a clot. The TF:FVIIa complex also participates as a secondary factor in extending the physiological

effects of the contact activation system.

[0107] The initiation and subsequent regulation of coagulation is complex, since maintenance of hemostasis is crucial for survival. There is an exquisite balance between hemostasis (normal clot formation and dissolution) and thrombosis (pathogenic clot formation). Serious clinical conditions involving aberrations in coagulation include deep vein thrombosis, myocardial infarction, pulmonary embolism, stroke and disseminated intravascular coagulation (in sepsis). There are also many bleeding coagulopathies where there is insufficient clot formation. These include hemophilia A (FVIII deficiency) or hemophilia B (FIX deficiency), where procoagulant therapy is required. The challenge in this therapeutic area is to operate in the narrow window between too much and too little coagulation.

[0108] The use of exogenous FVIIa as a therapeutic agent has been shown to induce hemostasis in patients with hemophilia A and B (Hedner, 2001, *Seminars Hematol.* 38 (suppl. 12): 43-7; Hedner, 2004, *Seminars Hematol.* 41 (suppl. 1): 35-9). It also has been used to treat bleeding in patients with liver disease, anticoagulation-induced bleeding, surgery, thrombocytopenia, thrombasthenia, Bernard-Soulier syndrome, von Willebrand disease, and other bleeding disorders (See e.g. Roberts et al., 2004, *Blood* 104: 3858-64).

[0109] Commercial preparations of human recombinant FVIIa are sold as NovoSeven™. NovoSeven™ is indicated for the treatment of bleeding episodes in hemophilia A or B patients and is the only recombinant FVIIa effective for bleeding episodes currently available. A circulating recombinant FVIIa half-life of 2.3 hours was reported in "Summary Basis for Approval for NovoSeven™" FDA reference number 96-0597. Moreover, the half-life of recombinant FVIIa is shorter in pediatric patients (~ 1.3 hours), suggesting that higher doses of recombinant FVIIa may be required in this population (Roberts et al., 2004, *Blood* 104: 3858-64). Accordingly, relatively high doses and frequent administration are necessary to reach and sustain the desired therapeutic or prophylactic effect. As a consequence, adequate dose regulation is difficult to obtain and the need of frequent intravenous administrations imposes restrictions on the patient's way of living.

[0110] A molecule with a longer circulation half-life would decrease the number of necessary administrations. Given the frequent injections associated with currently available FVIIa therapy and the potential for obtaining more optimal therapeutic FVIIa levels with concomitant enhanced therapeutic effect, there is a clear need for improved FVII or FVIIa-like molecules with a longer half-life *in vivo*.

[0111] Recombinant human coagulation factor VIIa (rFVIIa, NovoSeven; Novo Nordisk A/S, Copenhagen, Denmark) has proven to be efficacious for the treatment of bleeding episodes in hemophilia patients with inhibitors. A small fraction of patients may be refractory to rFVIIa treatment and could potentially benefit from genetically modified FVIIa molecules with increased potencies. To this end, FVIIa analogs with increased intrinsic activity have been investigated that exhibit superior hemostatic profiles *in vitro* (see e.g. WO02/077218 or WO05/074975, which are hereby incorporated by reference in their entirety, and Tranholm et al., 2003, *Blood* 102(10): 3615-20, which is also incorporated by reference). These analogs may also be used as more efficacious hemostatic agents in other indications where efficacy of rFVIIa has been observed, including in thrombocytopenia and trauma.

[0112] Thus, in some embodiments, the Factor VIIa analog that may be used in accordance with the invention is as described in WO02/077218 or WO05/074975. For example, the FVIIa analog may have a glutamine substituted for methionine at position 298 (i.e. M298Q-FVIIa). In certain exemplary embodiments, the FVIIa analog contains two additional mutations, valine at position 158 replaced by aspartic acid and glutamic acid at position 296 replaced by valine (i.e. V158D/E296V/M298Q-FVIIa). Additionally or alternatively, the Factor VIIa analog may have an alanine residue substitution for lysine at position 337 (i.e. V158D/E296V/M298Q/K337A-FVIIa). In still other embodiments, the Factor VIIa analog has a substitution or insertion selected from Q250C; P406C; and 407C, wherein a cysteine has also been introduced in the C-terminal sequence (see, e.g. US 7,235,638, which is hereby incorporated by reference in its entirety). The Factor VIIa analog may further comprise a substitution or insertion at one or more of positions 247, 260, 393, 396, and/or 405.

[0113] In these or other embodiments, the Factor VIIa analog comprises a substitution relative to the sequence of native Factor VIIa selected from: (a) a substitution of Lys157 with an amino acid selected from the group consisting of Gly, Val, Ser, Thr, Asp, and Glu; (b) a substitution of Lys337 with an amino acid selected from the group consisting of Ala, Gly, Val, Ser, Thr, Gln, Asp, and Glu; (c) a substitution of Asp334 with any amino acid other than Ala or Asn; and (d) a substitution of Ser336 with any amino acid other than Ala or Cys (see e.g. US 7,176,288, which is hereby incorporated by reference in its entirety). Additionally or alternatively, the Factor VIIa analog comprises a substitution of the Leu at position 305 of Factor VII with an amino acid residue selected from the group consisting of Val, Ile, Met, Phe, Trp, Pro, Gly, Ser, Thr, Cys, Tyr, Asn, Glu, Lys, Arg, His, Asp and Gln (see e.g. US 6,905,683, which is hereby incorporated by reference in its entirety).

[0114] Thus, in one aspect, the present invention provides therapeutic agents comprising an elastin-like peptide (ELP) and a Factor VII/VIIa, or functional analog thereof. For example, in certain embodiments, the Factor VII/VIIa is human Factor VII/VIIa (e.g., SEQ ID NO: 33). The Factor VII/VIIa may be a functional analog of human Factor VII/VIIa, including functional fragments truncated at the N-terminus and/or C-terminus by from 1 to 10 amino acids, including by 1, 2, 3, or about 5 amino acids. Functional analogs may contain from 1 to 10 amino acid insertions, deletions, and/or substitutions

(collectively) with respect to the native sequence (e.g., SEQ ID NO: 33), and in each case retaining the activity of the peptide. For example, such analogs may have from 1 to about 5 amino acid insertions, deletions, and/or substitutions (collectively) with respect to the native full length sequence, or with respect to one or both of the heavy and light chains. Such activity may be confirmed or assayed using any available assay, including those described herein. In these or

5 other embodiments, the Factor VII/VIIa component has at least about 75%, 80%, 85%, 90%, 95%, or 98% identity with the native sequence (SEQ ID NO:33). The determination of sequence identity between two sequences (e.g., between a native sequence and a functional analog) can be accomplished using any alignment tool, including Tatusova et al., Blast2 sequences - a new tool for comparing protein and nucleotide sequences, FEMS Microbiol Lett. 174:247-250(1999).

10 **[0115]** In exemplary embodiments, the FactorVII-ELP fusion has the amino acid sequence of SEQ ID NO:58. SEQ ID NO:58 further comprises a TEV protease cleavage site between the FactorVII and ELP sequences, which may be beneficial for removing the ELP sequence post expression where desired. However, in accordance with the invention, the tev sequence may be entirely removed, or replaced with another linking sequence as disclosed herein.

15 **[0116]** In another aspect, the present invention provides methods for the treatment or prevention of bleeding-related disorders. The method comprises administering an effective amount of the therapeutic agent comprising an elastin-like peptide (ELP) and a Factor VII/VIIa or functional analog thereof to a patient in need. In certain embodiments, the bleeding-related disorder is one or more of hemophilia (A or B), post-surgical bleeding, anticoagulation-induced bleeding, thrombocytopenia, Factor VII deficiency, Factor XI deficiency, bleeding in patients with liver disease, thrombasthenia, Bernard-Soulier syndrome, von Willebrand disease, and intracranial hemorrhage. Generally, the patient is a human or non-human animal (e.g., dog, cat, cow, or horse) patient. Preferably, the patient is human.

20 **[0117]** To characterize the *in vitro* binding properties of a suspected Factor VII/VIIa analog, or an ELP-containing Factor VIIa analog, TF binding assays can be performed as described previously (See, e.g., Chaing et al., 1994, Blood 83(12): 3524-35). Briefly, recombinant human TF can be coated onto Immulon II plates in carbonate antigen buffer overnight at 4°C. BSA is also coated onto the plates for use as a control. ELP-containing Factor VIIa analogs may be added at various concentrations in TBS-T buffer. After several washes, monospecific polyclonal rabbit anti-human FVIIa sera is added and incubated for approximately an hour at room temperature. Next, goat anti-rabbit IgG conjugated to alkaline phosphatase is added, followed by the alkaline phosphatase substrate PNPP, which is used for detection. After subtraction of background, the absorbance at ~ 405 nm is taken to be directly proportional to the degree of Factor VIIa binding to the immobilized TF. These values can then be compared to control plasma containing Factor VIIa.

25 **[0118]** The clotting ability of a Factor VII/VIIa analog or an ELP-containing Factor VIIa analog can be measured in human FVII deficient plasma. In this assay, the ELP-containing Factor VIIa analog diluted to varying concentrations directly into FVII deficient plasma. In a coagulometer, one part plasma $\pm$ a FVIIa analog can be mixed with 2 parts Innovin™ (Dade, Miami, Fla.) prothrombin time reagent (recombinant human tissue factor with phospholipids and CaCl₂). Clot formation is detected optically and time to clotting measured. Clotting time (seconds) is compared to the mean clotting time of FVII-deficient plasma alone and plotted as the fractional clotting time versus FVIIa analog concentration.

35 Therapeutic Proteins

[0119] The present invention further provides therapeutic agents comprising an ELP component and at least one therapeutic protein selected from Table 1. The ELP component and therapeutic protein may be coupled by recombinant fusion or chemical conjugation as described herein. Such therapeutic proteins are listed in Table 1 by protein name and GeneSeq Accession No. The amino acid sequence of each Therapeutic Protein, which is known in the art, is hereby incorporated by reference for each Therapeutic Protein listed in Table 1. Such therapeutic proteins are further described in US patent or PCT publications that are also listed in Table 1, and such US patent and PCT publications are hereby incorporated by reference, especially with respect to the structure of such therapeutic proteins and described functional analogs.

40 **[0120]** Table 1 further describes the biological activity of each listed Therapeutic Protein, as well as an exemplary assay for determining the activity of functional analogs or agents of the invention (e.g., fusion with an ELP component). Generally, functional analogs of therapeutic proteins listed in Table 1 may include functional fragments truncated at the N-terminus and/or C-terminus by from 1 to 10 amino acids, including by 1, 2, 3, 4 or about 5 amino acids. Functional analogs may contain from 1 to 10 amino acid insertions, deletions, and/or substitutions (collectively) with respect to the base sequence (e.g., as listed in Table 1), and in each case retaining the full or partial biological activity (as listed in Table 1) of the therapeutic protein. For example, functional analogs may have 1, 2, 3, 4, or 5 amino acid insertions, deletions, and/or substitutions (collectively) with respect to the base sequence. Such activity may be confirmed or assayed using any available assay, including those described in the Table. In these or other embodiments, the therapeutic protein has at least about 75%, 80%, 85%, 90%, 95%, or 98% identity with the corresponding base sequence. The molecules may further comprise additional chemical modifications known for each in the art.

55 **[0121]** In some embodiments, the therapeutic protein (e.g., as selected from Table 1) has a size of less than about 25 kDa, or less than about 10 kDa, or less than about 5 kDa, and the corresponding therapeutic agent of the invention

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(e.g., comprising the ELP component) has a molecular weight of less than about 60 kDa, 55 kDa, 50 kDa, or 40 kDa.

[0122] Table 1 further lists preferred indications for each therapeutic protein, for which the corresponding therapeutic agent finds use, such as in a method for treatment or prevention related to such indication.

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Table 1

Therapeutic Protein X	Exemplary Identifier (the sequences listed in this column are each hereby incorporated by reference)	PCT/Patent Reference (the patents and publications listed in this column are each hereby incorporated by reference)	Biological Activity	Exemplary Activity Assay (the publications listed in this column are each hereby incorporated by reference)	Preferred Indication Y
BMP-1	GeneSeq Accession P80618	WO8800205	BMP1 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic proteins induce cartilage and bone formation, play an important role in nephrogenesis, and play an important role in the development of many organs, including lung, heart, teeth, gut, skin, and particularly the kidney.	BMP-1 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan;27(1):84-8; Eur J Biochem 1996 Apr 1;237(1):295-302; J Biol Chem. Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-2	GeneSeq Accession P80619	WO8800205	BMP-2 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-2 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1;237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-2B	GeneSeq Accession W24850	US5631142	BMP-2b belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-2b activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1;237(1):295-302; J Biol Cbcre, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes

BMP-4	GeneSeq Accession B02796	WO0020591	BMP-4 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-4 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-5	GeneSeq Accession B02797	WO0020591	BMP-5 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-5 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-6	GeneSeq Accession R32904	US5187076	BMP-6 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-6 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
Osteogenic Protein-1; OP-1; BMP-7	GeneSeq Accession W34783	WO973462	OP-1 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	OP-1 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
Osteogenic Protein-2	GeneSeq Accession R57973	WO9406399	OP-2 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	OP-2 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes

GDP-1	GeneSeq Accession R60961	WO9406449	Members of the TGF-beta family of proteins initiate cell signaling by binding to heteromeric receptor complexes of type I (TbetaRI) and type II (TbetaRII) serine/threonine kinase receptors (reviewed by Massague, J. et al. (1994) Trends Cell Biol. 4:172-178; Miyazono, K. et al. (1994) Adv. Immunol. 55:181-220). Activation of this heteromeric receptor complex occurs when TGF-beta binds to TbetaRII, which then recruits and phosphorylates TbetaRI. Activated TbetaRI then propagates the signal to downstream targets (Chen, F. and Weinberg, R.A. (1995) PNA892:1565-1569; Wrana, J.L. et al. (1994) Nature 370:341-347).	The effect of GDF-1 on signaling can be assayed by treating Primary BAEs transferred with a construct called p3TP-Lux, containing a TGF-beta responsive promoter fused to a reporter gene, and measuring luciferase gene expression (Wrana et al., 1994, Nature 370: 341-347).	Developmental disorders, Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-9	GeneSeq Accession R86903	WO9533830	BMP-9 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-9 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-10	GeneSeq Accession R66202	WO9426893	BMP-10 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-10 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-12	GeneSeq Accession R78734	WO9516035	BMP-12 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-12 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes

BMP-15	GeneSeq Accession W11261	WO9636710	BMP-15 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-15 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-17	GeneSeq Accession Y17870	WO9929718	BMP-17 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-17 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
BMP-18	GeneSeq Accession Y17871	WO9929718	BMP-18 belongs to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP-18 activity can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Induction of Cartilage, Tissue and Bone Growth, and Diabetes
Inhibin alpha	GeneSeq Accession B02806	WO0020591	The inhibin beta A subunit joins the alpha subunit to form a pituitary FSH secretion inhibitor. Inhibin has been shown to regulate gonadal stromal cell proliferation negatively and to have tumour-suppressor activity. In addition, serum levels of inhibin have been shown to reflect the size of granulosa-cell tumors and can therefore be used as a marker for primary as well as recurrent disease.	Tumor suppressor activity of inhibin can be determined using assays known in the art: Matzuk et al., Nature 1992 Nov. 26: 360 (6402); 313-9.	Tumor suppression.
Inhibin beta	GeneSeq Accession H02808	WO0020591	The inhibin beta A subunit joins the alpha subunit to form a pituitary FSH secretion inhibitor. Inhibin has been shown to regulate gonadal stromal cell proliferation negatively and to have tumour-suppressor activity. In addition, serum levels of inhibin have been shown to reflect the size of granulosa-cell tumors and can therefore be used as a marker for primary as well as recurrent disease.	Tumor suppressor activity of inhibin can be determined using assays known in the art: Matzuk et al., Nature 1992 Nov. 26: 360 (6402); 313-9.	Tumor suppression.

Cerebus Protein	GeneSeq Accession W86032	WO9849296	Cerebus is believed to be involved in the inhibition of BMP activity	BMP activity, in the presence of the antagonist Cerebus, can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem. 1996 Apr 1; 237(1):295-302; J Biol Chem. Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	BMP Antagonist useful for Osteosarcoma, abnormal bone growth.
Soluble BMP Receptor Kinase Protein-3	GeneSeq Accession R95227	WO9614579	Soluble BMP receptor kinase protein-3 is involved in the binding of BMPs. Soluble BMP receptor kinase protein-3 is useful as an antagonist for the inhibition of BMP activity.	BMP activity, in the presence of the soluble antagonist BMP receptor kinase protein-3, can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem. 1996 Apr 1; 237(1):295-302; J Biol Chem. Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	BMP Antagonist useful for Osteosarcoma, abnormal bone growth.
BMP Processing Enzyme Furin	GeneSeq Accession W36099	WO9741250	BMPs belong to the transforming growth factor-beta (TGFB) superfamily. Bone morphogenic protein induces bone formation.	BMP activity, in the presence of the Furin, can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem. 1996 Apr 1; 237(1):295-302; J Biol Chem. Vol. 274, Issue 16, 10897-10902, April 16, 1999; and Hogan, B. L. M. (1996) Genes Dev. 10, 1580-1594.	Bone formation or Regeneration Abnormalities
TGF-beta 1	GeneSeq Accession R29657	WO9216228	Members of the TGF-beta family of proteins initiate cell signaling by binding to heteromeric receptor complexes of type I (TbetaRI) and type II (TbetaRII) serine/threonine kinase receptors (reviewed by Massague, J. et al. (1994) Trends Cell Biol. 4:172-178; Miyazono, K. et al. (1994) Adv. Immunol. 55:181-220). Activation of this heteromeric receptor complex occurs when TGF-beta. binds to TbetaRII, which then recruits and phosphorylates TbetaRI. Activated TbetaRI then propagates the signal to downstream targets (Chen, F. and Weinberg, R.A. (1995) PNA892:1565-1569; Wrana, J. L. et al. (1994) Nature 370:341.	The effect of TGF betas on signaling can be assayed by treating Primary BAECS transfected with a construct called p3TP-Lux, containing a TGF-beta responsive promoter fused to a reporter gene, and measuring luciferase gene expression (Wrana et al., 1994, Nature 370: 341-347).	Useful for treating cancer and to promote wound healing.

TGF-beta 2	GeneSeq Accession R39659	EP542679	Members of the TGF-beta family of proteins initiate cell signaling by binding to heteromeric receptor complexes of type I (TbetaRI) and type II (TbetaRII) serine/threonine kinase receptors (reviewed by Massague, J. et al. (1994) Trends Cell Biol. 4:172-178; Miyazono, K. et al. (1994) Adv. Immunol. 55:181-220). Activation of this heteromeric receptor complex occurs when TGF-beta. binds to TbetaRII, which then recruits and phosphorylates TbetaRI. Activated TbetaRI then propagates the signal to downstream targets (Chen, F. and Weinberg, R.A. (1995) PNA892:1565-1569; Wrana, J. L. et al. (1994) Nature 370:341.	The effect of TGF betas on signaling can be assayed by treating Primary BAEs transfected with a construct called p3TP-Lux, containing a TGF-beta responsive promoter fused to a reporter gene, and measuring luciferase gene expression (Wrana et al., 1994, Nature 370: 341-347).	Useful for treating cancer and to promote wound healing.
ZTGF-beta 9	GeneSeq Accession Y70654	WO0015798	Members of the TGF-beta family of proteins initiate cell signaling by binding to heteromeric receptor complexes of type I (TbetaRI) and type II (TbetaRII) serine/threonine kinase receptors (reviewed by Massague, J. et al. (1994) Trends Cell Biol. 4:172-178; Miyazono, K. et al. (1994) Adv. Immunol. 55:181-220). Activation of this heteromeric receptor complex occurs when TGF-beta. binds to TbetaRII, which then recruits and phosphorylates TbetaRI. Activated TbetaRI then propagates the signal to downstream targets (Chen, F. and Weinberg, R.A. (1995) PNA892:1565-1569; Wrana, J. L. et al. (1994) Nature 370:341.	The effect of TGF betas on signaling can be assayed by treating Primary BAEs transfected with a construct called p3TP-Lux, containing a TGF-beta responsive promoter fused to a reporter gene, and measuring luciferase gene expression (Wrana et al., 1994, Nature 370: 341-347).	Useful for treating cancer and to promote wound healing.

Anti-TGF beta family antibodies		GB2305921	Members of the TGF-beta family of proteins initiate cell signaling by binding to heteromeric receptor complexes of type I (TbetaRI) and type II (TbetaRII) serine/threonine kinase receptors (reviewed by Massague, J. et al. (1994) Trends Cell Biol. 4:172-178; Miyazono, K. et al. (1994) Adv. Immunol. 55:181-220). Activation of this heteromeric receptor complex occurs when TGF-beta. binds to TbetaRII, which then recruits and phosphorylates TbetaRI. Activated TbetaRI then propagates the signal to downstream targets (Chen, F. and Weinberg, R.A. (1995) PNA892:1565-1569; Wrana, J. L. et al. (1994) Nature 370:341.	The effect of TGF betas on signaling in the presence of an anti-TGF beta antibody, can be assayed by treating Primary BAECS transfected with a construct called p3TP-Lux, containing a TGF-beta responsive promoter fused to a reporter gene, and measuring luciferase gene expression (Wrana et al., 1994, Nature 370: 341-347).	Useful for control of fibrosis, immune, and inflammatory disease.
Latent TGF beta binding protein II	GeneSeq Accession Y70552	WO0012551	Members of the TGF-beta family of proteins initiate cell signaling by binding to heteromeric receptor complexes of type I (TbetaRI) and type II (TbetaRII) serine/threonine kinase receptors (reviewed by Massague, J. et al. (1994) Trends Cell Biol. 4:172-178; Miyazono, K. et al. (1994) Adv. Immunol. 55:181-220). Activation of this heteromeric receptor complex occurs when TGF-beta. binds to TbetaRII, which then recruits and phosphorylates TbetaRI. Activated TbetaRI then propagates the signal to downstream targets (Chen, F. and Weinberg, R.A. (1995) PNA892:1565-1569; Wrana, J. L. et al. (1994) Nature 370:341.	The effect of TGF betas on signaling in the presence of a TGF beta binding protein, can be assayed by treating Primary BAECS transfected with a construct called p3TP-Lux, containing a TGF-beta responsive promoter fused to a reporter gene, and measuring luciferase gene expression (Wrana et al., 1994, Nature 370: 341-347).	Useful for inhibiting tissue or tumor growth.

MP52	GeneSeq Accession W36100	WO9741250	Members of the TGF-beta family of proteins initiate cell signaling by binding to heteromeric receptor complexes of type I (TbetaRI) and type II (TbetaRII) serine/threonine kinase receptors (reviewed by Massague, J. et al. (1994) Trends Cell Biol. 4:172-178; Miyazono, K. et al. (1994) Adv. Immunol. 55:181-220). Activation of this heteromeric receptor complex occurs when TGF-beta. binds to TbetaRII, which then recruits and phosphorylates TbetaRI. Activated TbetaRI then propagates the signal to downstream targets (Chen, F. and Weinberg, R.A. (1995) PNA892:1565-1569; Wrana, J. L. et al. (1994) Nature 370:341.	The effect of TGF betas on signaling can be assayed by treating Primary BAEs transfected with a construct called p3TP-Lux, containing a TGF-beta responsive promoter fused to a reporter gene, and measuring luciferase gene expression (Wrana et al., 1994, Nature 370: 341-347).	Bone formation or Regeneration Abnormalities
b57 Protein	GeneSeq Accession W69293	WO9837195	BMPs are involved in the induction of bone formation. Specific antagonists are useful in preventing this activity from occurring.	BMP activity, in the presence of b57 protein, can be determined using the following assays known in the art: Nat Genet. 2001 Jan; 27(1):84-8; Eur J Biochem 1996 Apr 1; 237(1):295-302; J Biol Chem, Vol. 274, Issue 16, 1089-10902, April 16, 1999; and Hogan, B.L.M. (1996) Genes Deve. 10, 1580-1594.	BMP Antagonist useful for Osteosarcoma, abnormal bone growth.
Resistin	GeneSeq Accession W69293	WO0064920	This gene belongs to the family defined by mouse FIZZ1 and FIZZ3/Resistin genes. The characteristic feature of this family is the C-terminal stretch of 10 cys residues with identical spacing. The mouse homolog of this protein is secreted by adipocytes, may be the hormone potentially linking obesity to type II diabetes.	Ability of resistin to influence type II diabetes can be determined using assays known in the art: Pontoglio et al., J Clin Invest 1998 May 15; 101(10):2215-22.	Type II diabetes and Syndrome X.
Galectin-4	GeneSeq Accession W11841	WO9703190	Galectins are a family of carbohydrate-binding proteins characterized by an affinity for beta-galactoside containing glycoconjugates.	Ability of Galectin-4 polypeptides to bind lactose can be determined using assays known in the art: Wada, et al., J Biol Chem 1997 Feb 28; 272(9):6078-86.	Lactose intolerance.
APM-I; ACRP-30; Famoxin	GeneSeq Accession Y71035	W00026363	ACRP30 gene is exclusively expressed in adipose tissue. ACRP30 is thought to increase fatty acid oxidation by muscle tissue.	Ability of ACRP30 polypeptides to influence obesity and fat oxidation can be determined using assays known in the art: Fruebis et al., Proc Natl Acad Sci USA 2001 Feb 13; 98(4):2005-10.	Obesity, Metabolic disorders, Lipid Metabolism; Hormone Secretion.

ACRP-30 Homologue; Complement Clq C	GeneSeq Accession B30234	WO0063376	ACRP30 gene is exclusively expressed in adipose tissue. ACRP30 is thought to increase fatty acid oxidation by muscle tissue.	Ability of ACRP30 homologue polypeptides to influence obesity and fat oxidation can be determined using assays known in the art: Fruebis et al., Proc Natl Acad Sci USA 2001 Feb 13; 98(4):2005-10.	Obesity, Metabolic disorders, Lipid Metabolism; Hormone Secretion.
Calpain-10a	GeneSeq Accession Y79567	WO0023603	Calpain is believed to play a role in insulin secretion and insulin activity, and therefore may be useful in the treatment of type II diabetes.	Ability of Calpain-10 to influence type II diabetes can be determined using assays known in the art: Pontoglio et al., J Clin Invest 1998 May 15; 101(10):2215-22.	Diabetes mellitus; Regulation of Insulin secretory response; Insulin mediated glucose transport disorders.
Calpain-10b	GeneSeq Accession Y79568	WO0023603	Calpain is believed to play a role in insulin secretion and insulin activity, and therefore may be useful in the treatment of type II diabetes.	Ability of Calpain-10 to influence type II diabetes can be determined using assays known in the art: Pontoglio et al., J Clin Invest 1998 May 15; 101(10):2215-22.	Diabetes mellitus; Regulation of Insulin secretory response; Insulin mediated glucose transport disorders.
Calpain-10c	GeneSeq Accession Y79569	WO0023603	Calpain is believed to play a role in insulin secretion and insulin activity, and therefore may be useful in the treatment of type II diabetes.	Ability of Calpain-10 to influence type II diabetes can be determined using assays known in the art: Pontoglio et al., J Clin Invest 1998 May 15; 101(10):2215-22.	Diabetes mellitus; Regulation of Insulin secretory response; Insulin mediated glucose transport disorders.
PDGF-D	GeneSeq Accession Y71130	WO0027879	Vascular Endothelial Growth Factor.	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266).	Wound Healing; Atherosclerosis.
FasL	GeneSeq Accession Y28594	WO9936079	Activities associated with apoptosis and immune system functions.	Activity can be determined using Apoptosis assays known in the art: Walczak et al. (1996) EMBOJ 16: 5386-5397.	Apoptosis-related disorders; Autoimmune disorders; Graft v-Host disorders.
Chondro modulin-like protein	GeneSeq Accession Y71262	WO0029579	Chondromodulin proteins are cartilage proteins thought to confer resistance to angiogenesis, and thus are useful as anti-angiogenic agents that may have utility in combating cancer.	Ability of Chondromodulin-like protein to inhibit vascularization can be determined using assays known in the art: Hirakie et al., J Biol Chem 1997 Dec 19; 272(51):32419-26.	Antiangiogenic agent; Osteoblast proliferation stimulator; prevents vascularization of cartilage tissue; Useful to treat cancer.
Patched	GeneSeq Accession W72969	US5837538	Patched is a tumour-suppressor receptor for Sonic hedgehog (shh), which is a protein that controls developmental patterning and growth.	Ability of soluble Patched to bind to and inhibit the activities of shh can be determined using assays known in the art: Stone et al., Nature 1996 Nov 14; 384(6605):129-34.	Receptor for Hedgehog cellular proliferation signaling molecule. This receptor is useful as a means of preventing cellular proliferation via the shh signaling pathway, thus useful for cancers.

Patched-2	GeneSeq Accession Y43261	WO9953058	Patched is a tumour-suppressor receptor for Sonic hedgehog (shh), which is a protein that controls developmental patterning and growth.	Ability of soluble Patched to bind to and inhibit the activities of shh can be determined using assays known in the art: Stone et al., Nature 1996 Nov 14; 384(6605):129-34.	Receptor for Hedgehog cellular proliferation signaling molecule. This receptor is useful as a means of preventing cellular proliferation via the shh signaling pathway, thus useful for cancers.
Maspin; Protease Inhibitor 5	GeneSeq Accession R50938	WO9405804	Maspin is a member of the serpin family of serine protease inhibitors that is thought to suppress tumor metastasis.	The inhibitory effects of Maspin and other protease inhibitors can be assayed using methods known in the art such as a labeled protease substrate, for example, Universal Protease Substrate (casein, resorufin-labeled); Roche Molecular Biochemicals, Cat. No. 1080733.	Tumor suppressor which is down-regulated in breast cancers. The maspin protein has tumour suppressing and invasion suppressing activity.
Endostatin	GeneSeq Accession B28399	WO0064946	Endostatin is believed to inhibit effects of capillary endothelial cell proliferation.	The inhibitory effects of endostatin can be assayed using assays disclosed by Cao et al. (1996) J. Biol. Chem. 271 29461-29467.	Anti-angiogenic activity. Useful in the prevention and/or treatment of cancers.
aFGF; FGF-1	GeneSeq Accession P94037	EP298723	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti-cancer agents.
bFGF; FGF-2	GeneSeq Accession R06685	FR2642086	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti-cancer agents.
FGF-3; INT-2	GeneSeq Accession R07824	WO9503831	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti-cancer agents.
FGF-4; HST-1; HBGF-4	GeneSeq Accession R07825	WO9503831	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti-cancer agents.

FGF-5	GeneSeq Accession W22600	WO9730155	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti- cancer agents.
FGF-6; Heparin binding secreted transforming factor-2	GeneSeq Accession R58555	EP613946	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti- cancer agents.
FGF-8	GeneSeq Accession R80783	WO9524928	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti- cancer agents.
FGF-9; Gila activating factor	GeneSeq Accession R70822	WO9503831	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti- cancer agents.
FGF-12; Fibroblast growth factor homologous factor-1	GeneSeq Accession W06309	WO9635708	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti- cancer agents.
FGF-15	GeneSeq Accession Y08582	WO9927100	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti- cancer agents.
FGF-16	GeneSeq Accession Y05474	WO9918128	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti- cancer agents.

FGF-18	GeneSeq Accession Y08590	WO9927100	Fibroblast Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266); Examples 23 and 39 disclosed herein.	Promotion of growth and proliferation of cells, such as epithelial cells and keratinocytes. Antagonists may be useful as anti-cancer agents.
flt-3 ligand	GeneSeq Accession R67541	EP627487	Stem Cell Progenitor	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Promotion of immune cell growth and/or differentiation.
VEGF-110	GeneSeq Accession Y69417	WO0013702	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGB-121	GeneSeq Accession B50432	WO0071713	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGF-138	GeneSeq Accession Y43483	WO9940197	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGF-145	GeneSeq Accession Y69413	WO0013702	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.

VEGF-162	GeneSeq Accession Y43484	W09940197	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGF-165	GeneSeq Accession Y69414	W00013702	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGF-182	GeneSeq Accession Y43483	W09940197	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGF-189	GeneSeq Accession Y69415	W00013702	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGF-206	GeneSeq Accession Y69416	W00013702	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGF-D	GeneSeq Accession W53240	W09807832	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.

VEGF-E; VEGF-X	GeneSeq Accession Y33679	W09947677	Promotes the growth and/or proliferation of endothelial cells.	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
VEGF Receptor; KDR; flk-1	GeneSeq Accession W69679	W09831794	Receptor for VEGF polypeptides	VEGF activity, in the presence of flk-1 polypeptides, can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	VEGF Receptor. Fusion protein with the extracellular domain is useful as an anti- angiogenic agent. Antagonists may be useful in the promotion of angiogenesis.
Soluble VEGF Receptor	GeneSeq Accession W47037	US5712380	Receptor for VEGF polypeptides	VEGF activity, in the presence of VEGF Receptor polypeptides, can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	VEGF Receptor. Fusion protein with the extracellular domain is useful as an anti- angiogenic agent. Antagonists may be useful in the promotion of angiogenesis.
flt-1	GeneSeq Accession Y70751	W00021560	Receptor for VEGF polypeptides	VEGF activity, in the presence of flt-1 polypeptides, can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	VEGF Receptor. Fusion protein with the extracellular domain is useful as an anti- angiogenic agent. Antagonists may be useful in the promotion of angiogenesis.
VEGF R-3; flt-4	GeneSeq Accession B29047	W00058511	Receptor for VEGF polypeptides	VEGF activity, in the presence of flt-4 polypeptides, can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	VEGF Receptor. Fusion protein with the extracellular domain is useful as an anti- angiogenic agent. Antagonists may be useful in the promotion of angiogenesis.

Neuropilin-1	GeneSeq Accession Y06319	WO9929858	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
Neuropilin-2	GeneSeq Accession Y03618	WO9929858	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
Human fast twitch skeletal muscle troponin C	GeneSeq Accession W22597	W09730085	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis
Human fast twitch skeletal muscle troponin I	GeneSeq Accession W18054	W09730085	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis
Human fast twitch skeletal muscle troponin T	GeneSeq Accession W22599	W09730085	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis
fragment. myofibrillar protein troponin I	GeneSeq Accession W18053	W09719955	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis
myofibrillar protein troponin I	GeneSeq Accession W18054	W09719955	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis

Troponin peptides	GeneSeq Accessions Y29581, Y29582, Y29583, Y29584, Y29585, and Y29586	WO9933874	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis
Human fast twitch skeletal muscle Troponin subunit C	GeneSeq Accession B00134	WO0054770	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis
Human fast twitch skeletal muscle Troponin subunit I Protein	GeneSeq Accession B00135	WO0054770	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis
Human fast twitch skeletal muscle Troponin subunit T	GeneSeq Accession B00136	WO0054770	Troponins are contractile proteins that are thought to inhibit angiogenesis. High levels may contribute to the difficulty encountered in revascularizing the ischemic myocardium after cardiovascular injury.	Ability of soluble Troponins to inhibit angiogenesis can be determined using assays known in the art.: Proc Natl Acad Sci U S A 1999 Mar 16;96(6):2645-50.	Anti-angiogenesis
Activator Inhibitor-1; PAI-1	GeneSeq Accession R08411	WO9013648	PAIs are believed to play a role in cancer, and cardiovascular disease and blood-clotting disorders.	Methods that measure plasminogen activator inhibitor (PAI) activity are known in the art, for example, assay the ability of PAI to inhibit tissue plasminogen activator (tPA) or urokinase (uPA): J Biochem Biophys Methods 2000 Sep 11; 45(2):127-40, Breast Cancer Res Treat 1996;41(2):141-6. Methods that measure anti-angiogenesis activity are known in the art, for example, Proc Natl Acad Sci U S A 1999 Mar 16; 96(6):2645-50.	Anti-angiogenesis; blood-clotting disorders.

Plasminogen Activator Inhibitor- 2; PAI-2	GeneSeq Accession P94160	DE3722673	PAIs are believed to play a role in cancer, and cardiovascular disease and blood- clotting disorders.	Methods that measure plasminogen activator inhibitor (PAI) activity are known in the art, for example, assay the ability of PAI to inhibit tissue plasminogen activator (tPA) or urokinase (uPA): J Biochem Biophys Methods 2000 Sep 11; 45(2):127- 40, Breast Cancer Res Treat 1996;41(2):141-6. Methods that measure anti-angiogenesis activity are known in the art, for example, Proc Natl Acad Sci U S A 1999 Mar 16; 96(6):2645-50.	Anti-angiogenesis; blood- clotting disorders.
Activator Inhibitor- 2; PAI-2	GeneSeq Accession R10921	WO9102057	PAIs are believed to play a role in cancer, and cardiovascular disease and blood- clotting disorders.	Methods that measure plasminogen activator inhibitor (PAI) activity are known in the art, for example, assay the ability of PAI to inhibit tissue plasminogen activator (tPA) or urokinase (uPA): J Biochem Biophys Methods 2000 Sep 11; 45(2):127- 40, Breast Cancer Res Treat 1996;41(2):141-6. Methods that measure anti-angiogenesis activity are known in the art, for example, Proc Natl Acad Sci U S A 1999 Mar 16; 96(6):2645-50.	Anti-angiogenesis; blood- clotting disorders.
Human PAI-1 mutants	GeneSeq Accessions R11755, R11756, R11757, R11758, R11759, R11760, R11761, R11762 and R11763	WO9105048	PAIs are believed to play a role in cancer, and cardiovascular disease and blood- clotting disorders.	Methods that measure plasminogen activator inhibitor (PAI) activity are known in the art, for example, assay the ability of PAI to inhibit tissue plasminogen activator (tPA) or urokinase (uPA): J Biochem Biophys Methods 2000 Sep 11; 45(2):127- 40, Breast Cancer Res Treat 1996;41(2):141-6. Methods that measure anti-angiogenesis activity are known in the art, for example, Proc Natl Acad Sci U S A 1999 Mar 16; 96(6):2645-50.	Anti-angiogenesis; blood- clotting disorders.

CXCR3; CXC	GeneSeq Accession Y79372	WO0018431	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Soluble CXCR3 polypeptides may be useful for inhibiting chemokine activities and viral infection.
Modified Rantes	GeneSeq Accession W38129	WO9737005	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
RANTES	GeneSeq Accession Y05299	EP905240	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.

MCI-Ia	GeneSeq Accession R73914	WO9509232	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
MCP-Ib	GeneSeq Accession Y26176	WO9929728	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
MCP-I receptor	GeneSeq Accession R79165	WO9519436	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Soluble MCP-1 Receptor polypeptides may be useful for inhibiting chemokine activities and viral infection.

MCP-3	GeneSeq Accession R73915	W09509232	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
MCP-4 receptor	GeneSeq Accession W56689	W09809171	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Soluble MCP-4 Receptor polypeptides may be useful for inhibiting chemokine activities and viral infection.
RANTES receptor	GeneSeq Accession W29588	US5652133	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Soluble RANTES Receptor polypeptides may be useful for inhibiting chemokine activities and viral infection.

CCR5 variant	GeneSeq Accession W88238	WO9854317	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Soluble CCR5 polypeptides may be useful for inhibiting chemokine activities and viral infection.
CCR7	GeneSeq Accession B50859	US6153441	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Soluble CCR7 polypeptides may be useful for inhibiting chemokine activities and viral infection.
CXC3	GeneSeq Accession W23345	WO9727299	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.

Eotaxin	GeneSeq Accession W10099	WO9700960	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
Neurotactin	GeneSeq Accessions Y77537, W34307, Y53259, and, Y77539	US6013257 WO9742224	Neurotactin may play a role in chemotactic leukocyte migration and brain inflammation processes.	Chemotactic leukocyte migration assays are known in the art, for example: J. Immunol. Methods 33, ((1980)); Nature 1997 Jun 5; 387(6633):611-7.	Immune disorders.
Human CKbeta-9	GeneSeq Accession B50860	US6153441	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
Lymphotactin	GeneSeq Accession B50052	WO0073320	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.

MIP-3 alpha	GeneSeq Accession W44398	WO9801557	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
MIP-3 beta	GeneSeq Accession W44399	WO9801557	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
MIP-Gamma	GeneSeq Accession R70798	WO9504158	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.
Stem Cell Inhibitory Factor	GeneSeq Accession R11553	WO9104274	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Hematopoietic growth factors.

thrombopoietin	GeneSeq Accession R79905	WO9521920	Thrombopoietin is involved in the regulation of the growth and differentiation of megakaryocytes and preceptors thereof.	Thrombopoietin (TPO) can be assayed to determine regulation of growth and differentiation of megakaryocytes. Mol Cell Biol 2001 Apr; 21(8):2659-70; Exp Hematol 2001 Jan; 29(1):51-8 and within.	Hematopoietic growth factors.
c-kit ligand; SCF; Mast cell growth factor; MGF; Fibrosarcoma-derived stem cell factor	GeneSeq Accession Y53284, R83978 and R83977	EP992579 and EP676470	C-kit ligand is thought to stimulate the proliferation of mast cells, and is able to augment the proliferation of both myeloid and lymphoid hematopoietic progenitors in bone marrow culture. C-kit ligand is also thought to act synergistically with other cytokines.	Chemokine activities can be determined using assays known in the art. Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Hematopoietic growth factors.
Platelet derived growth factor	GeneSeq Accession B48653	WO0066736	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
Melanoma inhibiting protein	GeneSeq Accession R69811	WO9503328	Melanoma inhibiting protein has melanoma-inhibiting activity and can be used to treat cancer (melanoma, glioblastoma, neuroblastoma, small cell lung cancer, neuroectodermal tumors) or as an immunosuppressant (it inhibits IL-2 or phytohaemagglutinin induced proliferation of peripheral blood lymphocytes).	Tumor suppressor activity of melanoma inhibiting protein can be determined using assays known in the art. Matzuk et al., Nature 1992 Nov 26;360(6402):313-9.	Cancer; melanoma
Glioma-derived growth factor	GeneSeq Accession R08120	EP399816	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
Platelet derived growth factor precursor A	GeneSeq Accession R84759	EP682110	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.

Platelet derived growth factor precursor B	GeneSeq Accession R84760	EP682110	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
Platelet derived growth factor Bv-sis	GeneSeq Accession P80595 and P80596	EP282317	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
Placental Growth Factor	GeneSeq Accessions R23059 and R23060	WO9206194	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
Placental Growth Factor-2	GeneSeq Accession Y08289	DE19748734	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer.
Thrombopoietin derivative1	GeneSeq Accession Y77244	WO0000612	Thrombopoietin is involved in the regulation of the growth and differentiation of megakaryocytes and precursors thereof.	Thrombopoietin (TPO) can be assayed to determine regulation of growth and differentiation of megakaryocytes. Mol Cell Biol 2001 Apr; 21(8):2659-70; Exp Hematol 2001 Jan; 29(1):51-8 and within.	Thrombocytopenia, cancer.
Thrombopoietin derivative2	GeneSeq Accession Y77255	WO0000612	Thrombopoietin is involved in the regulation of the growth and differentiation of megakaryocytes and precursors thereof.	Thrombopoietin (TPO) can be assayed to determine regulation of growth and differentiation of megakaryocytes. Mol Cell Biol 2001 Apr; 21(8):2659-70; Exp Hematol 2001 Jan; 29(1):51-8 and within.	Thrombocytopenia, cancer.
Thrombopoietin derivative3	GeneSeq Accession Y77262	WO0000612	Thrombopoietin is involved in the regulation of the growth and differentiation of megakaryocytes and precursors thereof.	Thrombopoietin (TPO) can be assayed to determine regulation of growth and differentiation of megakaryocytes. Mol Cell Biol 2001 Apr; 21(8):2659-70; Exp Hematol 2001 Jan; 29(1):51-8 and within.	Thrombocytopenia, cancer.

Thrombopoietin derivative4	GeneSeq Accession Y77267	WO00000612	Thrombopoietin is involved in the regulation of the growth and differentiation of megakaryocytes and precursors thereof.	Thrombopoietin (TPO) can be assayed to determine regulation of growth and differentiation of megakaryocytes. Mol Cell Biol 2001 Apr; 21(8):2659-70; Exp Hematol 2001 Jan; 29(1):51-8 and within.	Thrombocytopenia, cancer.
Thrombopoietin derivative5	GeneSeq Accession Y77246	WO00000612	Thrombopoietin is involved in the regulation of the growth and differentiation of megakaryocytes and precursors thereof.	Thrombopoietin (TPO) can be assayed to determine regulation of growth and differentiation of megakaryocytes. Mol Cell Biol 2001 Apr; 21(8):2659-70; Exp Hematol 2001 Jan; 29(1):51-8 and within.	Thrombocytopenia, cancer.
Thrombopoietin derivative6	GeneSeq Accession Y77253	WO00000612	Thrombopoietin is involved in the regulation of the growth and differentiation of megakaryocytes and precursors thereof.	Thrombopoietin (TPO) can be assayed to determine regulation of growth and differentiation of megakaryocytes. Mol Cell Biol 2001 Apr; 21(8):2659-70; Exp Hematol 2001 Jan; 29(1):51-8 and within.	Thrombocytopenia, cancer.
Thrombopoietin derivative7	GeneSeq Accession Y77256	WO00000612	Thrombopoietin is involved in the regulation of the growth and differentiation of megakaryocytes and precursors thereof.	Thrombopoietin (TPO) can be assayed to determine regulation of growth and differentiation of megakaryocytes. Mol Cell Biol 2001 Apr; 21(8):2659-70; Exp Hematol 2001 Jan; 29(1):51-8 and within.	Thrombocytopenia, cancer.
Fractalkine	GeneSeq Accession Y53255	US6043086	Fractalkine is believed to play a role in chemotactic leukocyte migration and neurological disorders.	Fractalkine activity can be determined using Chemotactic leukocyte migration assays known in the art, for example: J. Immunol. Methods 33, ((1980)); Nature 1997 Jun 5;387(6633):611-7.	Immune disorders.
CXC3	GeneSeq Accession W23345	WO9757599	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Immune disorders.

CCR7	GeneSeq Accession B50859	US6153441	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ.	Soluble CCR7 polypeptides may be useful for inhibiting chemokine activities and viral infection.
Nerve Growth Factor-beta	GeneSeq Accession R11474	EP414151	Nerve Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266)	Neurological disorders, cancer
Nerve Growth Factor-beta2	GeneSeq Accession W69725	EP859056	Nerve Growth Factor	Proliferation assay using NR6R 3T3 cells (Rizzino 1988 Cancer Res. 48: 4266)	Neurological disorders, cancer
Neurotrophin-3	GeneSeq Accession W8889	WO9821234	Neurotrophins regulate neuronal cell survival and synaptic plasticity.	Trk tyrosine kinase activation assays known in the art can be used to assay for neurotrophin activity, for example, Proc Natl Acad Sci U S A 2001 Mar 13:98(6):3555-3560.	Neurological disorders, cancer
Neurotrophin-3	GeneSeq Accession R47100	WO9325684	Neurotrophins regulate neuronal cell survival and synaptic plasticity.	Trk tyrosine kinase activation assays known in the art can be used to assay for neurotrophin activity, for example, Proc Natl Acad Sci U S A 2001 Mar 13:98(6):3555-3560.	Neurological disorders, cancer
Neurotrophin-4a	GeneSeq Accession R47101	WO9325684	Neurotrophins regulate neuronal cell survival and synaptic plasticity.	Trk tyrosine kinase activation assays known in the art can be used to assay for neurotrophin activity, for example, Proc Natl Acad Sci U S A 2001 Mar 13:98(6):3555-3560.	Neurological disorders, cancer
Neurotrophin-4b	GeneSeq Accession R47102	WO9325684	Neurotrophins regulate neuronal cell survival and synaptic plasticity. tyrosine kinases.	Trk tyrosine kinase activation assays known in the art can be used to assay for neurotrophin activity, for example, Proc Natl Acad Sci U S A 2001 Mar 13:98(6):3555-3560.	Neurological disorders, cancer
Neurotrophin-4c	GeneSeq Accession R47103	WO9325684	Neurotrophins regulate neuronal cell survival and synaptic plasticity. tyrosine kinases.	Trk tyrosine kinase activation assays known in the art can be used to assay for neurotrophin activity, for example, Proc Natl Acad Sci U S A 2001 Mar 13:98(6):3555-3560.	Neurological disorders, cancer

Neurotrophin-4d	GeneSeq Accession R47102	WO9325684	Neurotrophins regulate neuronal cell survival and synaptic plasticity. tyrosine kinases.	Trk tyrosine kinase activation assays known in the art can be used to assay for neurotrophin activity, for example, Proc Natl Acad Sci U S A 2001 Mar 13;98(6):3555-3560.	Neurological disorders, cancer
Platelet-Derived Growth Factor A chain	GeneSeq Accession R38918	US5219739	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. W00045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Hematopoietic and immune disorders. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer
Platelet-Derived Growth Factor B chain	GeneSeq Accession R38919	US5219739	Vascular Endothelial Growth Factor	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. W00045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Hematopoietic and immune disorders. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer
Stromal Derived Factor-1 alpha	GeneSeq Accession Y39995	WO9948528	Stromal Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266)	Hematopoietic, immune disorders, cancer
Stromal Derived Factor-1 beta	GeneSeq Accession R75420	CA2117953	Stromal Growth Factor	Proliferation assay using NR6R-3T3 cells (Rizzino 1988 Cancer Res. 48: 4266)	Hematopoietic, immune disorders, cancer
Tarc	GeneSeq Accession W14917	WO9711969	Chemotactic for T lymphocytes. May play a role in T-cell development. Thought to bind CCR8 and CCR4	Chemotactic leukocyte migration assays are known in the art, for example: J. Immunol. Methods 33 ((1980))	Antiinflammatory. Immune disorders, cancer
Prolactin	GeneSeq Accession R78691	WO9521625	Prolactin is involved in immune cell proliferation and apoptosis.	Immune cell proliferation and suppression of apoptosis by prolactin can be assayed by methods well-known in the art, for example, Buckley, AR and Buckley DJ, Ann N Y Acad Sci 2000; 917:522-33, and within.	Reproductive system disorders, cancer.
Prolactin2	GeneSeq Accession Y31764	US5955346	Prolactin is involved in immune cell proliferation and apoptosis.	Immune cell proliferation and suppression of apoptosis by prolactin can be assayed by methods well-known in the art, for example, Buckley, AR and Buckley DJ, Ann N Y Acad Sci 2000; 917:522-33, and within.	Reproductive system disorders, cancer.

Follicle stimulating hormone Alpha subunit	GeneSeq Accession Y54160	EP974359	FSH stimulates secretion of interleukin-1 by cells isolated from women in the follicular phase	FSH activities can be determined using assays known in the art; J Gend Specif Med 1999 Nov-Dec;2(6):30-4; Mol Cell Endocrinol. 1997 Nov 15;134(2):109-18.	Reproductive system disorders, cancer.
Follicle stimulating hormone Beta subunit	GeneSeq Accession Y54161	EP974359	FSH stimulates secretion of interleukin-1 by cells isolated from women in the follicular phase	FSH activities can be determined using assays known in the art; J Gend Specif Med 1999 Nov-Dec;2(6):30-4; Mol Cell Endocrinol. 1997 Nov 15;134(2):109-18.	Reproductive system disorders, cancer.
Substance P (tachykinin)	GeneSeq Accession B23027	WO0054053	Substance P is associated with immunoregulation.	Immunoregulation and bone marrow, cell proliferation by substance P can be assayed by methods well-known in the art, for example, Lai et al. Proc Natl Acad Sci USA 2001 Mar 27; 98(7):3970-5; Jallat-Daloz et al. Allergy Asthma Proc 2001 Jan-Feb; 22(1):17-23; Kahler et al. Exp Lung Res 2001 Jan-Feb; 27(1):25-46; and Adamus MA and Dabrowski ZJ. J Cell Biochem 2001; 81(3):499-506.	diabetes mellitus, hypertension, cancer
Oxytocin (Neurophysin I)	GeneSeq Accession B24085 and B24086	WO0053755	Oxytocin is involved in the induction of prostaglandin (E2) release as well as an increased amount of calcium release by smooth muscle cells.	Oxytocin and prostaglandin E(2) release and Oxytocin (Ca2+) increase can be assayed by methods well-known in the art, for example, Pavan et al. AM J Obstet Gynecol 2000 Jul; 183(1):76-82 and Holda et al., Cell Calcium 1996 Jul;20(1):43-51.	inflammatory disorders immunologic disorders, cancer
Vasopressin (Neurophysin II)	GeneSeq Accession B24085 and B24086	WO0053755	Vasopressin is believed to have a direct antidiuretic action on the kidney, and it is thought to cause vasoconstriction of the peripheral vessels.	Vasopressin activity can be determined using assays known in the art, for example, Endocr Regul 1996 Mar;30(1):13-17.	inflammatory disorders immunologic disorders, cancer
IL-1	GeneSeq Accession P60326	EP165654	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) Cytokine 1, 14-20.	inflammatory disorders, immunologic disorders, cancer

IL-1 mature	GeneSeq Accession R14855	EP456332	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) Cytokine 1, 14-20.	inflammatory disorders, immunologic disorders, cancer
IL-1 beta	GeneSeq Accession Y08322	WO9922763	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) Cytokine 1, 14-20.	inflammatory disorders, immunologic disorders, cancer
IL-3 variants	GeneSeq Accession P80382, P80383, P80384, and P80381	WO8806161	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) J Cell Physiol. 140 323-334.	inflammatory disorders, immunologic disorders, cancer
IL-4	GeneSeq Accession P70615	WO8702990	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Siegel & Mostowski (1990) J Immunol Methods 132,287-295.	inflammatory disorders, immunologic disorders, cancer

IL-4 muteins	GeneSeq Accession W52151 W52152 W52153 W52154 W52155 W52156 W52157 W52158 W52159 W52160 W52161 W52162 W52163 W52164 and W52165	WO9747744	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Siegel & Mostowski (1990) <i>J Immunol Methods</i> 132,287-295.	inflammatory disorders, immunologic disorders, cancer
IL-1 alpha	GeneSeq Accession P90108	EP324447	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) <i>Cytokine</i> 1, 14-20.	inflammatory disorders, immunologic disorders, cancer
IL-3 variants	GeneSeq Accession R38561, R38562, R38563, R38564, R38565, R38566, R38567, R38568, R38569, R38570, R38571, and R38572	WO9307171	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Aarden et al (1987) <i>Eur. J. Immunol</i> 17, 1411-16.	inflammatory disorders, immunologic disorders, cancer
IL-6	GeneSeq Accession R45717 and R45718	WO9402512	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Aarden et al (1987) <i>Eur. J. Immunol</i> 17, 1411-16.	inflammatory disorders, immunologic disorders, cancer

IL-13	GeneSeq Accession R48624	WO9404680	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Boutelier et al (1995) J. Immunol. Methods 181,29.	inflammatory disorders, immunologic disorders, cancer
IL-4 mutein	GeneSeq Accession R47182	DE4137333	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Siegel & Mostowski (1990) J Immunol Methods 132,287-295.	inflammatory disorders, immunologic disorders, cancer
IL-4 mutein Y124X	GeneSeq Accession R47183	DE4137333	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Siegel & Mostowski (1990) J Immunol Methods 132,287-295.	inflammatory disorders, immunologic disorders, cancer
IL-4 mutein Y124G	GeneSeq Accession R47184	DE4137333	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Siegel & Mostowski (1990) J Immunol Methods 132,287-295.	inflammatory disorders, immunologic disorders, cancer

Human Interleukin-10 (precursor)	GeneSeq Accession R41664	WO9317698	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Thompson-Snipes et al (1991) J. Exp. Med. 173, 507-510.	inflammatory disorders, immunologic disorders, cancer
Human Interleukin-10	GeneSeq Accession R42642	WO9318783-A	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Thompson-Snipes et al (1991) J. Exp. Med. 173, 507-510.	inflammatory disorders, immunologic disorders, cancer
Human interleukin-1 beta precursor.	GeneSeq Accession R42447	EP569042	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) Cytokine 1, 14-20.	inflammatory disorders, immunologic disorders, cancer
Interleukin-1alpha	GeneSeq Accession R45364	EP578278	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer

Human interleukin-3 variant	GeneSeq Accession R22814	JP04063595	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Mathews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) <i>J Cell Physiol.</i> 140 323-334.	inflammatory disorders, immunologic disorders, cancer
IL-1 β fragments	GeneSeq Accession R35484 and R35485	EP541920	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Mathews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) <i>Cytokine</i> 1, 14-20.	inflammatory disorders, immunologic disorders, cancer
IL-1 inhibitor (IL-1 β)	GeneSeq Accession R35486 and R35484	EP541920	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Mathews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) <i>Cytokine</i> 1, 14-20.	inflammatory disorders, immunologic disorders, cancer
ICE 22kD subunit.	GeneSeq Accession R33780	EP533350	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Mathews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer

ICE 20kD subunit.	GeneSeq Accession R33781	EP5333350	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
ICE 10kD subunit	GeneSeq Accession R33782	EP5333350	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
Human Interleukin-10 (precursor)	GeneSeq Accession R41664	WO9317698	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Thompson-Snipes et al (1991) J. Exp. Med. 173, 507-510.	inflammatory disorders, immunologic disorders, cancer
Human Interleukin-10	GeneSeq Accession R42642	WO9318783	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Thompson-Snipes et al (1991) J. Exp. Med. 173, 507-510.	inflammatory disorders, immunologic disorders, cancer

Human Interleukin-1 beta precursor	GeneSeq Accession R42447	EP569042	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) <i>J Cell Physiol.</i> 140 323-334.	inflammatory disorders, immunologic disorders, cancer
Human interleukin-6	GeneSeq Accession R49041	WO9403492	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Aarden et al (1987) <i>Eur. J. Immunol</i> 17, 1411-16.	inflammatory disorders, immunologic disorders, cancer
Mutant Interleukin 6 S176R	GeneSeq Accession R54990	WO9411402	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Aarden et al (1987) <i>Eur. J. Immunol</i> 17, 1411-16.	inflammatory disorders, immunologic disorders, cancer
Interleukin 6	GeneSeq Accession R55256	JP06145063	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Aarden et al (1987) <i>Eur. J. Immunol</i> 17, 1411-16.	inflammatory disorders, immunologic disorders, cancer

Interleukin 8 (IL-8) receptor	GeneSeq Accession R53932	JP06100595	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Holmes et al (1991) <i>Science</i> 253, 1278-80.	Soluble IL-8 receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-7	GeneSeq Accession R59919	US5328988	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Park et al (1990) <i>J. Exp. Med.</i> 171, 1073-79.	inflammatory disorders, immunologic disorders, cancer
IL-3 containing fusion protein.	GeneSeq Accession R79342 and R79344	WO9521254	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) <i>J Cell Physiol.</i> 140 323-334.	inflammatory disorders, immunologic disorders, cancer

IL-3 mutant proteins	GeneSeq Accession R79254, R79255, R79256, R79257, R79258, R79259, R79260, R79261, R79262, R79263, R79264, R79265, R79266, R79267, R79268, R79269, R79270, R79271, R79272, R79273, R79274, R79275, R79276, R79277, R79278, R79279, R79280, R79281, R79282, R79283, R79284, and R79285	ZA9402636	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Giri et al (1994) EMBO J. 13 2822-2830.	inflammatory disorders, immunologic disorders, cancer
IL-12 p40 subunit.	GeneSeq Accession R63018	AU9466072	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
AGF	GeneSeq Accession R64240	WO9429344	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer

Human interleukin-12 40 kD subunit	GeneSeq Accession R79187	WO9519786	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Hori et al (1987), <i>Blood</i> 70, 1069-1078.	inflammatory disorders, immunologic disorders, cancer
Human interleukin-15 receptor from clone P1	GeneSeq Accession R90843	WO9530695	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Giri et al (1994) <i>EMBO J.</i> 13 2822-2830.	Soluble IL-8 receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-7	GeneSeq Accession R92796	WO9604306	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Park et al (1990) <i>J. Exp. Med.</i> 171, 1073-79.	inflammatory disorders, immunologic disorders, cancer
interleukin-9	GeneSeq Accession R92797	WO9604306	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yang et al (1989) <i>Blood</i> 74, 1880-84.	inflammatory disorders, immunologic disorders, cancer

interleukin-3	GeneSeq Accession R92801	WO9604306	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) <i>J Cell Physiol.</i> 140 323-334.	inflammatory disorders, immunologic disorders, cancer
Human interleukin-5	GeneSeq Accession R92802	WO9604306	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) <i>J Cell Physiol.</i> 140 323-334.	inflammatory disorders, immunologic disorders, cancer
Recombinant interleukin-16	GeneSeq Accession W33373	DE19617202	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Lim et al (1996) <i>J. Immunol.</i> 156, 2566-70.	inflammatory disorders, immunologic disorders, cancer
Human IL-16 protein	GeneSeq Accession W33234	DE19617202	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Lim et al (1996) <i>J. Immunol.</i> 156, 2566-70.	inflammatory disorders, immunologic disorders, cancer

Thrl 17 human interleukin 9	GeneSeq Accession W27521	WO9708321	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Mathews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
Meth 17 human interleukin 9	GeneSeq Accession W27522	WO9708321	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Mathews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yang et al (1989) <i>Blood</i> 74, 1880-84.	inflammatory disorders, immunologic disorders, cancer
Human intracellular IL-1 receptor antagonist.	GeneSeq Accession W77158	EP86-4585	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Mathews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) <i>Cytokine</i> 1, 14-20.	inflammatory disorders, immunologic disorders, cancer
Human interleukin-18 protein (IL-18)	GeneSeq Accession W77158	EP864585	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Mathews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and USHIO et al (1996) <i>J. Immunol.</i> 156, 4274-79.	inflammatory disorders, immunologic disorders, cancer

Human interleukin-18	GeneSeq Accession W77077	EP861663	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and USHIO et al (1996) J. Immunol. 156, 4274-79.	inflammatory disorders, immunologic disorders, cancer
Human interleukin 18 derivatives	GeneSeq Accessions W77083, W77084, W77085, W77086, W77087, W77088, and W77089	EP861663	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Ushio et al (1996) J. Immunol. 156, 4274-79.	inflammatory disorders, immunologic disorders, cancer
Interleukin-9 (IL-9) mature protein (Thr117 version).	GeneSeq Accession W68158	WO9827997	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yang et al (1989) Blood 74, 1880-84.	inflammatory disorders, immunologic disorders, cancer
IL-9 mature protein variant (Met117 version)	GenSeq Accession W68157	WO9827997	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yang et al (1989) Blood 74, 1880-84.	inflammatory disorders, immunologic disorders, cancer

Human IL-9 receptor protein variant #3.	GeneSeq Accession W64058	WO9824904	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yang et al (1989) Blood 74, 1880-84.	inflammatory disorders, immunologic disorders, cancer
Human IL-9 receptor protein variant fragment	GeneSeq Accession W64060	WO9824904	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yang et al (1989) Blood 74, 1880-84.	Soluble IL-9 receptor polypeptides may be useful for inhibiting interleukin activities.
Human IL-9 receptor protein variant #3.	GeneSeq Accession W64061	WO9824904	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yang et al (1989) Blood 74, 1880-84.	Soluble IL-9 receptor polypeptides may be useful for inhibiting interleukin activities.
Human Interleukin-12 p40 protein	GeneSeq Accession W51311	WO9817689	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Hori et al (1987), Blood 70, 1069-1078.	inflammatory disorders, immunologic disorders, cancer

Human Interleukin-12 p35 protein	GeneSeq Accession W51312	WO9817689	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Hori et al (1987), <i>Blood</i> 70, 1069-1078.	inflammatory disorders, immunologic disorders, cancer
Human protein with IL-16 activity	GeneSeq Accession W63753	DE19649233-	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Lim et al (1996) <i>J. Immunol.</i> 156, 2566-70.	inflammatory disorders, immunologic disorders, cancer
Human protein with IL-16 activity	GeneSeq Accession W59425	DE19649233-	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Lim et al (1996) <i>J. Immunol.</i> 156, 2566-70.	inflammatory disorders, immunologic disorders, cancer
Human interleukin-15	GeneSeq Accession W53878	US5747024	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Giri et al (1994) <i>EMBO J.</i> 13 2822-2830.	inflammatory disorders, immunologic disorders, cancer

Human wild-type interleukin-4 (hIL-4) protein	GeneSeq Accession W52149	WO9747744	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Siegel & Mostowski (1990) <i>J Immunol Methods</i> 132,287-295.	inflammatory disorders, immunologic disorders, cancer
interleukin-4 muteins	GeneSeq Accessions W52150, W52151, W52153, W52154, W52155, W52156, W52157, W52158, W52159, W52160, W52161, W52162, W52163, W52164, W52165, W52166, and W52167	WO9747744	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Siegel & Mostowski (1990) <i>J Immunol Methods</i> 132,287-295.	inflammatory disorders, immunologic disorders, cancer
Human interleukin 1 delta	GeneSeq Accession Y28408	WO9935268	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) <i>Cytokine</i> 1, 14-20.	inflammatory disorders, immunologic disorders, cancer

Human interleukin-1 receptor antagonist beta	GeneSeq Accession Y24395	WO9935268	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) Cytokine 1, 14-20.	inflammatory disorders, immunologic disorders, cancer
Human EDIRF II protein sequence	GeneSeq Accession Y22199	WO9932632	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
Human EDIRF I protein sequence	GeneSeq Accession Y22197	WO9932632	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
Human IL-1RD10 protein sequence	GeneSeq Accession Y14131	WO9919480	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) Cytokine 1, 14-20.	Soluble IL-1RD10 receptor polypeptides may be useful for inhibiting interleukin activities.

Human IL-1RD9	GeneSeq Accession Y14122	WO9919480	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) Cytokine 1, 14-20.	Soluble IL-1RD10 receptor polypeptides may be useful for inhibiting interleukin activities.
Human DNAX interleukin-40	GeneSeq Accession Y09196	WO9919491	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
(DIL-40) alternative sequence	GeneSeq Accession Y09197	WO9919491	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
IL-11	GeneSeq Accession R50176	WO9405318	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Lu et al (1994) J Immunol. Methods 173, 19.	inflammatory disorders, immunologic disorders, cancer

Human adipogenesis inhibitory factor	GeneSeq Accession R43260	EP566410	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
IL-11	GeneSeq Accession W02202	JP08127539	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Lu et al (1994) J Immunol. Methods 173, 19.	inflammatory disorders, immunologic disorders, cancer
IL-14	GeneSeq Accession R55800	WO9416074	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Ambrus et al (1993) PNAS 90, 63330-34.	inflammatory disorders, immunologic disorders, cancer
IL-17 receptor	GeneSeq Accession B03807	US6072033	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yao et al (1995) J. Immunol. 155, 5483-86.	Soluble IL-17 receptor polypeptides may be useful for inhibiting interleukin activities.

IL-17	GeneSeq Accession R76573	WO9518826	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yao et al (1995) J. Immunol. 155, 5483-86.	inflammatory disorders, immunologic disorders, cancer
CTLA-8	GeneSeq Accession W13651	WO9704097	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	inflammatory disorders, immunologic disorders, cancer
IL-19	GeneSeq Accession W37935	WO9808870	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Gallagher et al (2000) Genes Immun. 1, 442-50.	inflammatory disorders, immunologic disorders, cancer
IL-21 (TIF)	GeneSeq Accession Y92879	WO0024758	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Parrish-Novak et al (2000) Nature 408, 57-63.	inflammatory disorders, immunologic disorders, cancer

IL-8 receptor	GeneSeq Accession R33420	WO9306229	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Holmes et al (1991) Science 253, 1278-80..	Soluble IL-8 receptor polypeptides may be useful for inhibiting interleukin activities.
Human type II interleukin-1 receptor	GeneSeq Accession R85480	US5464937	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) Cytokine 1, 14-20.	Soluble type II interleukin-1 receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-12 receptor	GeneSeq Accession R69632	EP638644	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Hori et al (1987), Blood 70, 1069-1078.	Soluble IL-12 receptor polypeptides may be useful for inhibiting interleukin activities.
Interleukin 8 receptor B	GeneSeq Accession R80758	US5440021	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Holmes et al (1991) Science 253, 1278-80.	Soluble IL-8 receptor B polypeptides may be useful for inhibiting interleukin activities.

Human IL-8 receptor protein hIL8RA	GeneSeq Accession B09989	JP08103276	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Holmes et al (1991) <i>Science</i> 253, 1278-80.	Soluble IL-8 receptor A polypeptides may be useful for inhibiting interleukin activities.
Human IL-8 receptor protein hIL8R	GeneSeq Accession B09990	JP08103276	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Holmes et al (1991) <i>Science</i> 253, 1278-80.	Soluble IL-8 receptor polypeptides may be useful for inhibiting interleukin activities.
Interleukin-2 receptor associated protein p43	GeneSeq Accession R97569	WO9621732-	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Gillis et al (1978) <i>J. Immunol.</i> 120, 2027.	Soluble IL-2 receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-17 receptor	GeneSeq Accession W04185	WO9629408	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yao et al (1995) <i>J. Immunol.</i> 155, 5483-86.	Soluble IL-17 receptor polypeptides may be useful for inhibiting interleukin activities.

Human interleukin-11 receptor	GeneSeq Accession R99090	WO9619574	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Lu et al (1994) <i>J Immunol. Methods</i> 173, 19.	Soluble IL-11 receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-1 receptor accessory protein	GeneSeq Accession W01911	WO9623067	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) <i>Cytokine</i> 1, 14-20.	Inflammatory disorders, immunologic disorders, cancer
AGF Protein	GeneSeq Accession R92749	US5488032	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	Inflammatory disorders, immunologic disorders, cancer
Human interleukin-1 type-3 receptor	GeneSeq Accession R91064	WO9607739	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Orencole & Dinarello (1989) <i>Cytokine</i> 1, 14-20.	Soluble IL-type-3 receptor polypeptides may be useful for inhibiting interleukin activities

Human interleukin-13 beta receptor	GeneSeq Accession W24972	WO9720926	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Bouteller et al (1995) J. Immunol. Methods 181,29.	Soluble IL-13 beta receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-13 alpha receptor	GeneSeq Accession W24973	WO9720926	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Bouteller et al (1995) J. Immunol. Methods 181,29.	Soluble IL-13 alpha receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-4 receptor	GeneSeq Accession W13499	US5599905	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Siegel & Mostowski (1990) J Immunol Methods 132,287-295.	Soluble IL-4 receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-12 beta-2 receptor	GeneSeq Accession W12771	EP759466	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Hori et al (1987), Blood 70, 1069-1078.	Soluble IL-12 beta-2 receptor polypeptides may be useful for inhibiting interleukin activities.

Human interleukin-12 beta-1 receptor.	GeneSeq Accession W12772	EP759466	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Hori et al (1987), Blood 70, 1069-1078.	Soluble IL-12 beta-1 receptor polypeptides may be useful for inhibiting interleukin activities.
Human IL-9 receptor protein	GeneSeq Accessions W64055, W64056, and W64057	WO9824904	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yang et al (1989), Blood 74, 1880-84..	Soluble IL-9 receptor polypeptides may be useful for inhibiting interleukin activities.
IL-10 receptor	GeneSeq Accession W41804	US5716804	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Thompson-Snipes et al (1991) J. Exp. Med. 173, 507-510.	Soluble IL-10 receptor polypeptides may be useful for inhibiting interleukin activities.
Human IL-6 receptor	GeneSeq Accession Y30938	JP11196867	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Aarden et al (1987) Eur. J. Immunol 17, 1411-16.	Soluble IL-6 receptor polypeptides may be useful for inhibiting interleukin activities.

IL-17 receptor	GeneSeq Accession Y97181	US6096305	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yao et al (1995) <i>J. Immunol.</i> 155, 5483-86.	Soluble IL-17 receptor polypeptides may be useful for inhibiting interleukin activities.
IL-17 receptor	GeneSeq Accession Y97131	US6100235	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Yao et al (1995) <i>J. Immunol.</i> 155, 5483-86.	Soluble IL-17 receptor polypeptides may be useful for inhibiting interleukin activities.
Human interleukin-3 receptor	GeneSeq Accession R25300	EP509826	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) <i>J Cell Physiol.</i> 140 323-334.	Soluble IL-3 receptor polypeptides may be useful for inhibiting interleukin activities.
Human GM-CSF receptor	GeneSeq Accession R10919	WO9102063	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	Soluble GM-CSF receptor polypeptides may be useful for inhibiting interleukin activities.

Human IL-5 receptor alpha chain	GeneSeq Accession R25064	EP492214	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) J Cell Physiol. 140, 323-334.	Soluble IL-5 receptor alpha polypeptides may be useful for inhibiting interleukin activities.
IL-5 receptor	GeneSeq Accession W82842	WO9847923	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Kitamura et al (1989) J Cell Physiol. 140, 323-334.	Soluble IL-5 receptor polypeptides may be useful for inhibiting interleukin activities.
IL-6 receptor	GeneSeq Accession R37215	JP05091892	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Aarden et al (1987) Eur. J. Immunol 17, 1411-16.	Soluble IL-6 receptor polypeptides may be useful for inhibiting interleukin activities.
Human B cell stimulating factor- 2 receptor	GeneSeq Accession P90525	AU8928720	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225.	Soluble B cell stimulating factor-2 receptor polypeptides may be useful for inhibiting interleukin activities.

IL-7 receptor clone	GeneSeq Accession R08330	EP403114	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Park et al (1990) J. Exp. Med. 171, 1073-79.	Soluble IL-7 receptor polypeptides may be useful for inhibiting interleukin activities.
EPO receptor; EPOR	GeneSeq Accession R06512	WO9008822	EPO Receptor is involved in the proliferation and differentiation of erythroblasts.	EPO Receptor activity can be determined using assays known in the art, such as, J Biol Chem 2001 Mar 23;276(12:8995-9002; JAK2 protein tyrosine kinase activity: Blood 1994 Sep 1;84(5):1501-7 and Mol Cell Biol. 1994 Oct;14(10:6506-14.	Inflammatory disorders, immunologic disorders, cancer, erythroblast proliferation and differentiation
IL-15 receptor	GeneSeq Accession R90843	WO9530895	Interleukins are a group of multifunctional cytokines synthesized by lymphocytes, monocytes, and macrophages. Known functions include stimulating proliferation of immune cells (e.g., T helper cells, B cells, eosinophils, and lymphocytes), chemotaxis of neutrophils and T lymphocytes, and/or inhibition of interferons.	Interleukin activity can be determined using assays known in the art: Matthews et al., in <i>Lymphokines and Interferons: A Practical Approach</i> , Clemens et al., eds, IRL Press, Washington, D.C. 1987, pp. 221-225; and Giri et al (1994) EMBO J. 13 2822-2830.	Soluble IL-15 receptor polypeptides may be useful for inhibiting interleukin activities.
CD137/4-1BB Receptor Protein	GeneSeq Accession R70977	WO9507984	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble 4-1BB receptor polypeptides may be useful for inhibiting apoptosis, NF-kB activation, and/or co-stimulation of immune cells such as B and T cells.
BCMA	GeneSeq Accession Y71979	WO0068378	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble BCMA receptor polypeptides may be useful for inhibiting apoptosis, NF-kB activation, and/or co-stimulation of immune cells such as B and T cells.

CD27	GeneSeq Accession R20814	WO9201049	Activities associated with apoptosis, NF- kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble CD27 polypeptides may be useful for inhibiting apoptosis, NF-kB activation, and/or co- stimulation of immune cells such as B and T cells.
CD30	GeneSeq Accession R35478	DE4200043	Activities associated with apoptosis, NF- kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble CD30 polypeptides may be useful for inhibiting apoptosis NF-kB activation, and/or co- stimulation of immune cells such as B and T cells.
CD40	GeneSeq Accession Y33499	WO9945944	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble CD40 polypeptides may be useful for inhibiting apoptosis, NF-kB activation, and/or co- stimulation of immune cells such as B and T cells.
EDAR	Genbank Accession AAD50077		Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Immune Disorders, Lymphomas, X-linked hypohidrotic ectodermal dysplasia
OX40; ACT-4	GeneSeq Accession R74737	WO9512673	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Immune Disorders, Lymphomas, T cell disorders

TACI	GeneSeq Accession W75783	WO9839361	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble TACI receptor polypeptides may be useful for inhibiting apoptosis, NF- κ B activation, and/or co-stimulation of immune cells such as B and T cells.
TNF-R	GeneSeq Accession R10986	AU9058976	Activities associates with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble TNF-R receptor polypeptides may be useful for inhibiting apoptosis, NF- κ B activation, and/or co-stimulation of immune cells such as B and T cells.
TNF-RII; TNF p75 receptor; Death Receptor	GeneSeq Accession R11141	EP418014	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble TNFR-II receptor polypeptides may be useful for inhibiting apoptosis, NF- κ B activation, and/or co-stimulation of immune cells such as B and T cells.
hAPO-4; TROY	GeneSeq Accession W93581	WO9911791	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Immune Disorders, Cancers
TNF-alpha precursor	GeneSeq Accession P60074	EP205038	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer

Human TNF- alpha	GeneSeq Accession R62463	EP619372	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
Human TNF- alpha	GeneSeq Accession R42679	EP563714	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
Human TNF- beta (LT-alpha)	GeneSeq Accession B37799	WO0064479	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
LT-alpha	GeneSeq Accession P70107	EP250000	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
LT-beta	GeneSeq Accession R56869	WO9413808	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer

OPGL	GeneSeq Accession W83195	WO9846751	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer, loss of bone mass
FasL	GeneSeq Accession W98071	WO9903999	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore, et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
FasL	GeneSeq Accession W95041	WO9903998	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
CD27L	GeneSeq Accession R50121	WO9405691	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
CD30 ligand	GeneSeq Accession R45007	WO9324135	Activities associated with apoptosis, NF- κ B activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF- κ B activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer

CD40L	GeneSeq Accession R85486	WO9529935	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
4-1BB ligand	GeneSeq Accession W26657	US5674704	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
FAS Ligand Inhibitory Protein (DcR3)	GeneSeq Accession B19335	WO0058465	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Soluble DcR3 polypeptides may be useful for inhibiting apoptosis, NF-kB activation, and/or co-stimulation of immune cells such as B and T cells.
OX40L	GeneSeq Accession R79903	WO9521915	Activities associated with apoptosis, NF-kB activation, and co-stimulation of immune cells such as T and B cells.	Apoptosis activity, NF-kB activation, and B and T cell co-stimulation can be determined using assays known in the art: Moore et al., 1999, Science, 285(5425):260-3; Song HY et al., 1997 Proc Natl Acad Sci U S A 94(18):9792-6; Epsevik and Nissen-Meyer, 1986, J. Immunol. Methods.	Inflammatory disorders, immunologic disorders, cancer
Protease inhibitor peptides	GeneSeq Accessions R12435, R12436, R12437, R12438, R12439, R12440, and R12444	WO9106561	Peptides that inhibit the function/binding of HIV	HIV protease activities are known in the art: HIV protease assays: EP0387231. One can modify the assay to look for inhibition using any of the disclosed protease inhibitor polypeptides.	HIV, inflammatory disorders, immunologic disorders, cancer, viral infections

Retroviral protease inhibitors	GeneSeq Accessions R06660, R06661, R06662, R06663, R06664, R06665, R06666, R06667, R06668, R06669, R06670, R06671, R06672, R06673, R06674, R06675, and R06676	EP387231	Peptides that inhibit the function/binding of HIV	HIV protease activities are known in the art: HIV protease assays: EP0387231. One can modify the assay to look for inhibition using any of the disclosed protease inhibitor polypeptides.	HIV, inflammatory disorders, immunologic disorders, cancer, viral infections
HIV protease inhibiting peptides	GeneSeq Accessions R59293, R59294, R59295, R59296, R59297, R59298, R59299, R59300, R59301, R59302, R59303, R59304, R59305, R59306, R59307, R59308, R59309, R59310, R59311, R59312, R59313, R59314, R59315, R59316, R59317, R59318, R59319, R59320, R59321, R59322, R59323, R59324, R59325, R59326, R59327, R59328, R59329, R59330, R59331, R59332, R59333, R59334, R59335, R59336, R59337, R59338, R59339, R59340, R59341, R59342, R59343, R59344, R59345, R59346, R59347, R59348, R59349, and R59350	WO9301828	Peptides that inhibit the function/binding of HIV	HIV protease activities are known in the art: HIV protease assays: EP0387231. One can modify the assay to look for inhibition using any of the disclosed protease inhibitor polypeptides.	HIV, inflammatory disorders, immunologic disorders, cancer, viral infections

HIV-1 protease inhibitors	GeneSeq Accessions R86326, R86327, R86328, R86329, R86330, R86331, R86332, R86333, R86334, R86335, R86336, R86337, R86338, R86339, R86340, R86341, R86342, R86343, R86344, R86345, R86346, R86347, R86348, R86349, R86350, R86351, R86352, R86353, R86354, R86355, R86356, R86357, R86358, R86359, R86360, R86361, R86362, R86363, R86364, R86365, R86366, R86367, R86368, R86369, R86370, and R86371	DE4412174	Peptides that inhibit the function/binding of HIV	HIV protease activities are known in the art: HIV protease assays: EP0387231. One can modify the assay to look for inhibition using any of the disclosed protease inhibitor polypeptides.	HIV, inflammatory disorders, immunologic disorders, cancer, viral infections
HIV Inhibitor Peptide	GeneSeq Accession Y89687	WO9959615	Peptides that inhibit the function/binding of HIV	HIV protease activities are known in the art: HIV protease assays: EP0387231. One can modify the assay to look for inhibition using any of the disclosed protease inhibitor polypeptides.	HIV, inflammatory disorders, immunologic disorders, cancer, viral infections
HIV Inhibitor Peptide	GenSeq Accession Y31955	WO9948513	Peptides that inhibit the function/binding of HIV	HIV Protease activities are known in the art: HIV protease assays: EP0387231. One can modify the assay to look for inhibition using any of the disclosed protease inhibitor polypeptides.	HIV, inflammatory disorders, immunologic disorders, cancer, viral infections.
HIV Inhibitor Peptide	www.sciencedirect.com/science/article/pii/S0950268801001126 Published online 12 January 2001; 10.1126/science.1057453		Peptides that inhibit the function/binding of HIV	HIV protease activities are known in the art: HIV protease assays: EP0387231. One can modify the assay to look for inhibition using any of the disclosed protease inhibitor polypeptides.	HIV, inflammatory disorders, immunologic disorders, cancer, viral infections
Human monocyte chemoattractant	GeneSeq Accession	WO9509232	Chemokines are a family of small, secreted proteins involved in biological	Chemokine activities can be determined using assays known in the art: Methods in	Immune disorders, particularly useful for

factor hMCP-3	R73915		processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified.	Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	treating bacterial and/or viral meningitis
Human monocyte chemoattractant factor hMCP-1	GeneSeq Accession R73914	WO9509232	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, particularly useful for treating bacterial and/or viral meningitis
Human gro-beta chemokine	GeneSeq Accessions R6699 and W17671	WO9429341	Chemokines are a family of small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, inflammatory disorders, blood-related disorders, stem cell transplantation, cancer
Human gro-gamma chemokine	GeneSeq Accessions R66700 and W17672	WO9429341	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I.	Immune disorders, inflammatory disorders, blood-related disorders, stem cell transplantation,

				Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified	Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	cancer
Human gro- α chemokine	GeneSeq Accessions R66698 and W18024	WO9429341		Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, inflammatory disorders, blood-related disorders, stem cell transplantation, cancer
Human eosinophil-expressed chemokine (EEC)	GeneSeq Accession W05186	WO9632481		Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, particularly treatment of eosinophilia, inflammation, allergies, asthma, leukaemia and lymphoma
Chemokine-like protein PF4-414 Full-Length and Mature	GeneSeq Accessions R92318 and R99809	WO9613587		Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and blood-related disorders, particularly myelosuppression

Chemokine-like protein IL-8M3	GeneSeq Accession R99812	WO9613587	including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ; and Holmes et al (1991) Science 253, 1278-80.	Cancer and blood-related disorders, particularly myelosuppression
Human interleukin-8 (IL-8)	GeneSeq Accession R99814	WO9613587	including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ; and Holmes et al (1991) Science 253, 1278-80.	Cancer and blood-related disorders, particularly myelosuppression
Chemokine-like protein IL-8M1 Full-Length and Mature	GeneSeq Accessions R99815 and R99803	WO9613587	including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to – 17 receptors thus far identified	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ; and Holmes et al (1991) Science 253, 1278-80.	Cancer and blood-related disorders, particularly myelosuppression

Chemokine-like protein IL - 8M8 Full-Length and Mature	GeneSeq Accessions R99816 and R99805	WO9613587	biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot; T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ, and Holmes et al (1991) Science 253, 1278-80.	Cancer and blood-related disorders, particularly myelosuppression.
Chemokine-like protein IL - 8M8 Full-Length and Mature	GeneSeq Accessions R99817 and R99806	WO9613587	biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified. Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot; T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ, and Holmes et al (1991) Science 253, 1278-80.	Cancer and blood-related disorders, particularly myelosuppression.

Chemokine-like protein IL - 8/M8 Full-Length and Mature	GeneSeq Accessions R99818 and R99804	WO9613587	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ; and Holmes et al (1991) Science 253, 1278-80.	Cancer and blood-related disorders, particularly myelosuppression.
Chemokine-like protein IL - 8/M8 Full-Length and Mature	GeneSeq Accessions R99819 and R99807	WO9613587	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and blood-related disorders, particularly myelosuppression.
Chemokine-like protein IL - 8/M8 Full-Length and Mature	GeneSeq Accessions R99822 and R9807	WO9613587	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and blood-related disorders, particularly myelosuppression.

Human foetal spleen expressed chemokine, FSEC	GeneSeq Accession R98499	WO9622374	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot; T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders
Liver expressed chemokine-1 (LVEC-1)	GeneSeq Accession R95689	WO9616979	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot; T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inflammation of the liver
Liver expressed chemokine-2 (LVEC-2)	GeneSeq Accession R95690	WO9616979	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot; T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inflammation of the liver

Pituitary expressed chemokine (PGEc)	GeneSeq Accession R95691	WO9616879	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inflammation, particularly of the liver
Adenoid- expressed chemokine (ADEC)	GeneSeq Accession R97664	WO9617668	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inflammation, angiogenesis, tumorigenesis, musculoskeletal disorders
Human chemokineCC- 2	GeneSeq Accession W38170	WO9741230	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc. Totowa, NJ	Immune disorders, cell migration, proliferation and differentiation disorders

Human chemokine HCC-1	GeneSeq Accession W38171	WO9741230	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells and C.A. Power © Humana Press Inc., Totowa, NJ	Immune disorders, cell migration, proliferation and differentiation disorders
Human chemokine CC-3	GeneSeq Accession W38172	WO9741230	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune disorders, cell migration, proliferation and differentiation disorders
Novel betachemokine designated PTEC	GeneSeq Accession W27271	WO9739126	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune disorders, vascular disorders, cancer

Human CX3C 111 amino acid chemokine	GeneSeq Accession W23344	WO9727299	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune disorders, inflammatory diseases, abnormal proliferation, regeneration, degeneration, and atrophy
Human CCF18 chemokine	GeneSeq Accession W25942	WO9721812	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Abnormal physiology and development disorders, can also be used as an anti-viral agent
Human beta- chemokine H1305 (MCP-2)	GeneSeq Accession W26655	WO9725427	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Chemotaxis, blood-related disorders, viral infection, HIV, wound healing, cancer

Human eosinocyte CC type chemokine eotaxin	GeneSeq Accession W14990	WO9712914	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Inflammatory and immune disorders
Human thymus and activated regulated cytokine (TARC)	GeneSeq Accession W14018	WO9711969	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Inflammatory and immune disorders
Human chemokine beta-8 short forms	GeneSeq Accession W16315	WO9712041	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Cancer, wound healing, immune disorders
Microphage derived	GeneSeq Accession	WO9640923	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Inflammatory diseases, wound healing

chemokine, MDC	W20058		processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. ©Humana Press Inc., Totowa, NJ	angiogenesis
Human chemokine ZSIG-35	GeneSeq Accession W30565	WO9844117	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. ©Humana Press Inc., Totowa, NJ	Inflammatory and immune diseases
Primate CC chemokine "ILINCK"	GeneSeq Accession W69990	WO98328658	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. ©Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders, abnormal proliferation, regeneration, generation and atrophy disorders
Primate CXC chemokine "IBICK"	GeneSeq Accession W69989	WO9832858	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. ©Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders, abnormal proliferation, regeneration, generation and atrophy disorders

Human CC-type chemokine protein designated SLC (secondary lymphoid chemokine)	GeneSeq Accession W69163	WO9831809	have been described, which bind to ~17 receptors thus far identified. Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. ©Humana Press Inc., Totowa, NJ	Immune, inflammatory, and infectious disorders, cancer
Human CC chemokine ELC protein	GeneSeq Accession W62542	WO9826071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and infectious diseases, particularly herpes virus
Human DVic-1 C-C chemokine	GeneSeq Accession W60649	Wo9823750	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Abnormal proliferation, regeneration, degeneration, and atrophy disorders, including cancer

Human C-C chemokine DGWCC	GeneSeq Accession W60650	WO9823750	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, cell proliferation disorders, cancer
Human STCP-1	GeneSeq Accession W62783	WO9824907	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, particularly T cell related disorders, viral infection, and inflammation, especially joint
Exodua protein	GeneSeq Accession W61279	WO9821330	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders, angiogenesis, cancer, and proliferation disorders, particularly myeloproliferative diseases

Human Chr19Kine protein	GeneSeq Accession W50887	W09814581	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods of Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and degenerative disorders
Human T cell mixed lymphocyte reaction expressed chemokine (TMEC)	GeneSeq Accession W58703	US5780268	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods of Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune, inflammatory, and infectious disorders, cancer
Human 6CKine protein	GeneSeq Accession W50885	W09814581	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods of Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Cancer and degenerative disorders

human liver and activation regulated chemokine (LARC)	GeneSeq Accession W57475	WO9817800	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune, inflammatory, and infectious disorders, cancer
RANTES peptide	GeneSeq Accession W29538	WO9744462	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Infectious diseases, particularly HIV

RANTES 8-68	GeneSeq Accession W29529	WO9744462	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Infectious diseases, particularly HIV
RANTES 9-68	GeneSeq Accession W29528	WO9744462	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Infectious diseases, particularly HIV

Human chemokine protein 331D5	GeneSeq Accession W59433	WO9811226	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Abnormal proliferation, regeneration, degeneration or atrophy, including cancer
Human chemokine protein 611b4	GeneSeq Accession W59430	WO9811226	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Abnormal proliferation, regeneration, degeneration or atrophy, including cancer

Chemokine MCP-4	GeneSeq Accession W56690	WO9809171	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune, Inflammatory, and infectious diseases
Human stromal cell-derived chemokine, SDF-1	GeneSeq Accession W50766	FR2751658	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	HIV infections
Thymus expressed chemokine (TECK)	GeneSeq Accession W44397	WO9801557	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders

Human chemokine MIP-3alpha	GeneSeq Accession W44398	WO9801557	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders
Human chemokine MIP-3beta	GeneSeq Accession W44399	WO9801557	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders
Human monocyte chemotactic protein (MCP) sequence	GeneSeq Accession W42072	WO9802459	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, respiratory disorders, cancer

Macrophage-derived chemokine (MDC)	GeneSeq Accessions W40811 and Y24414	US5688927/ US5932703	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune, and inflammatory disorders, cancer
Macrophage derived chemokine analogue MDC-eyf	GeneSeq Accession Y24416	US5932703	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders
Macrophage derived chemokine analogue MDC (n+1)	GeneSeq Accession Y24413	US5932703	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders

Macrophage derived chemokine analogue MDC-yl	GeneSeq Accession Y24415	US5932703	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders
Human type CC chemokine eotaxin 3 protein sequence	GeneSeq Accession Y43178	JP11243960	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Allergic diseases and HIV infection
Human MCP-3 and human Muc-1 core epitope (VNT) fusion protein	GeneSeq Accession Y29893	WO9946392	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and immune disorders, particularly HIV infection

Human IP-10 and human Muc-1 core epitope (VNT) fusion protein	GeneSeq Accession Y29894	WO9946392	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and immune disorders, particularly HIV infection
Human IP-10 and HIV-1 gp 120 hypervariable region fusion protein	GeneSeq Accession Y29897	WO9946392	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and immune disorders, particularly HIV infection
Human mammary associated chemokine (MACK) protein Full-Length and Mature	GeneSeq Accessions Y29092 and Y29093	WO9936540	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Breast disease, including cancer

Tim-1 protein	GeneSeq Accession Y28290	WO99333990	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inflammation due to stimuli such as heart attacks and stroke, infection, physical trauma, UV or ionizing radiation, burns, frostbite or corrosive chemicals
Human Lkn-1 Full-Length and Mature protein	GeneSeq Accessions Y17280, Y17274, Y17281, and Y17275	WO9928473 and WO9928472	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	HIV infection and cancer, particularly leukemia
N-terminal modified chemokine met-hSDF-1 alpha	GeneSeq Accession Y05818	WO9920759	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inhibit or stimulate angiogenesis, inhibit the binding of HIV

N-terminal modified chemokine mel-hSDF-1 beta	GeneSeq Accession Y05819	WO9920759	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inhibit or stimulate angiogenesis, inhibit the binding of HIV, antiinflammatory; immunosuppressant
N-terminal modified chemokine GroHEK/hSDF-1alpha	GeneSeq Accession Y05820	WO9920759	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inhibit or stimulate angiogenesis, inhibit the binding of HIV, antiinflammatory; immunosuppressant
N-terminal modified chemokine GroHEK/hSDF-1beta	GeneSeq Accession Y05821	WO9920759	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inhibit or stimulate angiogenesis, inhibit the binding of HIV, antiinflammatory; immunosuppressant

Chemokine Eotaxin	GeneSeq Accession Y14230	WO9912968	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Increase or enhance an inflammatory response, an immune response or haematopoietic cell-associated activity; treat a vascular indication; Cancer; enhance wound healing, to prevent or treat asthma, organ transplant rejection, rheumatoid arthritis or allergy
Chemokine hMCP1a	GeneSeq Accession Y14225	WO9912968	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, Vascular disorders, Wound healing, cancer, prevent organ transplant rejection, increase or enhance an inflammatory response,
Chemokine hMCP1b	GeneSeq Accession Y14226	WO9912968	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, Vascular disorders, Wound healing, cancer, prevent organ transplant rejection, increase or enhance an inflammatory response,

Chemokine hSDF1b	GeneSeq Accession Y14228	WO9912968	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, Vascular disorders, Wound healing, cancer, prevent organ transplant rejection, Increase or enhance an inflammatory response,
Chemokine hIL-8	GeneSeq Accession Y14229	WO9912968	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ; and Holmes et al (1991) Science 253, 1278-80.	Immune disorders, Vascular disorders, Wound healing, cancer, prevent organ transplant rejection, Increase or enhance an inflammatory response,
Chemokine hMCP1	GeneSeq Accession Y14222	WO9912968	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, Vascular disorders, Wound healing, cancer, prevent organ transplant rejection, Increase or enhance an inflammatory response,

Chemokine hMCP2	GeneSeq Accession Y14223	WO9912968	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, Vascular disorders, Wound healing, cancer, prevent organ transplant rejection, increase or enhance an inflammatory response.
Chemokine hMCP3	GeneSeq Accession Y14224	WO9912968	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, Vascular disorders, Wound healing, cancer, prevent organ transplant rejection, increase or enhance an inflammatory response.
C-C chemokine, MCP2	GeneSeq Accession Y05300	EP905240	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inflammatory, Immune and infectious diseases; pulmonary diseases and skin disorders; tumours, and angiogenesis-related haematopoiesis-related diseases

Wild type monocyte chemotactic protein 2	GeneSeq Accession Y07233	EP906954	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Inflammatory, immune and infectious diseases; pulmonary diseases and skin disorders; tumours, and angiogenesis-and haematopoiesis-related diseases
Truncated monocyte chemotactic protein 2 (6-76)	GeneSeq Accession Y07234	EP906954	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. Humana Press Inc., Totowa, NJ	Inflammatory, immune and infectious diseases; pulmonary diseases and skin disorders; tumours, and angiogenesis-and haematopoiesis-related diseases
Truncated RANTES protein (3-68)	GeneSeq Accessions Y07236 and Y07232	EP905241; EP906954	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. Humana Press Inc., Totowa, NJ	Inflammatory, immune and infectious diseases; pulmonary diseases and skin disorders; tumours, and angiogenesis-and haematopoiesis-related diseases

Wild type monocyte chemotactic protein 2	GeneSeq Accession Y07237	EP905241	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, Humana Press Inc., Totowa, NJ	Inflammatory, immune and infectious diseases; pulmonary diseases and skin disorders; tumours, and angiogenesis-and haematopoiesis-related diseases
Truncated monocyte chemotactic protein 2 (6-76)	GeneSeq Accession Y07238	EP905241	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, Humana Press Inc., Totowa, NJ	Inflammatory, immune and infectious diseases; pulmonary diseases and skin disorders; tumours, and angiogenesis-and haematopoiesis-related diseases
A partial CXCR4B protein	GeneSeq Accession W97363	EP897980	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, Humana Press Inc., Totowa, NJ	Soluble CXCR4B receptor polypeptides may be useful for inhibiting chemokine activities and viral infection.

Interferon gamma-inducible protein (IP-10)	GeneSeq Accession W96709	US5871723	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders
A monokine induced by gamma-interferon (MIG)	GeneSeq Accession W96710	US5871723	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders
Interleukin-8 (IL-8) protein.	GeneSeq Accession W96711	US5871723	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, Humana Press Inc., Totowa, NJ; and Holmes et al (1991) Science 253, 1278-80.	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders

Epithelial neutrophil activating protein-78 (ENA-78)	GeneSeq Accession W96712	US5871723	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders
Growth related oncogene-alpha (GRO-alpha).	GeneSeq Accession W96713	US5871723	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders
Growth related oncogene-beta (GRO-beta).	GeneSeq Accession W96714	US5871723	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders

Growth related oncogene-gamma (GRO-gamma)	GeneSeq Accession W96715	US5871723	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders
A platelet basic protein (PBP)	GeneSeq Accession W96716	US5871723	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders
Connective tissue activating protein- III (CTAP-III)	GeneSeqAccession S96717	US5871723	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders

Beta-thromboglobulin protein (beta-TG)	GeneSeq Accession W96718	US5871723	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders
Neutrophil activating peptide-2 (NAP-2)	GeneSeq Accession W96719	US5871723	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders
Granulocyte chemotactic protein-2 (GCP-2)	GeneSeq Accession W96720	US5871723	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Angiogenesis, Cancer, Inflammatory and Immune disorders, Cardio-Vascular disorders, Musco-skeletal disorders

Human chemokine MIG-beta protein	GeneSeq Accession W90124	EP887409	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Immune disorders, viral, parasitic, fungal or bacterial infections, Cancer; autoimmune diseases or transplant rejection
Human ZCHEMO-8	GeneSeq Accession W82716	WO9854326	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Immune disorders, cancer, myelopietic disorders, autoimmune disorders, immunodeficiencies, Inflammatory and infectious diseases, Vascular disorders, wound healing
Human Act-2 protein	GeneSeq Accession W82717	WO9854326	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power, © Humana Press Inc., Totowa, NJ	Immune disorders, cancer, myelopietic disorders, autoimmune disorders, immunodeficiencies, Inflammatory and infectious diseases, Vascular disorders, wound healing

Human S1SD protein	GeneSeq Accession W82720	WO9854326	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune disorders, cancer, myelopietic disorders, autoimmune disorders and immunodeficiencies, Inflammatory and infectious diseases, Vascular disorders, wound healing
Human M110 protein	GeneSeq Accession W82721	WO9854326	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune disorders, cancer, myelopietic disorders, autoimmune disorders and immunodeficiencies, Inflammatory and infectious diseases, Vascular disorders, wound healing
Human M11A protein	GeneSeq Accession W82722	WO9854326	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols, Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune disorders, cancer, myelopietic disorders, autoimmune disorders and immunodeficiencies, Inflammatory and infectious diseases, Vascular disorders, wound healing

Human CCC3 protein	GeneSeq Accession W82723	WO9854326	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokines activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Immune disorders, cancer, myelopietic disorders, autoimmune disorders and immunodeficiencies, inflammatory and infectious diseases, vascular disorders, wound healing
A human L105 chemokine designated hul105_3.	GeneSeq Accession W87588	WO9856818	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Cancer, wound healing

A human L105 chemokine designated huL105_7.	GeneSeq Accession W87589	WO9856818	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Cancer, wound healing
Human mature gro-alpha polypeptide used to treat sepsis	GeneSeq Accession W81498	WO9848828	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Infectious diseases, sepsis

Human mature gro-gamma polypeptide used to treat sepsis	GeneSeq Accession W81500	WO9848828	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G- protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Infectious diseases, sepsis
Human thymus expressed chemokine TECK and TECK variant	GeneSeq Accessions B19607 and B19608	WO0053635	Chemokines area family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G- protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Mehtods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Inflammatory disorders, cancer, immune and vascular disorders

Human chemokine SDF1alpha	GeneSeq Accession B15791	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods of Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine GROalpha	GeneSeq Accession B15793	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine eotaxin	GeneSeq Accession B15794	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders

Human chemokine MIG	GeneSeq Accession B15803	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine PF4	GeneSeq Accession B15804	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine I-309	GeneSeq Accession B15805	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders

Human chemokine HCC-1	GeneSeq Accession B15806	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine C10	GeneSeq Accession B15807	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine CCR-2	GeneSeq Accession B15808	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders

Human chemokine ENA-78	GeneSeq Accession B15809	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine GRObeta	GeneSeq Accession B15810	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine IP-10	GeneSeq Accession B15811	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders

Human chemokine SDF1beta	GeneSeq Accession B15812	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine GRO alpha	GeneSeq Accession B15813	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders
Human chemokine MIP1beta	GeneSeq Accession B15831	WO0042071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Autoimmune disorders, Immune, Vascular and Inflammatory disorders

A human C-C chemokine designated exodus	GeneSeq Accession B07939	US6096300	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer
Human chemokine L105_7	GeneSeq Accession Y96922	US6084071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Chemotaxis, Gene Therapy, Wound healing
Human chemokine L105_3	GeneSeq Accession Y96923	US6084071	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Chemotaxis, Gene Therapy, Wound healing

Human secondary lymphoid chemokine (SLC)	GeneSeq Accession B01434	WO0038706	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer, Vascular and Immune disorders
Human non-ELR CXC chemokine H174	GeneSeq Accession Y96310	WO0029439	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and Inflammatory disorders, Cancer, Haemostatic and thrombolytic activity
Human non-ELR CXC chemokine IP10	GeneSeq Accession Y96311	WO0029439	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~ 17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and Inflammatory disorders, Cancer, haemostatic and thrombolytic activity

Human non-ELR CXC chemokine Mig	GeneSeq Accession Y96313	WO0029439	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Immune and inflammatory disorders, Cancer, haemostatic and thrombolytic activity
Human chemokine Ckbeta-7	GeneSeq Accession Y96280	WO0028035	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer, wound healing, inflammatory and immunoregulatory disorders
Human chemokine MIP-1alpha	GeneSeq Accession Y96281	WO0028035	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer, wound healing, inflammatory and immunoregulatory disorders

Human mature chemokine Ckbeta-7 (optionally truncated)	GenSeq Accession Y96282	WO0028035	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer, wound healing, inflammatory and immunoregulatory disorders
Human chemokine receptor CXCR3	GeneSeq Accession Y79372	WO0018431	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Soluble CXCR3 polypeptides may be useful for inhibiting chemokine activities and viral infection.
Human neurotactin chemokine like domain	GeneSeq Accession Y53259	US6043086	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified.	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Neurological disorders, Immune and respiratory disorders

Human CC type chemokine interleukin C	GeneSeq Accession Y57771	JP11302298	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer and infectious diseases
Human CKbeta-9	GeneSeq Accession B50860	US6153441	Chemokines are a family of related small, secreted proteins involved in biological processes ranging from hematopoiesis, angiogenesis, and leukocyte trafficking. Members of this family are involved in a similarly diverse range of pathologies including inflammation, allergy, tissue rejection, viral infection, and tumor biology. The chemokines exert their effects by acting on a family of seven transmembrane G-protein-coupled receptors. Over 40 human chemokines have been described, which bind to ~17 receptors thus far identified	Chemokine activities can be determined using assays known in the art: Methods in Molecular Biology, 2000, vol. 138: Chemokine Protocols. Edited by: A.E.I. Proudfoot, T.N.C. Wells, and C.A. Power. © Humana Press Inc., Totowa, NJ	Cancer, Auto-immune and inflammatory disorders, Cardiovascular disorders
Preproapolipoprotein "paris" variant	GeneSeq Accession W08602	WO9637608	Apoa-1 participates in the reverse transport of cholesterol from tissues to the liver for excretion by promoting cholesterol efflux from tissues and by acting as a cofactor for the lecithin cholesterol acyltransferase (lcatal).	Lipid binding activity can be determined using assays known in the art, such as, for example, the Cholesterol Efflux Assays of Takahashi et al., P.N.A.S., Vol. 96, Issue 20, 11358-11363, September 28, 1999.	Useful for cardiovascular disorders, cholesterol disorders, and Hyperlipidaemia
Preproapolipoprotein "milano" variant		5,721,114	Apoa-1 participates in the reverse transport of cholesterol from tissues to the liver for excretion by promoting cholesterol efflux from tissues and by acting as a cofactor for the lecithin cholesterol acyltransferase (lcatal).	Lipid binding activity can be determined using assays known in the art, such as, for example, the Cholesterol Efflux Assays of Takahashi et al., P.N.A.S., Vol. 96, Issue 20, 11358-11363, September 28, 1999.	Useful for cardiovascular disorders, cholesterol disorders, and Hyperlipidaemia

Glycodelin-A; Progesterone- associated endometrial protein	GeneSeq Accession W00289	WO9628169	Naturally produced female contraceptive that is removed rapidly from the body following 2-3 days production. Uses include contraception	Glycodelin-A activity can be determined using the hemizona assay as described in Oehninger, S., Coddington, C.C., Hodgen, G.D., and Seppala, M (1995) Fertil. Steril. 63, 377-383.	Naturally derived contraceptive useful for the prevention of pregnancy.
NOGO-A	Genbank Accession CAB99248		NOGO polypeptides are potent inhibitors of neurite growth.	Inhibition of Neurite outgrowth. Antagonists to NOGO polypeptides may promote the outgrowth of neurites, thus inducing regeneration of neurons.	NOGO-A polypeptide antagonists are useful for the promotion of neural growth, which could be useful in the treatment of neural disorders and dysfunction due to degenerative diseases or trauma; useful in the treatment of neoplastic diseases of the CNS; induce regeneration of neurons or to promote the structural plasticity of the CNS.
NOGO-B	Genbank Accession CAB99249		NOGO polypeptides are potent inhibitors of neurite growth.	Inhibition of Neurite outgrowth. Antagonists to NOGO polypeptides may promote the outgrowth of neurites, thus inducing regeneration of neurons.	NOGO-B polypeptide antagonists are useful for the promotion of neural growth, which could be useful in the treatment of neural disorders and dysfunction due to degenerative diseases or trauma; useful in the treatment of neoplastic diseases of the CNS; induce regeneration of neurons or to promote the structural plasticity of the CNS.

NOGO-C	Genbank Accession CAB99250		NOGO polypeptides are potent inhibitors of neurite growth.	Inhibition of Neurite outgrowth. Antagonists to NOGO polypeptides may promote the outgrowth of neurites, thus inducing regeneration of neurons.	NOGO-C polypeptide antagonists are useful for the promotion of neural growth, which could be useful in the treatment of neural disorders and dysfunction due to degenerative diseases or trauma; useful in the treatment of neoplastic diseases of the CNS; induce regeneration of neurons or to promote the structural plasticity of the CNS.
NOGO-66 Receptor	Genbank Accession AAG53612		NOGO polypeptides are potent inhibitors of neurite growth, and are thought to mediate their effects through the NOGO-66 Receptor.	Inhibition of Neurite outgrowth by mediating the biological effects of NOGO polypeptides. Soluble NOGO-66 receptor polypeptides may promote the outgrowth of neurites, thus inducing regeneration of neurons.	NOGO-66 receptor polypeptides are useful for the promotion of neural growth, which could be useful in the treatment of neural disorders and dysfunction due to degenerative diseases or trauma; useful in the treatment of neoplastic diseases of the CNS; induce regeneration of neurons or to promote the structural plasticity of the CNS.
Antibodies specific for collapsin		US5416197	These antibodies are useful for the promotion of neurite outgrowth	Collapsin activity, which is thought to inhibit the outgrowth of neurites, can be assayed in the presence of antibodies specific for collapsing using assays known in the art, such as, for example, the collapse assay disclosed by Luo et al., Cell 1993 Oct 22;75(2):217-27	Useful for the promotion of neural growth, which could be useful in the treatment of neural disorders and dysfunction due to degenerative diseases or trauma.
Humanized Anti-VEGF Antibodies, and fragments thereof		WO9845331	These agents have anti-inflammatory and anti-cancer applications	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer

Humanized Anti-VEGF Antibodies, and fragments thereof	WO0029584	These agents have anti-inflammatory and anti-cancer applications	VEGF activity can be determined using assays known in the art, such as those disclosed in International Publication No. WO0045835, for example.	Promotion of growth and proliferation of cells, such as vascular endothelial cells. Antagonists may be useful as anti-angiogenic agents, and may be applicable for cancer
Membrane bound proteins	GeneSeq. Accession Y66631-Y66765	Cancer, Immune Disorders	These proteins can be used for linking bioactive molecules to cells and for modulating biological activities of cells, using the polypeptides for specific targeting. The polypeptide targeting can be used to kill the target cells, e.g. for the treatment of cancers. These proteins are useful for the treatment of immune system disorders.	Activities can be determined using assay known in the art, such as, for example, the assays disclosed in International Publication No. WO0121658.
Secreted and Transmembrane polypeptides	GeneSeq Accession B44241-B44334	Cancer, Immune Disorders	These proteins can be used for linking bioactive molecules to cells and for modulating biological activities of cells, using the polypeptides for specific targeting. The polypeptide targeting can be used to kill the target cells, e.g. for the treatment of cancers. These proteins are useful for the treatment of immune system disorders.	Activities can be determined using assay known in the art, such as, for example, the assays disclosed in International Publication No. WO0121658
Secreted and Transmembrane polypeptides	GeneSeq Accession Y41685-Y41774	Cancer, Immune Disorders	These proteins can be used for linking bioactive molecules to cells and for modulating biological activities of cells, using the polypeptides for specific targeting. The polypeptide targeting can be used to kill the target cells, e.g. for the treatment of cancers. These proteins are useful for the treatment of immune system disorders.	Activities can be determined using assay known in the art, such as, for example, the assays disclosed in International Publication No. WO0121658

Conjugation and Coupling

[0123] The present invention provides therapeutic agents comprising an ELP component and a therapeutic component, such as therapeutic proteins listed in Table 1, as well as a GLP-1 receptor agonists, insulin, Factor VII/VIIa, and functional analogs as described. Such agents may be prepared by recombinant technology and/or chemical coupling (e.g., conjugation).

[0124] A recombinantly-produced ELP fusion protein, in accordance with certain embodiments of the invention, includes the ELP component and the therapeutic component associated with one another by genetic fusion. For example, the fusion protein may be generated by translation of a polynucleotide encoding the therapeutic component cloned in-frame with the ELP component (or vice versa). Such an ELP fusion protein may contain one or more copies of the therapeutic component attached to the N-terminus and/or the C-terminus of the ELP component. In some embodiments, the therapeutic proteinaceous component is attached to both the N- and C-terminus of the ELP component and the fusion protein may contain one or more equivalents of the therapeutic component on either or both ends of the ELP component.

[0125] In certain embodiments, the ELP component and the therapeutic components can be fused using a linker peptide of various lengths to provide greater physical separation and allow more spatial mobility between the fused portions, and thus maximize the accessibility of the therapeutic component, 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 flexible linker may include amino acids having relatively small side chains, and which may be hydrophilic. Without limitation, the flexible linker may contain a stretch of glycine and/or serine residues. More rigid linkers may contain, for example, more sterically hindering amino acid side chains, such as (without limitation) tyrosine or histidine. The linker may be less than about 50, 40, 30, 20, 10, or 5 amino acid residues. The linker can be covalently linked to and between an ELP component and a therapeutic component, for example, via recombinant fusion.

[0126] The linker or peptide spacer may be protease-cleavable or non-cleavable. By way of example, cleavable peptide spacers include, without limitation, a peptide sequence recognized by proteases (*in vitro* or *in vivo*) of varying type, such as Tev, thrombin, factor Xa, plasmin (blood proteases), metalloproteases, cathepsins (e.g., GFLG, etc.), and proteases found in other corporeal compartments. In some embodiments employing cleavable linkers, the fusion protein ("the therapeutic agent") may be inactive, less active, or less potent as a fusion, which is then activated upon cleavage of the spacer *in vivo*. Alternatively, where the therapeutic agent is sufficiently active as a fusion, a non-cleavable spacer may be employed. The non-cleavable spacer may be of any suitable type, including, for example, non-cleavable spacer moieties having the formula [(Gly)*n*-Ser]*m* (SEQ ID NO.: 22) where *n* is from 1 to 4, inclusive, and *m* is from 1 to 4, inclusive. Alternatively, a short ELP sequence different than the backbone ELP could be employed instead of a linker or spacer, while accomplishing the necessary effect.

[0127] In still other embodiments, the therapeutic agent is a recombinant fusion having a therapeutic component flanked on each terminus by an ELP component. At least one of said ELP components may be attached via a cleavable spacer, such that the therapeutic component is inactive, but activated *in vivo* by proteolytic removal of a single ELP component. The resulting single ELP fusion being active, and having an enhanced half-life (or other property described herein) *in vivo*.

[0128] In other embodiments, the present invention provides chemical conjugates of the ELP component and the therapeutic component. The conjugates can be made by chemically coupling an ELP component to a therapeutic component by any number of methods well known in the art (See e.g. Nilsson et al., 2005, Ann Rev Biophys Bio Structure 34: 91-118). In some embodiments, the chemical conjugate can be formed by covalently linking the therapeutic component to the ELP component, directly or through a short or long linker moiety, through one or more functional groups on the therapeutic proteinaceous component, e.g., amine, carboxyl, phenyl, thiol or hydroxyl groups, to form a covalent conjugate. Various conventional linkers can be used, e.g., diisocyanates, diisothiocyanates, carbodiimides, bis (hydroxysuccinimide) esters, maleimide- hydroxysuccinimide esters, glutaraldehyde and the like.

[0129] Non-peptide chemical spacers can additionally be of any suitable type, including for example, by functional linkers described in Bioconjugate Techniques, Greg T. Hermanson, published by Academic Press, Inc., 1995, and those specified in the CrossLinking Reagents Technical Handbook, available from Pierce Biotechnology, Inc. (Rockford, Illinois), the disclosures of which are hereby incorporated by reference, in their respective entireties. Illustrative chemical spacers include homobifunctional linkers that can attach to amine groups of Lys, as well as heterobifunctional linkers that can attach to Cys at one terminus, and to Lys at the other terminus.

[0130] In certain embodiments, relatively small ELP components (e.g., ELP components of less than about 30 kDa, 25 kDa, 20 kDa, 15 kDa, or 10 kDa), that do not transition at room temperature (or human body temperature, e.g., T_t >37°C), are chemically coupled or crosslinked. For example, two relatively small ELP components, having the same or different properties, may be chemically coupled. Such coupling, in some embodiments, may take place *in vivo*, by the addition of a single cysteine residue at or around the C-terminus of the ELP. Such ELP components may each be fused to one or more therapeutic components, so as to increase activity or avidity at the target.

Polynucleotides, Vectors, and Host Cells

[0131] In another aspect, the invention provides polynucleotides comprising a nucleotide sequence encoding the therapeutic agent of the invention. Such polynucleotides further comprise, in addition to sequences encoding the ELP and therapeutic components, one or more expression control elements. For example, the polynucleotide, may comprise one or more promoters or transcriptional enhancers, ribosomal binding sites, transcription termination signals, and polyadenylation signals, as expression control elements. The polynucleotide may be inserted within any suitable vector, which may be contained within any suitable host cell for expression.

[0132] A vector comprising the polynucleotide can be introduced into a cell for expression of the therapeutic agent. The vector can remain episomal or become chromosomally integrated, as long as the insert encoding the therapeutic agent can be transcribed. Vectors can be constructed by standard recombinant DNA technology. Vectors can be plasmids, phages, cosmids, phagemids, viruses, or any other types known in the art, which are used for replication and expression in prokaryotic or eukaryotic cells. It will be appreciated by one of skill in the art that a wide variety of components known in the art (such as expression control elements) may be included in such vectors, including a wide variety of transcription signals, such as promoters and other sequences that regulate the binding of RNA polymerase onto the promoter. Any promoter known to be effective in the cells in which the vector will be expressed can be used to initiate expression of the therapeutic agent. Suitable promoters may be inducible or constitutive. Examples of suitable promoters include the SV40 early promoter region, the promoter contained in the 3' long terminal repeat of Rous sarcoma virus, the HSV-1 (herpes simplex virus-1) thymidine kinase promoter, the regulatory sequences of the metallothionein gene, etc., as well as the following animal transcriptional control regions, which exhibit tissue specificity and have been utilized in transgenic animals: elastase I gene control region which is active in pancreatic acinar cells; insulin gene control region which is active in pancreatic beta cells, immunoglobulin gene control region which is active in lymphoid cells, mouse mammary tumor virus control region which is active in testicular, breast, lymphoid and mast cells, albumin gene control region which is active in liver, alpha-fetoprotein gene control region which is active in liver, alpha 1 -antitrypsin gene control region which is active in the liver, beta-globin gene control region which is active in erythroid cells, myelin basic protein gene control region which is active in oligodendrocyte cells in the brain, myosin light chain-2 gene control region which is active in skeletal muscle, and gonadotropin releasing hormone gene control region which is active in the hypothalamus.

Pharmaceutical Compositions

[0133] The present invention further provides pharmaceutical compositions comprising the therapeutic agents of the invention (as described above) together with a pharmaceutically acceptable carrier or excipient. Such pharmaceutical compositions may be employed in the methods of treatment as described above, for each of the therapeutic proteins, e.g., the therapeutic proteins listed in Table 1, GLP-1 receptor agonists, insulin, and Factor VII/VIIa embodiments.

[0134] The therapeutic agents of the invention may overcome certain deficiencies of peptide agents when administered (e.g., parenterally), including in some embodiments, the limitation that such peptides may be easily metabolized by plasma proteases or cleared from circulation by kidney filtration. Traditionally, the oral route of administration of peptide agents may also be problematic, because in addition to proteolysis in the stomach, the high acidity of the stomach destroys such peptide agents before they reach their intended target tissue. Peptides and peptide fragments produced by the action of gastric and pancreatic enzymes are cleaved by exo and endopeptidases in the intestinal brush border membrane to yield di- and tripeptides, and even if proteolysis by pancreatic enzymes is avoided, polypeptides are subject to degradation by brush border peptidases. Any of the peptide agents that survive passage through the stomach are further subjected to metabolism in the intestinal mucosa where a penetration barrier prevents entry into the cells. In certain embodiments, the therapeutic agents of the invention may overcome such deficiencies, and provide compositional forms having enhanced efficacy, bioavailability, therapeutic half-life, persistence, degradation assistance, etc. The therapeutic agents of the invention thus include oral and parenteral dose forms, as well as various other dose forms, by which peptide agents can be utilized in a highly effective manner. For example, in some embodiments, such agents may achieve high mucosal absorption, and the concomitant ability to use lower doses to elicit an optimum therapeutic effect.

[0135] The therapeutic agents of the present invention may be administered in smaller doses and/or less frequently than unfused or unconjugated counterparts. While one of skill in the art can determine the desirable dose in each case, a suitable dose of the therapeutic agent for achievement of therapeutic benefit, may, for example, be in a range of about 1 microgram (μg) to about 100 milligrams (mg) per kilogram body weight of the recipient per day, preferably in a range of about 10 μg to about 50 mg per kilogram body weight per day and most preferably in a range of about 10 μg to about 50 mg per kilogram body weight per day. The desired dose may be presented as one dose or two or more sub-doses administered at appropriate intervals throughout the day. These sub-doses can be administered in unit dosage forms, for example, containing from about 10 μg to about 1000 mg, preferably from about 50 μg to about 500 mg, and most preferably from about 50 μg to about 250 mg of active ingredient per unit dosage form. Alternatively, if the condition of the recipient so requires, the doses may be administered as a continuous infusion.

[0136] The mode of administration and dosage forms will of course affect the therapeutic amount of the peptide active therapeutic agent that is desirable and efficacious for a given treatment application. For example, orally administered dosages can be at least twice, e.g., 2-10 times, the dosage levels used in parenteral administration methods.

[0137] The therapeutic agents of the invention may be administered per se as well as in various forms including pharmaceutically acceptable esters, salts, and other physiologically functional derivatives thereof. The present invention also contemplates pharmaceutical formulations, both for veterinary and for human medical use, which include therapeutic agents of the invention. In such pharmaceutical and medicament formulations, the therapeutic agents can be used together with one or more pharmaceutically acceptable carrier(s) therefore and optionally any other therapeutic ingredients. The carrier(s) must be pharmaceutically acceptable in the sense of being compatible with the other ingredients of the formulation and not unduly deleterious to the recipient thereof. The therapeutic agents are provided in an amount effective to achieve the desired pharmacological effect, as described above, and in a quantity appropriate to achieve the desired daily dose.

[0138] The formulations of the therapeutic agent include those suitable for parenteral as well as non-parenteral administration, and specific administration modalities include oral, rectal, buccal, topical, nasal, ophthalmic, subcutaneous, intramuscular, intravenous, transdermal, intrathecal, intra-articular, intra-arterial, sub-arachnoid, bronchial, lymphatic, vaginal, and intra-uterine administration. Formulations suitable for oral and parenteral administration are preferred.

[0139] When the therapeutic agent is used in a formulation including a liquid solution, the formulation advantageously can be administered orally or parenterally. When the therapeutic agent is employed in a liquid suspension formulation or as a powder in a biocompatible carrier formulation, the formulation may be advantageously administered orally, rectally, or bronchially.

[0140] When the therapeutic agent is used directly in the form of a powdered solid, the active agent can be advantageously administered orally. Alternatively, it may be administered bronchially, via nebulization of the powder in a carrier gas, to form a gaseous dispersion of the powder which is inspired by the patient from a breathing circuit comprising a suitable nebulizer device.

[0141] The formulations comprising the therapeutic agent of the present invention may conveniently be presented in unit dosage forms and may be prepared by any of the methods well known in the art of pharmacy. Such methods generally include the step of bringing the therapeutic agents into association with a carrier which constitutes one or more accessory ingredients. Typically, the formulations are prepared by uniformly and intimately bringing the therapeutic agent into association with a liquid carrier, a finely divided solid carrier, or both, and then, if necessary, shaping the product into dosage forms of the desired formulation.

[0142] Formulations suitable for oral administration may be presented as discrete units such as capsules, cachets, tablets, or lozenges, each containing a predetermined amount of the active ingredient as a powder or granules; or a suspension in an aqueous liquor or a non-aqueous liquid, such as a syrup, an elixir, an emulsion, or a draught.

[0143] A tablet may be made by compression or molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine, with the therapeutic agent being in a free-flowing form such as a powder or granules which optionally is mixed with a binder, disintegrant, lubricant, inert diluent, surface active agent, or discharging agent. Molded tablets comprised of a mixture of the powdered peptide active therapeutic agent-ELP construct(s) with a suitable carrier may be made by molding in a suitable machine.

[0144] A syrup may be made by adding the peptide active therapeutic agent-ELP construct(s) to a concentrated aqueous solution of a sugar, for example sucrose, to which may also be added any accessory ingredient(s). Such accessory ingredient(s) may include flavorings, suitable preservative, agents to retard crystallization of the sugar, and agents to increase the solubility of any other ingredient, such as a polyhydroxy alcohol, for example glycerol or sorbitol.

[0145] Formulations suitable for parenteral administration conveniently comprise a sterile aqueous preparation of the therapeutic agent, which preferably is isotonic with the blood of the recipient (e.g., physiological saline solution). Such formulations may include suspending agents and thickening agents or other microparticulate systems which are designed to target the peptide active therapeutic agent to blood components or one or more organs. The formulations may be presented in unit-dose or multi-dose form.

[0146] Nasal spray formulations comprise purified aqueous solutions of the therapeutic agent with preservative agents and isotonic agents. Such formulations are preferably adjusted to a pH and isotonic state compatible with the nasal mucus membranes.

[0147] Formulations for rectal administration may be presented as a suppository with a suitable carrier such as cocoa butter, hydrogenated fats, or hydrogenated fatty carboxylic acid.

[0148] - Topical formulations comprise the therapeutic agent dissolved or suspended in one or more media, such as mineral oil, petroleum, polyhydroxy alcohols, or other bases used for topical pharmaceutical formulations.

[0149] In addition to the aforementioned ingredients, the formulations of this invention may further include one or more accessory ingredient(s) selected from diluents, buffers, flavoring agents, disintegrants, surface active agents, thickeners, lubricants, preservatives (including antioxidants), and the like.

[0150] The features and advantages of the present invention are more fully shown with respect to the following non-

limiting examples.

EXAMPLES

EXAMPLE 1: CONSTRUCTION OF VARIOUS ELP COMPONENT CONSTRUCTS

[0151] Cloning steps were conducted in *Escherichia coli* strain XL1-Blue (rec A1, *endA1*, *gyrA96*, *thi-1*, *hsdR17* (r_k^- , m_k^+), *supE44*, *relA1*, *lac[F']*, *proAB*, *lacI^qΔM15*, Tn10 (Tet^r)) (Stratagene La Jolla, CA). pUC19 (NEB, Beverly, MA) was used as the cloning vector for the ELP construction (Meyer and Chilkoti, Nat. Biotechnol., 17(11):1112-5, 1999). Modified forms of pET15b and pET24d vectors (Novagen) were used to express ELP and ELP-fusion proteins in BL21 Star (DE3) strain (*F⁻*, *ompT*, *hsdS_B* ($r_B^- m_B^-$), *gal*, *dcm*, *rne131*, (DE3)) (Invitrogen Carlsbad, CA) or BLR(DE3) (*F⁻*, *ompT*, *hsdS_B* ($r_B^- m_B^-$), *gal*, *dcm*, $\Delta(srl-recA)$ 306::Tn10(TcR)(DE3)) (Novagen Madison, WI). Synthetic DNA oligos were purchased from Integrated DNA Technologies, Coralville, IA. All vector constructs were made using standard molecular biology protocols (e.g., Current Protocols in Molecular Biology, ed. Ausubel, et al., 1995).

Construction of ELP1 [V₅A₂G₃] Gene Series

[0152] The ELP1 [V₅A₂G₃] series designate polypeptides containing multiple repeating units of the pentapeptide VPGXG (SEQ ID NO: 3), where X is valine, alanine, and glycine at a relative ratio of 5:2:3.

[0153] The ELP1 [V₅A₂G₃] series monomer, ELP1 [V₅A₂G₃-10], was created by annealing four 5' phosphorylated, PAGE purified synthetic oligos to form double stranded DNA with *EcoRI* and *HindIII* compatible ends (Meyer and Chilkoti, Nat. Biotechnol., 17(11):1112-5, 1999). The oligos were annealed in a 1 μ M mixture of the four oligos in 50 μ l IX ligase buffer (Invitrogen) to 95°C in a heating block then the block was allowed to cool slowly to room temperature. The ELP1 [V₅A₂G₃-10]/*EcoRI-HindIII* DNA segment was ligated into a pUC19 vector digested with *EcoRI* and *HindIII* and CIAP dephosphorylated (Invitrogen) to form pUC19-ELP1 [V₅A₂G₃-10]. Building of the ELP1 [V₅A₂G₃] series library began by inserting ELP1 [V₅A₂G₃-10] *Pf*MI/*Bgl*I fragment from pUC19-ELP1 [V₅A₂G₃-10] into pUC19-ELP1 [V₅A₂G₃-10] linearized with *Pf*MI and dephosphorylated with CIAP to create pUC19-ELP1 [V₅A₂G₃-20]. pUC19-ELP1 [V₅A₂G₃-20] was then built up to pUC19-ELP1 [V₅A₂G₃-30] and pUC19-ELP1 [V₅A₂G₃-40] by ligating ELP1 [V₅A₂G₃-10] or ELP1 [V₅A₂G₃-20] *Pf*MI/*Bgl*I fragments respectively into *Pf*MI digested pUC 19-ELP1 [V₅A₂G₃-20]. This procedure was used to expand the ELP1 [V₅A₂G₃] series to create pUC19-ELP1 [V₅A₂G₃-60], pUC19-ELP1 [V₅A₂G₃-90] and pUC19-ELP1 [V₅A₂G₃-180] genes.

Construction of ELP1 [K₁V₂F₁] Gene Series

[0154] The ELP1 [K₁V₂F₁] series designate polypeptides containing multiple repeating units of the pentapeptide VPGXG (SEQ ID NO: 3), where X is lysine, valine, and phenylalanine at a relative ratio of 1:2:1.

[0155] The ELP1 [K₁V₂F₁] series monomer, ELP1 [K₁V₂F₁-4], was created by annealing two 5' phosphorylated, PAGE purified synthetic oligos to form double stranded DNA with *EcoRI* and *HindIII* compatible ends (Meyer and Chilkoti, 1999). The oligos were annealed in a 1 μ M mixture of the four oligos in 50 μ l 1X ligase buffer (Invitrogen) to 95°C in a heating block then the block was allowed to cool slowly to room temperature. The ELP1 [K₁V₂F₁-4]/*EcoRI-HindIII* DNA segment was ligated into a pUC19 vector digested with *EcoRI* and *HindIII* and CIAP dephosphorylated (Invitrogen) to form pUC19-ELP1 [K₁V₂F₁-4]. Building of the ELP1 [K₁V₂F₁] series library began by inserting ELP1 [K₁V₂F₁-4] *Pf*MI/*Bgl*I fragment from pUC19-ELP1 [K₁V₂F₁-4] into pUC19-ELP1 [K₁V₂F₁-4] linearized with *Pf*MI and dephosphorylated with CIAP to create pUC19-ELP1 [K₁V₂F₁-8]. Using the same procedure the ELP1 [K₁V₂F₁] series was doubled at each ligation to form pUC19-ELP1 [K₁V₂F₁-16], pUC19-ELP1 [K₁V₂F₁-32], pUC19-ELP1 [K₁V₂F₁-64] and pUC19-ELP1 [K₁V₂F₁-128].

Construction of ELP1 [K₁V₇F₁] Gene Series

[0156] The ELP1 [K₁V₇F₁] series designate polypeptides containing multiple repeating units of the pentapeptide VPGXG (SEQ ID NO: 3), where X is lysine, valine, and phenylalanine at a relative ratio of 1:7:1.

[0157] The ELP1 [K₁V₇F₁] series monomer, ELP1 [K₁V₇F₁-9], was created by annealing four 5' phosphorylated, PAGE purified synthetic oligos to form double stranded DNA with *Pf*MI and *HindIII* compatible ends. The ELP1 [K₁V₇F₁-9] DNA segment was then ligated into *Pf*MI/*HindIII* dephosphorylated PUC19-ELP1 [V₅A₂G₃-180] vector thereby substituting ELP1 [V₅A₂G₃-180] for ELP1 [K₁V₇F₁-9] to create the pUC19-ELP1 [K₁V₇F₁-9] monomer. The ELP1 [K₁V₇F₁] series was expanded in the same manner as the ELP1 [K₁V₂F₁] series to create pUC19-ELP1 [K₁V₇F₁-18], pUC19-ELP1 [K₁V₇F₁-36], pUC19-ELP1 [K₁V₇F₁-72] and pUC19-ELP1 [K₁V₇F₁-144].

Construction of ELP1 [V] Gene Series

[0158] The ELP1 [V] series designate polypeptides containing multiple repeating units of the pentapeptide VPGXG (SEQ ID NO: 3), where X is exclusively valine.

[0159] The ELP1 [V] series monomer, ELP1 [V-5], was created by annealing two 5' phosphorylated, PAGE purified synthetic oligos to form double stranded DNA with *EcoRI* and *HindIII* compatible ends. The ELP1 [V-5] DNA segment was then ligated into *EcoRI/HindIII* dephosphorylated pUC19 vector to create the pUC19-ELP1 [V-5] monomer. The ELP1 [V] series was created in the same manner as the ELP1 [$V_5A_2G_3$] series, ultimately expanding pUC19-ELP1 [V-5] to pUC19-ELP1 [V-60] and pUC19-ELP1 [V-120].

Construction of ELP2 Gene Series

[0160] The ELP2 series designate polypeptides containing multiple repeating units of the pentapeptide AVGVV.

[0161] The ELP2 series monomer, ELP2 [5], was created by annealing two 5' phosphorylated, PAGE purified synthetic oligos to form double stranded DNA with *EcoRI* and *HindIII* compatible ends. The ELP2 [5] DNA segment was then ligated into *EcoRI/HindIII* dephosphorylated pUC19 vector to create the pUC19-ELP2[5] monomer. The ELP2 series was expanded in the same manner as the ELP1 [$K_1V_2F_1$] series to create pUC19-ELP2[10], pUC19-ELP2 [30], pUC19-ELP2 [60] and pUC19-ELP2 [120].

Construction of ELP3 [V] Gene Series

[0162] The ELP3 [V] series designate polypeptides containing multiple repeating units of the pentapeptide IPGXG (SEQ ID NO: 5), where X is exclusively valine.

[0163] The ELP3 [V] series monomer, ELP3 [V-5], was created by annealing two 5' phosphorylated, PAGE purified synthetic oligos to form double stranded DNA with PflM1 amino terminal and GGC carboxyl terminal compatible ends due to the lack of a convenient carboxyl terminal restriction site but still enable seamless addition of the monomer. The ELP3 [V-5] DNA segment was then ligated into PflM1/BglI dephosphorylated pUC19-ELP4[V-5], thereby substituting ELP4 [V-5] for ELP3 [V-5] to create the pUC19-ELP3 [V-5] monomer. The ELP3 [V] series was expanded by ligating the annealed ELP3 oligos into pUC19-ELP3[V-5] digested with PflMI. Each ligation expands the ELP3 [V] series by 5 to create ELP3 [V-10], ELP3 [V-15], etc.

Construction of the ELP4 [V] Gene Series

[0164] The ELP4 [V] series designate polypeptides containing multiple repeating units of the pentapeptide LPGXG (SEQ ID NO: 7), where X is exclusively valine.

[0165] The ELP4 [V] series monomer, ELP4 [V-5], was created by annealing two 5' phosphorylated, PAGE purified synthetic oligos to form double stranded DNA with *EcoRI* and *HindIII* compatible ends. The ELP4 [V-5] DNA segment was then ligated into *EcoRI/HindIII* dephosphorylated pUC19 vector to create the pUC19-ELP4[V-5] monomer. The ELP4 [V] series was expanded in the same manner as the ELP1 [$K_1V_2F_1$] series to create pUC19-ELP4[V-10], pUC19-ELP4[V-30], pUC19-ELP4[V-60] and pUC19-ELP4[V-120].

[0166] The ELP genes were also inserted into other vectors such as pET15b-SD0, pET15b-SD3, pET15b-SD5, pET15b-SD6, and pET24d-SD21. The pET vector series are available from Novagen, San Diego, CA.

[0167] The pET15b-SD0 vector was formed by modifying the pET15b vector using SD0 double-stranded DNA segment containing the multicloning restriction site (*SacI-NdeI-NcoI-XhoI-SnaBI-BamHI*). The SD0 double-stranded DNA segment had *XbaI* and *BamHI* compatible ends and was ligated into *XbaI/BamHI* linearized and 5'-dephosphorylated pET15b to form the pET15b-SD0 vector.

[0168] The pET15b-SD3 vector was formed by modifying the pET15b-SD0 vector using SD3 double-stranded DNA segment containing a *SfiI* restriction site upstream of a hinge region- thrombin cleavage site followed by the multicloning site (*NdeI-NcoI-XhoI-SnaBI-BamHI*). The SD3 double-stranded DNA segment had *SacI* and *NdeI* compatible ends and was ligated into *SacI/NdeI* linearized and 5'-dephosphorylated pET15b-SD0 to form the pET15b-SD3 vector.

[0169] The pET15b-SD5 vector was formed by modifying the pET15b-SD3 vector using the SD5 double-stranded DNA segment containing a *SfiI* restriction site upstream of a thrombin cleavage site followed by a hinge and the multicloning site (*NdeI-NcoI-XhoI-SnaBI-BamHI*). The SD5 double-stranded DNA segment had *SfiI* and *NdeI* compatible ends and was ligated into *SfiI/NdeI* linearized and 5'-dephosphorylated pET15b-SD3 to form the pET15b-SD5 vector.

[0170] The pET15b-SD6 vector was formed by modifying the pET15b-SD3 vector using the SD6 double-stranded DNA segment containing a *SfiI* restriction site upstream of a linker region-TEV cleavage site followed by the multicloning site (*NdeI-NcoI-XhoI-SnaBI-BamHI*). The SD6 double-stranded DNA segment had *SfiI* and *NheI* compatible ends and was ligated into *SfiI/NdeI* linearized and 5'-dephosphorylated pET15b-SD3 to form the pET15b-SD6 vector.

[0171] The pET24d-SD21 vector was formed by modifying the pET24d vector using the SD21 double-stranded DNA segment with *NcoI* and *NheI* compatible ends. The SD21 double-stranded DNA segment was ligated into *NcoI/NheI* linearized and 5' dephosphorylated pET24d to create the pET24d-SD21 vector, which contained a new multicloning site *NcoI-SfiI-NheI-BamHI-EcoRI-SacI-Sall-HindIII-NotI-XhoI* with two stop codons directly after the *SfiI* site for insertion and

expression of ELP with the minimum number of extra amino acids.
[0172] The pUC19-ELP1 [V₅A₂G₃-60], pUC19-ELP1 [V₅A₂G₃-90], and pUC19-ELP1 [V₅A₂G₃-180] plasmids produced in XL1-Blue were digested with *PflMI* and *BglI*, and the ELP-containing fragments were ligated into the *SfiI* site of the pET15b-SD3 expression vector as described hereinabove to create pET15b-SD3-ELP1 [V₅A₂G₃-60], pET15b-SD5-ELP1 [V₅A₂G₃-90] and pET15b-SD5-ELP1 [V₅A₂G₃-180], respectively.

[0173] The pUC19-ELP1 [V₅A₂G₃-90], pUC19-ELP1 [V₅A₂G₃-180], pUC19-ELP1 [V-60] and pUC19-ELP1 [V-120] plasmids produced in XL1-Blue were digested with *PflMI* and *BglI*, and the ELP-containing fragments were ligated into the *SfiI* site of the pET15b-SD5 expression vector as described hereinabove to create pET15b-SD5-ELP1 [V₅A₂G₃-90], pET15b-SD5-ELP1 [V₅A₂G₃-180], pET15b-SD5-ELP1 [V-60] and pET15b-SD5-ELP1 [V-120], respectively.

[0174] The pUC19-ELP1 [V₅A₂G₃-90] plasmid produced in XL1-Blue was digested with *PflMI* and *BglI*, and the ELP-containing fragment was ligated into the *SfiI* site of the pET15b-SD6 expression vector as described hereinabove to create pET15b-SD6-ELP1 [V₅A₂G₃-90].

[0175] The pUC19-ELP1 [K₁V₂F₁-64], and pUC19-ELP1 [K₁V₂F₁-128] plasmids produced in XL1-Blue were digested with *PflMI* and *BglI*, and the ELP-containing fragments were ligated into the *SfiI* site of the pET24d-SD21 expression vector as described hereinabove to create pET24d-SD21-ELP1 [K₁V₂F₁-64] and pET24d-SD21-ELP1 [K₁V₂F₁-128], respectively.

[0176] The pUC19-ELP1 [K₁V₇F₁-72] and pUC19-ELP1 [K₁V₇F₁-144] plasmids produced in XL1-Blue were digested with *PflMI* and *BglI*, and the ELP-containing fragments were ligated into the *SfiI* site of the pET24d-SD21 expression vector as described hereinabove to create pET24d-SD21-ELP1 [K₁V₇F₁-72], pET24d-SD21-ELP1 [K₁V₇F₁-144], respectively.

[0177] The pUC19-ELP2[60] and pUC19-ELP2[120] plasmids produced in XL1-Blue were digested with *NcoI* and *HindIII*, and the ELP-containing fragments were ligated into the *NcoI* and *HindIII* sites of the pET24d-SD21 expression vector as described hereinabove to create pET24d-SD21-ELP2[60], pET24d-SD21-ELP2[120], respectively.

[0178] The pUC19-ELP4[V-60] and pUC19-ELP4[V-120] plasmids produced in XL1-Blue were digested with *NcoI* and *HindIII*, and the ELP-containing fragments were ligated into the *NcoI* and *HindIII* sites of the pET24d-SD21 expression vector as described hereinabove to create pET24d-SD21-ELP4[V-60], pET24d-SD21-ELP4[V-120], respectively.

EXAMPLE 2: ISOLATION AND PURIFICATION OF FUSION PROTEINS CONTAINING INSULIN A PEPTIDE (InsA)

[0179] ELP-InsA fusion proteins included the following:

[0180] Insulin A peptide and ELP1 [V-60] polypeptide with an enterokinase protease cleavage site therebetween.

[0181] Insulin A peptide and ELP1 [V₅A₂G₃-90] polypeptide with an enterokinase protease cleavage site therebetween.

[0182] Insulin A peptide and ELP1 [V-120] polypeptide with an enterokinase protease cleavage site therebetween.

[0183] Insulin A peptide and ELP1 [V₅A₂G₃-180] polypeptide with an enterokinase protease cleavage site therebetween.

[0184] A single colony of *E. coli* strain BLR (DE3) (Novagen) containing the respective ELP-InsA fusion protein was inoculated into 5 ml CircleGrow (Q-BIOgene, San Diego, CA) supplemented with 100 µg/ml ampicillin (Sigma) and grown at 37°C with shaking at 250 rpm for 5 hours. The 5 ml culture was then inoculated into a 500 ml culture and allowed to grow at 25°C for 16 hours before inducing with 1 mM IPTG for 4 hours at 25°C. The culture was harvested and suspended in 40 ml 20 mM Tris-HCl pH 7.4, 50 mM NaCl, 1 mM DTT and 1 Complete EDTA free Protease inhibitor pellet (Roche, Indianapolis, IN). Cells were lysed by ultrasonic disruption on ice for 3 minutes, which consisted of 10 seconds bursts at 35% power separated by 30 second cooling down intervals. Cell debris was removed by centrifugation at 20,000g, 4°C for 30 minutes.

[0185] Inverse phase transition was induced by adding NaCl to the cell lysate at room temperature to achieve a final concentration of 1.0 M therein, followed by centrifugation at 20,000g for 15 minutes at room temperature. The resulting pellet contained the respective ELP-InsA fusion protein and non-specifically NaCl precipitated proteins.

[0186] The pellet was re-suspended in 40 ml ice-cold 20 mM Tris-HCl pH 7.4, 50 mM NaCl, 1 mM DTT and re-centrifuged at 20,000 g, 4°C for 15 minutes to remove the non-specifically NaCl precipitated proteins. The inverse transition cycle was repeated two additional times to increase the purity of the respective ELP-InsA fusion protein and reduce the final volume to 0.5 ml.

EXAMPLE 3: HALF-LIFE OF ELP1

[0187] The pharmacokinetics of ELP1 were determined by intravenously administering [¹⁴C]ELP1 to nude mice (Balb/c

nu/nu) bearing a leg/flank FaDu xenograft and collecting blood samples at various time intervals after administration. The blood pharmacokinetics exhibited a characteristic distribution and elimination response for large macromolecules, which was well described by a bi-exponential process.

[0188] The plasma concentration time-course curve was fit to the analytical solution of a two-compartment model to approximate both an elimination and distribution response. Certain pharmacokinetic parameters are shown in Table 1 below. The distribution volume of the ELP (1.338 μL) was nearly identical to the hypothetical plasma volume of 1.363 μL (Barbee, R.W., et al., Am. J. Physio. 263(3) (1992) R728-R733), indicating that the ELP did not rapidly distribute or bind to specific organs and tissues directly after administration. The AUC is a measure of the cumulative exposure to ELP in the central compartment or the blood plasma. The body clearance is defined as the rate of ELP elimination in the body relative to its plasma concentration and is the summation of clearance through all organs including the kidney, liver and others.

Table 1: Pharmacokinetic parameters calculated for [^{14}C]ELP1

	k_1 (hr^{-1})	k_2 (hr^{-1})	k_e (hr^{-1})	V_d (μL)	AUC (mg ELP hr/ml)	Cl_B ($\mu\text{L/hr}$)
ELP1-150	3.54	1.99	0.24	1,338	7.1	317

[0189] The mass transfer rate constants are from a standard two-compartment model (k_1 ; from central to peripheral compartment; k_2 , from peripheral to central compartment; and k_e , elimination from central compartment). The distribution volume (V_d), central compartment concentration time-course area under the curve (AUC) and body clearance (Cl_B) are displayed. Data are shown as the mean values ($n = 5$, except V_d and initial plasma concentration (C_0) was calculated from a similar cohort with $n = 3$).

EXAMPLE 4: BIODISTRIBUTION OF ELPs IN NUDE MICE

^{14}C labeled ELP1-150 and/or ^{14}C labeled ELP2-160

[0190] ^{14}C labeled ELP1-150 and/or ^{14}C labeled ELP2-160 were administered to nude mice with a FaDu tumor (mean $\pm$ SD, $n=6$). The tumor was heated post administration of the ELP in a water bath at 41.5°C . The distribution was highest to the organs with the highest blood content: liver, kidneys, spleen, and lungs.

^{14}C labeled ELP2-[$\text{V}_1\text{A}_8\text{G}_7$ -160]

[0191] ^{14}C labeled ELP2-[$\text{V}_1\text{A}_8\text{G}_7$ -160] ($T_t > 60^\circ\text{C}$) was administered to nude mice for a plasma concentration of $15\mu\text{M}$. ELP concentrations were determined following 1 hour of heating (41°C) of an implanted FaDu tumor, located in the right hind leg of the nude mouse. Data are shown as the mean, plus the 95% confidence interval. $N=6$.

[0192] ELP concentration was measured 1.5 hours following systemic administration of ^{14}C labeled ELP2-[$\text{V}_1\text{A}_8\text{G}_7$ -160]. The highest distribution is seen in organs with the highest blood content: liver, kidneys, spleen, and lungs.

EXAMPLE 5: EXENDIN-4 ELP FUSION

[0193] The DNA sequence for Exendin-4 (Ex-4) (SEQ ID NO: 14) was reverse translated from the amino acid sequence using codons optimized for *E. coli* expression. The DNA sequence encoding Exendin-4 was constructed by annealing together synthetic oligonucleotides with overhanging 5' and 3' ends compatible with the restriction sites *NdeI* and *XhoI* in the plasmid pET24d-ELP1-90 (Figure 1). This plasmid was digested with the restriction enzymes *NdeI* and *XhoI* and the annealed DNA sequence was ligated into the cut vector. Insertion was confirmed by restriction digest and DNA sequencing. The resulting plasmid was designated as pET24d-Ex-4 ELP1-90 (Figure 2A), and the sequence of the resulting Exendin-4-ELP fusion shown in Figure 2B. Primers for construction of the fusion are also indicated.

[0194] pET24d-Ex-4 ELP1-90 was used to transform the *E. coli* strain BRL (Invitrogen) and selected transformants were grown in media 3 (1.2% Tryptone Peptone, 2.4% yeast extract, 5g/L casamino acids, 2% glycerol, 2.313g Potassium phosphate dibasic/L, 12.541g Potassium phosphate monobasic/L) in shake flasks. Production proceeded by autoinduction by inoculating 1 OD cells into 1 L of media 3 and allowing growth to proceed for 17 hr at 37°C without addition of inducer. The product was recovered by collection of the cell pellet, sonicated to disrupt the cells and recovered by thermal and/or salt induced transition modulated by the ELP moiety (Improved Non-chromatographic Purification of a Recombinant Protein by Cationic Elastin-like Polypeptides, Dong Woo Lim, Kimberly Trabbic-Carlson, J. Andrew MacKay, and Ashutosh Chilkoti. Biomacromolecules 2007, 8, 1417-1424).

[0195] This example is with the ELP designated 1-90. This is based on the VPGXG (SEQ ID NO: 3) motif where X is

a V, G or A in the ratio 5:3:2 in a 10 unit repeat, repeated 8x with a final (C-terminal) 10-unit repeat where X is a V, G, A and W in the ratio 4:3:2:1.

[0196] [(VPGXG)₁₀]₉ where the X residue in the ten sequential iterations of the repeat unit (numerical subscript) can be described as [(V_{1, 4, 5, 6, 10}G_{2, 7, 9}A_{3, 8})₈(V_{1, 4, 5, 6}G_{2, 7, 9}A_{3, 8}W₁₀)].

[0197] The ELP may be any combination of VPGXG (SEQ ID NO: 3) units where X is any of the 20 natural amino acids, except proline, in any combination of repeat units of any length. In addition, the amino acid may be an unnatural amino acid for which the host strain has been engineered to accept an engineered tRNA for incorporation at specific codon (Wang L, Brock A, Herberich B, Schultz PG. Expanding the genetic code of Escherichia coli. [2001] Science 292, 498-500).

[0198] This construct was produced in the cytosol with an N-terminal methionine, which is normally removed by methionine aminopeptidase. Complete and accurate processing of the methionine, however, cannot be assumed; this enzyme may also remove the N-terminal histidine of the Exendin-4 moiety. This could result in a mixture of, unprocessed, processed and incorrectly processed products. Consequently, further constructs were developed to generate products with correctly processed N-termini.

[0199] Primers were designed to add a Tev protease (Tobacco Etch Virus cysteine protease) cleavage site between the N-terminal methionine and the histidine at the N-terminus of Exendin-4. This allows for removal of the methionine and the Tev recognition sequence to give the mature N-terminus of Exendin-4 (histidine). This can be done post-production or the Tev protease can be co-expressed to cleave the recognition sequence during production, for instance, as an intein (Ge, X., Yang, D.S.C., Trabbic-Carlson, K., Kim, B., Chilkoti, A. and Filipe, C.D.M. Self-Cleavable Stimulus Responsive Tags for Protein Purification without Chromatography. J. Am. Chem. Soc. 127, 11228-11229, 2005). The Tev Exendin-4 sequence is shown in **Figure 3A**. **Figure 3B** shows additional sequences added, labeled as "Linker Tev," provide a better target for the Tev protease.

[0200] An alternative route to obtaining a correctly processed N-terminus for Ex-4 is to use a leader or signal sequence that directs the product to the periplasm and which is cleaved by a signal peptidase in the process. In this instance, a signal sequence, DsbA, that directs the transcript to the signal recognition particle for direct secretion of the polypeptide into the periplasm is given. (See **Figure 4A**). The plasmid pET24d-DsbA-Ex-4 ELP1-90 is shown in **Figure 4B**.

[0201] While this example illustrates the preparation of therapeutic agents with Exendin-4 sequences, such sequences can be replaced with GLP-1, insulin, Factor VII/VIIa, or other therapeutic protein listed in Table 1, generated in exactly or a similar manner as detailed for Exendin-4.

EXAMPLE 6: GLP1-ELP FUSION PROTEIN

[0202] The ELP plasmid constructs were used to prepare two GLP1-ELP fusion proteins, GLP1(A8G,7-37)ELP1-90 and GLP1(A8G,7-37)ELP1-120. The plasmid constructs, fusion-encoding nucleotide sequence, as well as the amino acid sequence of the resulting fusion proteins are shown in **Figures 5 and 6**.

[0203] Both constructs contain an N-terminal Tev protease site to allow processing to the mature form where His⁷ of GLP1 is at the N-terminus. The processed fusion proteins have calculated molecular weights of about 39,536 and about 50,828, respectively.

EXAMPLE 7: FVII ELP FUSION PROTEIN

[0204] The coagulation factor VII (FVII) gene was modified by PCR from a cDNA clone (Oragene) to add restriction sites at the 5' and 3' ends for cloning into the ELP-containing vector. At the 5' end an *NheI* site was added and at the 3' end a *NotI* site was added. The DNA and amino acid sequences of the Factor VII gene are shown in the accompanying Sequence Listing as SEQ ID NOS: 34 and 33, respectively. The DNA sequences of the 5' and 3' primers used to PCR amplify the factor VII (FVII) gene were:

P13: CTAGCTAGCATGGTCTCCCAGGCCCTC (SEQ ID NO.: 49)

P14: TATTCTTGCGGCCGCGGAAATGGGGCTCGCAG (SEQ ID NO.: 50)

[0205] The resulting PCR fragment was digested with the restriction enzymes *NheI* and *NotI* and ligated into the plasmid pcDNA3.1+ ELP1-90 previously digested with the restriction enzymes *NheI* and *NotI* (**Figure 7A**).

[0206] The resulting plasmid, pcDNA3.1+ FVII-ELP1-90, was transiently transfected into HEK293 cells and culture media harvested. The ELP fusion was purified by phase transition (**Figures 9 and 10**).

[0207] The nucleotide and amino acid sequences of the FactorVII-ELP fusion is shown in Figure 7B. As shown, the FactorVII-ELP fusion protein contains a Tev protease linker between the FactorVII component and the ELP component. This linker is optional.

EXAMPLE 8: INSULIN ELP FUSION PROTEIN

[0208] The cDNA for the human insulin gene is modified at the 5' and 3' ends for insertion in to pET24d-ELP1-90. The 5' primer adds an N-terminal methionine for bacterial expression and an *NdeI* restriction enzyme site. The 3' primer adds an *XhoI* restriction enzyme site. The PCR product and the plasmid are both digested with the restriction enzymes *NdeI* and *XhoI* and ligated together. The sequence of the insulin (Chains B, C, and A fused to ELP1 is shown in **Figure 8A**.

[0209] Correct insertion is determined by restriction digest and DNA sequencing. The resulting plasmid, designated pET24d Insulin-ELP1-90, is shown in **Figure 8B**.

[0210] The native insulin form is generated after recovery from *E. coli* by treatment with trypsin and carboxypeptidase B to remove the C-peptide chain.

[0211] For correct processing of the N-terminus of the B-chain similar modifications to those made for the Exendin-4 fusion (protease cleavage site, signal sequence) can be implemented (see Example 4). Alternatively, the first two residues can be Met - Arg, which can also be removed by trypsin digestion in production of the final material (R. M. Belagaje, S. G. Reams, S. C. Ly and W. F. Prouty, Increased production of low molecular weight recombinant proteins in *Escherichia coli*. Protein Sci. 6, 1953-1962, 1997).

[0212] Additional constructs would place the insulin cDNA at the 3' end of the ELP for a C-terminal fusion, add linkers between the Insulin and ELP sequences, and/or use modified forms of insulin which have no C-peptide (single chain insulins as described) removing the need for additional processing.

EXAMPLE 9: SYNTHESIS OF THE ELP GENE FOR CONJUGATION

[0213] A gene encoding a 50 amino acid sequence was constructed from chemically-synthesized oligonucleotides using standard molecular biology protocols. The 50 amino acid sequence contained 10 repeats of the pentapeptide VPGXG (SEQ ID NO: 3), where the guest residues (V, G, and A in a 5:3:2 molar ratio) were selected to provide a T_m of 40°C. The gene was oligomerized end-to-end by standard molecular biology techniques, to produce an oligomeric ELP gene. Additionally a single 50 amino acid sequence was constructed containing the 10 repeat pentapeptide VPGXG (SEQ ID NO: 3) polypeptide where the guest residues were V, G, A and C in a 4:3:2:1 molar ratio. This sequence could be added at any cycle of the oligomerization process to introduce a single cysteine residue into the final construct at a chosen point along the length of the construct.

[0214] The example given here is with the ELP designated 1-90. This is based on the VPGXG (SEQ ID NO: 3) motif where X is a V, G or A in the ratio 5:3:2 in a 10-unit repeat, repeated 8x with a final (C-terminal) 10-unit repeat where X is a V, G, A and C in the ratio 4:3:2:1, i.e., [(VPGXG)₁₀]₉ (SEQ ID NO.: 3).

[0215] Alternatively, the residue could be one of either arginine, lysine, aspartic acid or glutamic acid. The purpose of these amino acids is to provide a reactive side chain for the chemical conjugation of, for example, insulin. In this particular case the use of an ELP would be to extend the circulating half-life of the therapeutic protein (e.g., insulin) to provide prolonged basal glucose control. Conjugated to an ELP that transitions at body temperature, the insulin would form a precipitated depot at the site of injection in a similar manner to Lantus® (Sanofi Aventis) but without the requirement for formulation in acidic (pH 4.0) conditions with m-cresol for a more tolerable injection.

EXAMPLE 10: POTENCY AND HALF-LIFE OF FACTOR VII-ELP

[0216] **Figure 11** shows the activation of Factor X by FactorVIIa-ELP1-90, and by Factor VIIa as a comparison. Factor VII-ELP was produced in HEK cells. Factor VIIa was derived from human plasma. As shown, FactorVIIa-ELP retains full activity.

[0217] When administered to rats by i.v., Factor VII-ELP demonstrated a half-life of about 690 minutes. In contrast, Factor VII demonstrated a half-life of 45-60 minutes. Half-life in this example was measured by sandwich ELISA for FactorVII. **Figure 12**.

[0218] Also in contrast, the reported half-life for NovoSeven™ is 45 minutes, the reported half-life for FactorVIIa-albumin fusion is 263 minutes, and the reported half-life for Factor VIIa-PEG is 300 minutes in mice and 600 minutes in dog.

EXAMPLE 11: GLP-1 (OR EXENDIN-4) IN VITRO BIOASSAY

[0219] Activation of the GLP-1 receptor (GLP1 R) results in production of cAMP secondary messenger within the cell. Therefore, GLP-1 or Exendin-4 analogs and corresponding therapeutic agents may be tested by their ability to activate GLP1 R on the cell surface and produce cAMP.

[0220] For this bioassay CHO cells transfected with cDNA coding for GLP1R are used. These cells respond to stimulation by GLP-1 and produce high levels of cAMP. Log phase growing cells are plated and increasing concentrations of test compounds (e.g., therapeutic agent of the invention, or GLP-1 or exendin-4 functional analog) are added to the

cells. After an appropriate incubation period (usually 15-60 min) in physiological buffer at 37°C the cAMP produced is measured using a CatchPoint cAMP assay kit from Molecular Devices (Sunnyvale, CA). The EC₅₀ of each test compound as compared to GLP-1 peptide or Exendin-4 peptide (or as compared to an unfused or unconjugated counterpart of a therapeutic agent of the invention) is indicative of the changes in activity due to a specific modifications introduced into the peptide, or due to particular chemical or recombinant coupling to an ELP component.

[0221] As shown in **Figure 13**, both GLP1-ELP (PB0868) and Exendin-4-ELP (PB 0859) maintain high activity *in vitro*, shown in comparison to Exendin alone. It is of note that the specific activity of Albugon® and Liraglutide® run 50-100 fold less than the exendin peptide.

EXAMPLE 12: GLP-1 (OR EXENDIN-4) IN VIVO BIOASSAY

[0222] The activity of GLP-1 or Exendin analogues or corresponding therapeutic agents may be tested in animals. For this assay, normal or diabetic animals may be used. Diabetic animals with blood glucose concentration 300-500 mg/dl are injected with different doses of GLP-1 or Exendin analogues or corresponding therapeutic agent, and changes in blood glucose monitored with a glucometer. The drop in glucose at different times points post administration is compared to that resulting with standard amounts of GLP-1 or Exendin-4 peptide, or compared to an unfused or unconjugated counterpart of a therapeutic agent of the invention. Alternatively, the blood glucose excursion in normal or diabetic animals during specific time period after administration of exogenous glucose is compared to GLP-1 or Exendin-4 (or to unfused or unconjugated counterparts of therapeutic agents). In this way the activity of the analogues and fusion proteins can be compared to the natural peptides.

[0223] **Figure 14** shows the pharmacokinetics of GLP1-ELP1-120 in rats administered both by i.v. and subcutaneously. Three rats were used for each time point. The dose was ~10mg/kg. The T_{1/2} when administered by i.v. was about 12.9 hours. The T_{1/2} when administered subcutaneously was about 8.6 hours.

[0224] **Figure 15** shows the pharmacokinetics of GLP1-ELP1-120 in rabbits administered both by i.v. and subcutaneously. Three rabbits were used for each time point. The dose was ~1mg/kg. The T_{1/2} when administered by i.v. was about 20 hours. The T_{1/2} when administered subcutaneously was about 24 hours.

[0225] **Figure 16** shows the sustained glycemic control in diabetic mice with GLP1-ELP1-90.

[0226] All reference cited herein are hereby incorporated by reference in their entireties. While the invention has been described herein in reference to specific aspects, features and illustrative embodiments of the invention, it will be appreciated that the utility of the invention is not thus limited, but rather extends to and encompasses numerous other variations, modifications and alternative embodiments, as will suggest themselves to those of ordinary skill in the field of the present invention, based on the disclosure herein

The following numbered clauses, describing aspects of the invention, are part of the description.

1. A therapeutic agent comprising an elastin-like peptide (ELP) component and a therapeutic component, the therapeutic component having an extended circulatory half-life and/or a lower therapeutically effective dose when compared to the therapeutic component alone.

2. The therapeutic agent of clause 1, wherein the therapeutic component is a therapeutic protein listed in Table 1 or a functional portion or functional analog thereof.

3. The therapeutic agent of clause 2, wherein an ELP component is covalently bonded to the therapeutic protein at an N- and/or C-terminus thereof.

4. The therapeutic agent of any one of clauses 1-3, wherein a therapeutic component is covalently bonded to the ELP component at the N-terminus of the ELP component, and a second therapeutic component is covalently bonded to the ELP component at the C-terminus of the ELP component.

5. The therapeutic agent of any one of clauses 1-4, wherein a therapeutic component is covalently bonded to the ELP component at an N- and/or C-terminus thereof.

6. The therapeutic agent of any one of clauses 1-5, wherein a first ELP component is covalently bonded to the therapeutic component at the N- terminus of the therapeutic component, and a second ELP component is covalently bonded to the therapeutic component at the C- terminus of the therapeutic component.

7. The therapeutic agent of any one of clauses 1-6, further comprising a spacer moiety between the ELP component and the therapeutic component.

8. The therapeutic agent of clause 7, wherein the spacer moiety comprises one or more of a protease-resistant moiety, a non-peptide chemical moiety, and a protease cleavage site.

9. The therapeutic agent of clause 8, wherein the protease cleavage site is a thrombin cleavage site, a factor Xa cleavage site, a metalloprotease cleavage site, an enterokinase cleavage site, a Tev cleavage site, and a cathepsin cleavage site.

10. The therapeutic agent of clause 9, wherein the spacer moiety comprises a non-cleavable moiety having the formula $[(\text{Gly})_n\text{-Ser}]_m$ (SEQ ID NO: 22) where n is from 1 to 4, inclusive, and m is from 1 to 4, inclusive.

11. The therapeutic agent of any one of clauses 1-10, wherein the ELP comprises at least one repeating unit selected from SEQ ID NOS: 1-12.

12. The therapeutic agent of clause 11, wherein said repeating unit is VPGXG (SEQ ID NO: 3).

13. The therapeutic agent of clause 12, wherein X is any natural or non-natural amino acid residue, and wherein X varies among at least two units.

14. The therapeutic agent of clause 13, wherein each X is independently selected from alanine, arginine, asparagine, aspartic acid, glutamic acid, glutamine, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, serine, threonine, tryptophan, tyrosine and valine residues.

15. The therapeutic agent of any one of clauses 1-14, wherein the therapeutic agent has a molecular weight of less than about 50 kDa.

16. The therapeutic agent of any one of clauses 1-15, wherein the ELP component has a Tt greater than 37°C.

17. The therapeutic agent of any one of clauses 1-16, wherein the therapeutic agent is a genetically encoded fusion protein.

18. A polynucleotide comprising a nucleotide sequence encoding the therapeutic agent of clause 17.

19. The polynucleotide of clause 18, further comprising an expression control element operably linked to said nucleotide sequence.

20. The polynucleotide of clause 19, wherein said expression control element comprises a promoter.

21. A vector comprising the polynucleotide of any one of clauses 18-20.

22. An isolated host cell containing the vector of clause 21 or the polynucleotide of any one of clauses 18-20.

23. A pharmaceutical composition comprising the therapeutic agent of any one of clauses 1-18, and a pharmaceutically acceptable carrier and/or excipient.

24. A method of treating or preventing a disease, disorder, or condition in a subject, comprising administering an effective amount of the therapeutic agent of any one of clauses 1-17 or the pharmaceutical composition of clause 23 to a subject in need thereof.

25. The method of clause 24, wherein the subject has an indication listed in Table 1.

26. A therapeutic agent comprising an elastin-like peptide (ELP) component and a glucagon-like peptide (GLP)-1 receptor agonist.

27. The therapeutic agent of clause 26, wherein the GLP-1 receptor agonist is GLP-1 or functional analog thereof.

28. The therapeutic agent of clause 26, wherein the GLP-1 receptor agonist is an exendin-4 or functional analog thereof.

29. The therapeutic agent of any one of clauses 26-28, wherein the ELP component is covalently bonded to the GLP-1 receptor agonist at an N- and/or C-terminus thereof.

30. The therapeutic agent of any one of clauses 26-29, wherein a first GLP-1 receptor agonist is covalently bonded to the ELP component at the N-terminus of the ELP component, and a second GLP-1 receptor agonist is covalently bonded to the ELP component at the C-terminus of the ELP component.

31. The therapeutic agent of any one of clauses 26-30, wherein the GLP-1 receptor agonist is covalently bonded to the ELP component at an N- and/or C-terminus thereof.

32. The therapeutic agent of any one of clauses 26-31, wherein a first ELP component is covalently bonded to the GLP-1 receptor agonist at the N-terminus of the GLP-1 receptor agonist, and a second ELP component is covalently bonded to the GLP-1 receptor agonist at the C-terminus of the GLP-1 receptor agonist.

33. The therapeutic agent of any one of clauses 26-32, further comprising a spacer moiety between the ELP component and GLP-receptor agonist.

34. The therapeutic agent of clause 33, wherein the spacer moiety comprises one or more of a protease-resistant moiety, a non-peptide chemical moiety, and a protease cleavage site.

35. The therapeutic agent of clause 34, wherein the protease cleavage site is a thrombin cleavage site, a factor Xa cleavage site, a metalloprotease cleavage site, an enterokinase cleavage site, a Tev cleavage site, and a cathepsin cleavage site.

36. The therapeutic agent of clause 34, wherein the spacer moiety comprises a non-cleavable moiety having the formula $[(\text{Gly})_n\text{-Ser}]_m$ (SEQ ID NO.: 22) where n is from 1 to 4, inclusive, and m is from 1 to 4, inclusive.

37. The therapeutic agent of any one of clauses 26-36, wherein the ELP comprises at least one repeating unit selected from SEQ ID NOS: 1-12.

38. The therapeutic agent of clause 37, wherein said repeating unit is VPGXG (SEQ ID NO: 3).

39. The therapeutic agent of clause 38, wherein X is any natural or non-natural amino acid residue, and wherein X varies among at least two units.

40. The therapeutic agent of clause 38, wherein each X is independently selected from alanine, arginine, asparagine, aspartic acid, glutamic acid, glutamine, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, serine, threonine, tryptophan, tyrosine and valine residues.

41. The therapeutic agent of any one of clauses 26-40, wherein the therapeutic agent has a molecular weight of less than about 50 kDa.

42. The therapeutic agent of any one of clauses 26-40, wherein the ELP component has a Tt greater than 37°C.

43. The therapeutic agent of any one of clauses 26-42, wherein the therapeutic agent is a genetically encoded fusion protein.

44. A polynucleotide comprising a nucleotide sequence encoding the therapeutic agent of clause 43.

45. The polynucleotide of clause 44, further comprising an expression control element operably linked to said nucleotide sequence.

46. The polynucleotide of clause 45, wherein said expression control element comprises a promoter.

47. A vector comprising the polynucleotide of any one of clause 44-46.

48. An isolated host cell containing the vector of clause 47 or the polynucleotide of any one of clauses 44-46.

49. A pharmaceutical composition comprising the therapeutic agent of any one of clauses 26-43, and a pharmaceutically acceptable carrier and/or excipient.

50. A method of treating or preventing a biological condition, disorder, or disease in a subject, comprising administering an effective amount of the therapeutic agent of any one of clauses 26-43 or the pharmaceutical composition of clause 49 to a subject in need thereof.

51. The method of clause 50, wherein the subject has diabetes.

52. A therapeutic agent comprising an elastin-like peptide (ELP) component and an insulin or functional analog thereof.

53. The therapeutic agent of clause 52, wherein the ELP component is covalently bonded to the insulin or functional analog at an N- and/or C-terminus thereof.

54. The therapeutic agent of clause 52 or 53, wherein a first insulin or functional analog is covalently bonded to the ELP component at the N-terminus of the ELP component, and a second insulin or functional analog is covalently bonded to the ELP component at the C-terminus of the ELP component.

55. The therapeutic agent of clause 52, wherein the insulin or functional analog is covalently bonded to the ELP component at an N- and/or C-terminus thereof.

56. The therapeutic agent of clause 52 or 53, wherein a first ELP component is covalently bonded to the insulin or functional analog at an N- terminus thereof, and a second ELP component is covalently bonded to the insulin or functional analog at a C- terminus thereof.

57. The therapeutic agent of any one of clauses 52-56, further comprising a spacer moiety between the ELP component and the insulin or functional analog.

58. The therapeutic agent of clause 57, wherein the spacer moiety comprises one or more of a protease-resistant moiety, a non-peptide chemical moiety, and a protease cleavage site.

59. The therapeutic agent of clause 58, wherein the protease cleavage site is a thrombin cleavage site, a factor Xa cleavage site, a metalloprotease cleavage site, an enterokinase cleavage site, a Tev cleavage site, or a cathepsin cleavage site.

60. The therapeutic agent of clause 58, wherein the spacer moiety comprises a non-cleavable moiety having the formula $[(\text{Gly})_n\text{-Ser}]_m$ (SEQ ID NO: 22) where n is from 1 to 4, inclusive, and m is from 1 to 4, inclusive.

61. The therapeutic agent of any one of clauses 52-60, wherein the ELP component comprises at least one repeating unit selected from SEQ ID NOS: 1-12.

62. The therapeutic agent of clause 61, wherein said repeating unit is VG PXG (SEQ ID NO: 3).

63. The therapeutic agent of clause 61, wherein the repeating unit is Ile-Pro-Gly-X-Gly (SEQ ID NO: 5) or Leu-Pro-Gly-X-Gly (SEQ ID NO: 7), wherein X is any natural or non-natural amino acid residue, and wherein X varies among at least two units.

64. The therapeutic agent of clause 62 or 63, wherein each X is independently selected from alanine, arginine, asparagine, aspartic acid, glutamic acid, glutamine, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, serine, threonine, tryptophan, tyrosine and valine residues.

65. The therapeutic agent of any one of clauses 52-64, wherein the therapeutic agent has a molecular weight of less than about 50 kDa.

66. The therapeutic agent of any one of clauses 52-65, wherein the ELP component has a T_m greater than 37°C.

67. The therapeutic agent of any one of clauses 52-66, wherein the therapeutic agent is a genetically encoded fusion

protein.

68. A polynucleotide comprising a nucleotide sequence encoding the therapeutic agent of clause 67.

69. The polynucleotide of clause 68, further comprising an expression control element operably linked to said nucleotide sequence.

70. The polynucleotide of clause 69, wherein said expression control element comprises a promoter.

71. A vector comprising the polynucleotide of any one of clauses 68-70.

72. An isolated host cell containing the vector of clause 71 or the polynucleotide of any one of clauses 68-70.

73. A pharmaceutical composition comprising the therapeutic agent of any one of clauses 52-67, and a pharmaceutically acceptable carrier and/or excipient.

74. A method of preventing or treating a biological condition, disorder, or disease in a subject, comprising administering an effective amount of the therapeutic agent of any one of clauses 52-67 or the pharmaceutical composition of clause 73 to a subject in need thereof.

75. The method of clause 74, wherein the subject has diabetes.

76. A therapeutic agent comprising an elastin-like peptide (ELP) component and a Factor VII/VIIa or functional analog thereof.

77. The therapeutic agent of clause 76, wherein the ELP component is covalently bonded to the Factor VII/VIIa or functional analog at an N- and/or C-terminus of the Factor VII/VIIa.

78. The therapeutic agent of clause 76 or 77, wherein a first Factor VII/VIIa or functional analog is covalently bonded to the ELP at the N-terminus of the ELP component, and a second Factor VII/VIIa or functional analog is covalently bonded to the ELP component at the C-terminus of the ELP component.

79. The therapeutic agent of clause 76, wherein the Factor VII/VIIa or functional analog is covalently bonded to the ELP component at an N- and/or C-terminus of the ELP component.

80. The therapeutic agent of clause 76 or 77, wherein a first ELP component is covalently bonded to the Factor VII/VIIa or functional analog at an N-terminus thereof, and a second ELP component is covalently bonded to the Factor VII/VIIa or functional analog at a C-terminus thereof.

81. The therapeutic agent of any one of clauses 76-80, further comprising a spacer moiety between the ELP component and factor VII/VIIa or functional analog.

82. The therapeutic agent of clause 81, wherein the spacer moiety comprises one or more of a protease-resistant moiety, a non-peptide chemical moiety, and a protease cleavage site.

83. The therapeutic agent of clause 82, wherein the protease cleavage site is a thrombin cleavage site, a factor Xa cleavage site, a metalloprotease cleavage site, an enterokinase cleavage site, a Tev cleavage site, and a cathepsin cleavage site.

84. The therapeutic agent of clause 82, wherein the spacer moiety comprises a non-cleavable moiety having the formula $[(\text{Gly})_n\text{-Ser}]_m$ (SEQ ID NO: 22) where n is from 1 to 4, inclusive, and m is from 1 to 4, inclusive.

85. The therapeutic agent of any one of clauses 76-84, wherein the ELP component comprises at least one repeating unit selected from SEQ ID NOS: 1-12.

86. The therapeutic agent of clause 85, wherein said repeating unit is VPGXG (SEQ ID NO: 3).

87. The therapeutic agent of clause 86, wherein X is any natural or non-natural amino acid residue, and wherein X

varies among at least two units.

88. The therapeutic agent of clause 86, wherein each X is independently selected from alanine, arginine, asparagine, aspartic acid, glutamic acid, glutamine, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, serine, threonine, tryptophan, tyrosine and valine residues.

89. The therapeutic agent of any one of clauses 76-88, wherein the therapeutic agent has a molecular weight of less than about 70 kDa.

90. The therapeutic agent of any one of clause 76-89, wherein the ELP component has a Tt greater than 37°C.

91. The therapeutic agent of any one of clauses 76-90, wherein the therapeutic agent is a genetically encoded fusion protein.

92. A polynucleotide comprising a nucleotide sequence encoding the therapeutic agent of clause 91.

93. The polynucleotide of clauses 92, further comprising an expression control element operably linked to said nucleotide sequence.

94. The polynucleotide of clause 93, wherein said expression control element comprises a promoter.

95. A vector comprising the polynucleotide of any one of clauses 92-94.

96. An isolated host cell containing the vector of clause 95 or the polynucleotide of any one of clauses 92-95.

97. A pharmaceutical composition comprising the therapeutic agent of any one of clauses 76-91, and a pharmaceutically acceptable carrier and/or excipient.

98. A method of preventing or treating a subject for a biological condition, disorder, or disease, comprising administering an effective amount of the therapeutic agent of any one of clauses 76-91 or the pharmaceutical composition of clause 97 to a subject in need thereof.

99. The method of clause 98, wherein the subject has a hemorrhage.

100. The method of clause 98 or 99, wherein the subject has hemophilia.

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<211> 44
<212> PRT
<213> Unknown

<220>
<223> GLP-1 receptor agonist

<400> 18

45 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

50 Ser Gly Ala Pro Pro Ser Lys Lys Lys Lys Lys Lys
35 40

55 <210> 19
<211> 31
<212> PRT
<213> Unknown

<220>

<223> GLP-1 receptor agonist

<400> 19

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

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

<210> 20

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<212> PRT

<213> Artificial Sequence

<220>

<223> Peptide linker sequence

<400> 20

Lys Asp Asp Asn Pro Asn Leu Pro Arg Leu Val Arg
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<210> 21

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<223> Peptide linker sequence

<400> 21

Gly Ala Gly Ser Ser Ser Arg Arg Ala Pro Gln Thr
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<210> 22

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<223> Non-cleavable spacer sequence

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<221> MISC_FEATURE

<222> (2)..(4)

<223> Gly residue is optional

<220>

<221> MISC_FEATURE

<222> (6)..(8)

<223> Ser residue is optional

<400> 22

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Gly Gly Gly Gly Ser Ser Ser Ser
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5 <210> 23
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10 <220>
<223> Exendin-4/ELP sequence

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tctgagcaaa cagatggaag aagaagcggc ggcgctgttt attgaatggc tgaaaaacgg 180
cgccccgagc agcggcgcg cgcgcgcgag cctcgagggc atgggtgggc cgggcgtggg 240
20 tgttccgggc gtgggtgttc cgggtggcgg tgtgccgggc gcaggtgttc ctggtgtagg 300

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<211> 69
<212> PRT
<213> Artificial Sequence

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<223> Exendin-4/ELP sequence

30 <400> 24

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1 5 10 15

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

40 Ser Ser Gly Ala Pro Pro Pro Ser Leu Glu Gly Met Gly Gly Pro Gly
35 40 45

Val Gly Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala
50 55 60

45 Gly Val Pro Gly Val
65

50 <210> 25
<211> 140
<212> DNA
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55 <220>
<223> Exendin-4/ELP sequence with N-terminal Tev cleavage site

<400> 25

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gatggaagaa gaagcgggtgc gcctgtttat tgaatggctg aaaaacggcg gcccgagcag 120
5 cggcgcgccg ccgccgagcc 140

<210> 26
<211> 46
<212> PRT
10 <213> Artificial Sequence

<220>
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15 <400> 26

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1 5 10 15

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

Leu Lys Asn Gly Gly Pro Ser Ser Gly Ala Pro Pro Pro Ser
25 35 40 45

<210> 27
<211> 155
<212> DNA
30 <213> Artificial Sequence

<220>
<223> Exendin-4/ELP sequence with N-terminal Tev cleavage site and
additional N-terminal sequence

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cgatctgagc aaacagatgg aagaagaagc ggtgcgcctg tttattgaat ggctgaaaaa 120
40 cggcggcccg agcagcggcg cgccgccgcc gagcc 155

<210> 28
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<212> PRT
45 <213> Artificial Sequence

<220>
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additional N-terminal sequence

50 <400> 28

Met Asp Ile Pro Thr Thr Glu Asn Leu Tyr Phe Gln His Gly Glu Gly
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55 Thr Phe Thr Ser Asp Leu Ser Lys Gln Met Glu Glu Glu Ala Val Arg
20 25 30

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Leu Phe Ile Glu Trp Leu Lys Asn Gly Gly Pro Ser Ser Gly Ala Pro
35 40 45

5 Pro Pro Ser
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10 <210> 29
<211> 176
<212> DNA
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15 <220>
<223> Exendin-4/ELP sequence with DsbA leader sequence

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tggcgaaggc acctttacca gcgatctgag caaacagatg gaagaagaag cgggtgcgcct 120
20 gtttattgaa tggctgaaaa acggcggccc gagcagcggc gcgccgccgc cgagcc 176

25 <210> 30
<211> 58
<212> PRT
<213> Artificial Sequence

30 <220>
<223> Exendin-4/ELP sequence with DsbA leader sequence

<400> 30

35 Met Lys Lys Ile Trp Leu Ala Leu Ala Gly Leu Val Leu Ala Phe Ser
1 5 10 15

Ala Ser Ala His Gly Glu Gly Thr Phe Thr Ser Asp Leu Ser Lys Gln
20 25 30

40 Met Glu Glu Glu Ala Val Arg Leu Phe Ile Glu Trp Leu Lys Asn Gly
35 40 45

45 Gly Pro Ser Ser Gly Ala Pro Pro Pro Ser
50 55

50 <210> 31
<211> 400
<212> DNA
<213> Artificial Sequence

<220>
<223> Insulin/ELP sequence

55 <400> 31
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 acacacccaa gaccgcggtg gaggcagagg acctgcaggt ggggcaggtg gagctgggag 180
 5 ggggccctgg tgcaggcagc ctgcagccct tggccctgga ggggtccctg cagaagcgtg 240
 gcattgtgga acaatgctgt accagcatct gctccctcta ccagctggag aactactgca 300
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<210> 32
 <211> 119
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<220>
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 20 <400> 32

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25 Tyr Leu Val Cys Gly Glu Arg Gly Phe Phe Tyr Thr Pro Lys Thr Arg
 20 25 30

30 Arg Glu Ala Glu Asp Leu Gln Val Gly Gln Val Glu Leu Gly Gly Gly
 35 40 45

Pro Gly Ala Gly Ser Leu Gln Pro Leu Ala Leu Glu Gly Ser Leu Gln
 50 55 60

35 Lys Arg Gly Ile Val Glu Gln Cys Cys Thr Ser Ile Cys Ser Leu Tyr
 65 70 75 80

40 Gln Leu Glu Asn Tyr Cys Asn Leu Glu Gly Met Gly Gly Pro Gly Val
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Gly Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly
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45 Val Pro Gly Val Gly Val Pro
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50 <210> 33
 <211> 406
 <212> PRT
 <213> Homo sapiens

55 <400> 33

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Asp	Ala	Glu	Arg	Thr	Lys	Leu	Phe	Trp	Ile	Ser	Tyr	Ser	Asp	Gly	Asp
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Gln	Cys	Ala	Ser	Ser	Pro	Cys	Gln	Asn	Gly	Gly	Ser	Cys	Lys	Asp	Gln
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Leu	Gln	Ser	Tyr	Ile	Cys	Phe	Cys	Leu	Pro	Ala	Phe	Glu	Gly	Arg	Asn
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Cys	Glu	Thr	His	Lys	Asp	Asp	Gln	Leu	Ile	Cys	Val	Asn	Glu	Asn	Gly
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Lys	Gly	Glu	Cys	Pro	Trp	Gln	Val	Leu	Leu	Leu	Val	Asn	Gly	Ala	Gln
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Leu	Cys	Gly	Gly	Thr	Leu	Ile	Asn	Thr	Ile	Trp	Val	Val	Ser	Ala	Ala
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His	Cys	Phe	Asp	Lys	Ile	Lys	Asn	Trp	Arg	Asn	Leu	Ile	Ala	Val	Leu
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Val	Ala	Gln	Val	Ile	Ile	Pro	Ser	Thr	Tyr	Val	Pro	Gly	Thr	Thr	Asn
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His	Asp	Ile	Ala	Leu	Leu	Arg	Leu	His	Gln	Pro	Val	Val	Leu	Thr	Asp
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	Asp	Arg	Gly	Ala	Thr	Ala	Leu	Glu	Leu	Met	Val	Leu	Asn	Val	Pro	Arg	
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	Glu	Trp	Leu	Gln	Lys	Leu	Met	Arg	Ser	Glu	Pro	Arg	Pro	Gly	Val	Leu	
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	accctgatca acaccatctg ggtggtctcc gcgcccact gtttcgacaa aatcaagaac	600
	tggaggaacc tgatcgcggt gctgggcgag cacgacctca gcgagcacga cggggatgag	660
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 <212> DNA
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<211> 59
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 Arg Glu Ala Glu Asp Leu Gln Val Gly Gln Val Glu Leu Gly Gly Gly
 35 40 45
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 Pro Gly Ala Gly Ser Leu Gln Pro Leu Ala Leu Glu Gly Ser Leu Gln
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 Lys Arg Gly Ile Val Glu Gln Cys Cys Thr Ser Ile Cys Ser Leu Tyr
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 <210> 54
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 Leu Val Lys Gly Arg Gly Leu Glu Gly Met Gly Gly Pro Gly Val Gly
 25 35 40 45

 Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val
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 Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly Val Gly Val Pro
 65 70 75 80

 Gly Gly Gly Val Pro Gly Ala Gly Val Pro Gly Gly Gly Val Pro Gly
 35 85 90 95

 Val Gly Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala
 40 100 105 110

 Gly Val Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly Val Gly
 115 120 125
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 Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val Pro Gly Gly Gly Val
 130 135 140

 Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro
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 Gly Ala Gly Val Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly
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 Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val Pro Gly Gly

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45	Val	Pro	Gly	Val	Gly	Val	Pro	Gly	Val	Gly	Val	Pro	Gly	Gly	Gly	Val					
		370					375					380									
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	Gly	Val	Gly	Val	Pro	Gly	Gly	Gly	Val	Pro	Gly	Ala	Gly	Val	Pro	Gly					
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Gly Val Pro Gly Ala Gly Val Pro Gly Gly Gly Val Pro Gly Val Gly
435 440 445

5 Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val
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10 Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly Val Gly Val Pro
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Gly Gly Gly Val Pro Gly Ala Gly Val Pro Gly Gly Gly Val Pro Gly
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15 Trp Pro

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<220>
25 <223> pPB1022 DNA sequence

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

55 Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val Pro Gly Val Gly Val
 50 55 60

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60	Ala Gly Val	Pro Gly Val Gly Val	Pro Gly Val Gly Val	Pro Gly Val
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65	Gly Val	Pro Gly Gly Gly Val	Pro Gly Ala Gly Val	Pro Gly Gly Gly
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70	Val Pro Gly Val Gly Val	Pro Gly Val Gly Val	Pro Gly Gly Gly Val	Pro
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	Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val	370	375	380
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35	Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro	465	470	475
				480
40	Gly Ala Gly Val Pro Gly Gly Gly Val Pro Gly Val Gly Val Pro Gly	485	490	495
	Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val Pro Gly Val	500	505	510
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	Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val Pro	545	550	555
				560
55	Gly Val Gly Val Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly	565	570	575

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Gly Gly Val Pro Gly Ala Gly Val Pro Gly Gly Gly Val Pro Gly Val
580 585 590

5 Gly Val Pro Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly
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10 Val Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly Val Gly Val
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15 Pro Gly Gly Gly Val Pro Gly Ala Gly Val Pro Gly Gly Gly Val Pro
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Gly Trp Pro

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25 <220>
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 55 Ser Tyr Ser Asp Gly Asp Gln Cys Ala Ser Ser Pro Cys Gln Asn Gly
 85 90 95

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60	Gly	Val	Gly	Val	Pro	Gly	Val	Gly	Val	Pro	Gly	Gly	Gly	Val	Pro	Gly
					790						795					800
65	Ala	Gly	Val	Pro	Gly	Gly	Gly	Val	Pro	Gly	Val	Gly	Val	Pro	Gly	Val
					805					810					815	
70	Gly	Val	Pro	Gly	Gly	Gly	Val	Pro	Gly	Ala	Gly	Val	Pro	Gly	Val	Gly
				820					825					830		
75	Val	Pro	Gly	Val	Gly	Val	Pro	Gly	Val	Gly	Val	Pro	Gly	Gly	Gly	Val
			835					840					845			

Pro Gly Ala Gly Val Pro Gly Gly Gly Val Pro Gly Val Gly Val Pro
850 855 860

Gly Val Gly Val Pro Gly Gly Gly Val Pro Gly Ala Gly Val Pro Gly
865 870 875 880

Val Gly Val Pro Gly Val Gly Val Pro Gly Val Gly Val Pro Gly Gly
885 890 895

Gly Val Pro Gly Ala Gly Val Pro Gly Gly Gly Val Pro Gly Trp Pro
900 905 910

<210> 59

<211> 31

<212> PRT

<213> Unknown

<220>

<223> GLP-1 receptor agonist

<400> 59

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 Gly
20 25 30

Claims

1. A therapeutic agent for use in a method of treating diabetes, the method comprising administering the therapeutic agent to a subject, wherein said therapeutic agent is a fusion protein comprising an elastin-like peptide (ELP) component and an insulin A chain and an insulin B chain.
2. The therapeutic agent for the use according to claim 1, wherein the insulin A chain comprises an amino acid sequence having from 1 to 10 amino acid insertions, deletions, and/or substitutions relative to SEQ ID NO: 15, and/or wherein the insulin B chain comprises an amino acid sequence having from 1 to 10 amino acid insertions, deletions, and/or substitutions relative to SEQ ID NO: 16.
3. The therapeutic agent for the use according to any one of the preceding claims, wherein the insulin A chain is attached to the N-terminus of the ELP component.
4. The therapeutic agent for the use according to any one of the preceding claims, wherein the ELP component comprises at least one repeating unit selected from SEQ ID NOs: 1-12.
5. The therapeutic agent for the use according to claim 4, wherein the repeating unit is VPGXG (SEQ ID NO: 3), and wherein X is any natural amino acid residue.
6. The therapeutic agent for the use according to claim 5, wherein each X is V, G, or A at a ratio of 5:3:2, optionally wherein the ELP component comprises 10 repeats, 60 repeats, 90 repeats, 120 repeats or 180 repeats of VPGXG (SEQ ID NO: 3).

7. The therapeutic agent for the use according to claim 5, wherein each X is K, V, or F at a ratio of 1:2:1, optionally wherein the ELP component comprises 4 repeats, 8 repeats, 16 repeats, 32 repeats or 64 repeats of VPGXG (SEQ ID NO: 3).
- 5 8. The therapeutic agent for the use according to claim 5, wherein each X is K, V, or F at a ratio of 1:7:1, optionally wherein the ELP component comprises 18 repeats, 36 repeats, 72 repeats or 144 repeats of VPGXG (SEQ ID NO: 3).
9. The therapeutic agent for the use according to claim 5, wherein each X is V, optionally wherein the ELP component comprises 5 repeats, 60 repeats or 120 repeats of VPGXG (SEQ ID NO: 3).
- 10 10. The therapeutic agent for the use according to claim 4, wherein the repeating unit is AVGVVP (SEQ ID NO: 4), optionally wherein the ELP component comprises 5 repeats, 10 repeats, 30 repeats, 60 repeats or 120 repeats of AVGVVP (SEQ ID NO: 4).
- 15 11. The therapeutic agent for the use according to claim 4, wherein the repeating unit is IPGXG (SEQ ID NO: 5), wherein X is V, optionally wherein the ELP component comprises 5 repeats, 10 repeats or 15 repeats of IPGXG (SEQ ID NO: 5).
12. The therapeutic agent for the use according to claim 4, wherein the repeating unit is LPGXG (SEQ ID NO: 7), wherein X is V, optionally wherein the ELP component comprises 5 repeats, 10 repeats, 30 repeats, 60 repeats or 120 repeats of LPGXG (SEQ ID NO: 7).
- 20 13. The therapeutic agent for the use according to any one of the preceding claims, wherein the ELP has a Tt of from about 35°C to about 60°C, optionally wherein the Tt is about 35°C.
- 25 14. The therapeutic agent for the use according to any one of the preceding claims, wherein, in the method, the therapeutic agent is administered with a pharmaceutically acceptable carrier and/or excipient.
15. A therapeutic agent for use in a method of treating diabetes, the method comprising administering the therapeutic agent to a subject,
30 wherein said therapeutic agent is a fusion protein of an insulin A chain and an elastin-like peptide (ELP) component comprising 120 repeats of VPGXG (SEQ ID NO: 3), where each X is V, G, or A at a ratio of 5:3:2, wherein the insulin A chain amino acid sequence is positioned at the N-terminal side of the ELP component.

FIGURE 1

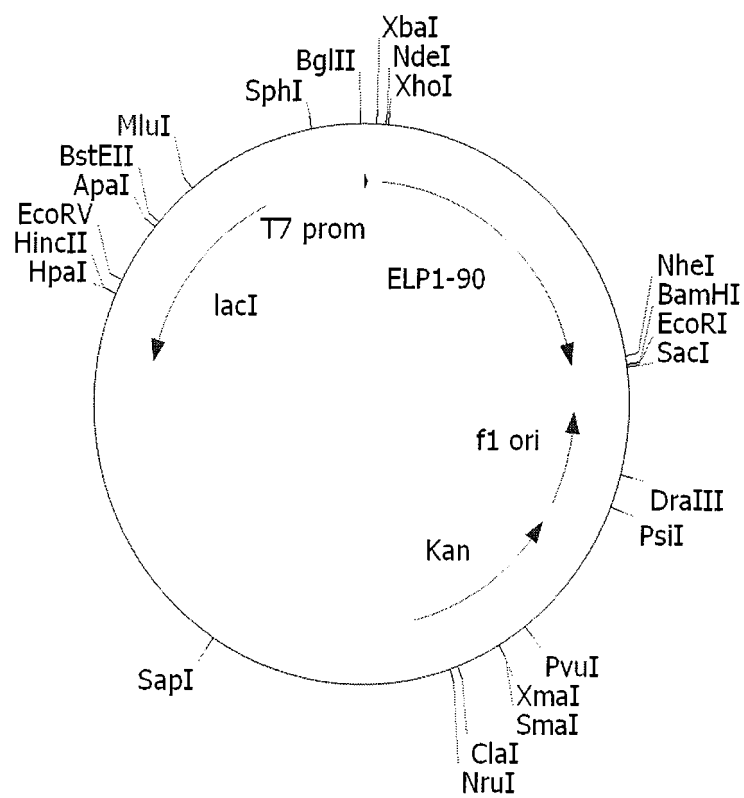


FIGURE 2A

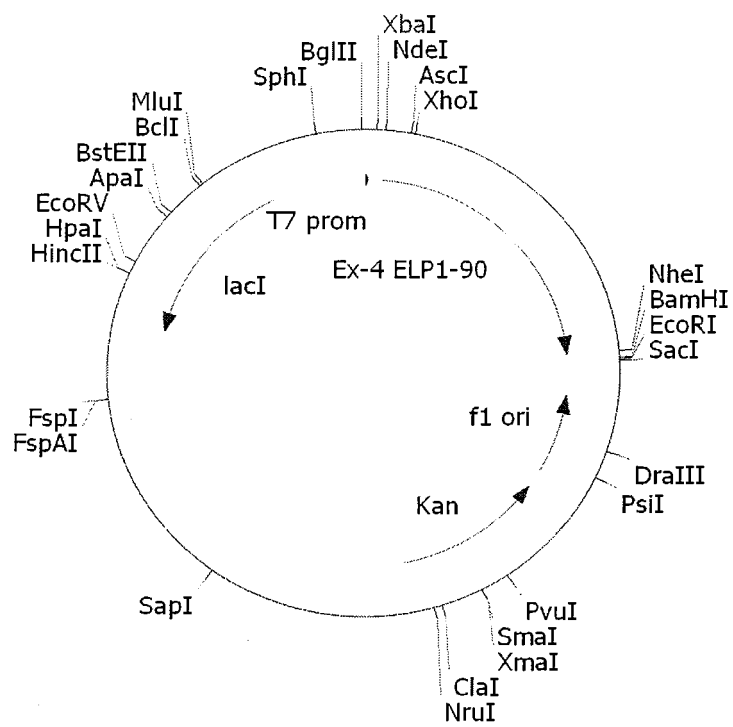


FIGURE 2B

```

1      aattaatacg actcactata ggggaattgt gagcggataa caattccct
      ttaattatgc tgagtgatat cccottaaca ctgcctatt gttaagggga
      >>...T7 prom....>>

      XbaI                      NdeI
      -+-----                ----+---
                                  P1-----
51     ctagaaataa ttttgtttaa ctttaagaag gagatataca tatgcatggc
      gatctttatt aaaacaaatt gaaattcttc ctctatatgt atacgtaccg
                                  <-----
                                  m h g
                                  > Ex-4>

      ----->P2-----
101    gaaggcaoct ttaccagcga tctgagcaaa cagatggaag aagaagcggc
      ottcogtgga aatggtcgct agaactggtt gtctacottc ttcttcgcca
      -----P6<-----
      >.....Ex-4.....>
      e g t f t s d l s k q m e e e a

                                  AscI
                                  ---+-----
      ----->P3-----
151    gcgcctgttt attgaatggc tgaaaaacgg cggcccgcgc agcggcgcgc
      cgcggaacaaa taacttacgc actttttgcc gccgggctcg tcgccgcgcg
      -----P5<-----
      >.....Ex-4.....>
      v r l f i e w l k n g g p s s g a

      XhoI
      -+-----
      ----->
201    cgcgcgcgag cctcgagggc atgggtgggc cgggcgtggg tgttcggggc
      gcggcggctc ggagctcccg taccaccccg gccgcaccc acaaggcccg
      -----P4
      >.....>.....>ELP1-90>
      p p p s l e g m g g p g v g v p g

251    gtgggtgttc cgggtggcgg tgtgccgggc gcaggtgttc ctggtgtagg (SEQ ID NO.23)
      caaccacaag gccaccgcc acacggcccg cgtccacaag gaccacatcc
      >.....ELP1-90.....>
      v g v p g g g v p g a g v p g v (SEQ ID NO.24)

P1: SEQ ID NO.: 35
P2: SEQ ID NO.: 36
P3: SEQ ID NO.: 37
P4: SEQ ID NO.: 38
P5: SEQ ID NO.: 39
P6: SEQ ID NO.: 40

```

FIGURE 3A

```

NdeI
-----
P7-----
1  tatggaaaac ctgtatttcc aacatggcga aggcaccttt accagcgatc
   accttttg gacataaagg ttgtaccgct tccgtggaaa tggtcgctag
   <-----P8
   >>.....Ex-4 ELP1-90.....>
     m e n l y f q h g e g t f t s d
     >>.....Tev.....>>

>
51  tgagcaaaca gatggaagaa gaagcgggtgc gcctgtttat tgaatggctg
   actogtttgt ctaccttctt cttcgccacg cggacaaaata acttaccgac
   >.....Ex-4 ELP1-90'.....>
     l s k q m e e e a v r l f i e w l

                               AscI
                               --+-----
101  aaaaacggcg gcccgagcag cggcgcgccg ccgccgagcc (SEQ ID NO.: 25)
     tttttgccgc cgggctogtc gccgcgcggc ggcggctcgg agct
     <-----P4
     >.....Ex-4 ELP1-90.....>>
       k n g g p s s g a p p p s (SEQ ID NO.: 26)

P4: SEQ ID NO.: 38
P7: SEQ ID NO.: 41
P8: SEQ ID NO.: 42

```

FIGURE 3B

```

NdeI
----
P9-----
1  tatggatata ccaacgaccg aaaacctgta tttccaacat ggccaaggca
   acctatag ggttgctggc ttttgacat aaagggttgta ccgcttccgt
   <-----
   >>.....Ex-4 ELP1-90.....>
     m d i p t t e n l y f q h g e g
   >>.....Linker Tev.....>>

----->
51  cctttaccag cgatctgagc aaacagatgg aagaagaagc ggtgcgcctg
   ggaaatggtc gctagactcg tttgtctacc ttcttcttcg ccacgcggac
   -----P10
   >.....Ex-4 ELP1-90.....>
     t f t s d l s k q m e e e a v r l

                                           AscI
                                           --+-----
101 tttattgaat ggctgaaaaa cggcgggcccg agcagcggcg cgccgccgcc
   aaataactta ccgacttttt gccgcggggc tcgtcgccgc gcggcgggcg
   <-----
   >.....Ex-4 ELP1-90.....>
     f i e w l k n g g p s s g a p p

151 gagcc (SEQ ID NO.: 27)
   ctcggagct
   -----P4
   >..>> Ex-4 ELP1-90
   p s (SEQ ID NO.: 28)

P4: SEQ ID NO.: 38
P9: SEQ ID NO.: 43
P10: SEQ ID NO.: 44

```

FIGURE 4A

```

NdeI
----
P11-----
1  tatgaaaaag atttggctgg cgctggctgg tttagtttta gcgttttagcg
   acttttttc taaaccgacc gcgaccgacc aaatcaaaat cgcaaatcgc
   <-----
>>.....DsbA.....>
   m k k i w l a l a g l v l a f s
   ----->
51  catcggcgca tggcgaaggc acctttacca gcgatctgag caaacagatg
   gtagccgcgt accgcttccg tggaaatggc cgctagactc gtttgctctac
   -----P12
>.DsbA>>>>.....Ex-4 ELP1-90.....>
   a s a h g e g t f t s d l s k q m
   -----
101 gaagaagaag cggcgcgcct gtttattgaa tggctgaaaa acggcgccccc
   cttcttcttc gccacgcgga caaataactt accgactttt tgccgcgcggg
   <-----
>.....Ex-4 ELP1-90.....>
   e e e a v r l f i e w l k n g g
   -----
           AscI
           --+-----
151  gagcagcggc gcgccgccgc cgagcc (SEQ ID NO.: 29)
   ctcgtcgccg cgcggcggcg gctcggagct
   -----P4
>.....Ex-4 ELP1-90.....>>
   p s s g a p p p s (SEQ ID NO.: 30)

P4: SEQ ID NO.: 38
P11: SEQ ID NO.: 45
P12: SEQ ID NO.: 46

```

FIGURE 4B

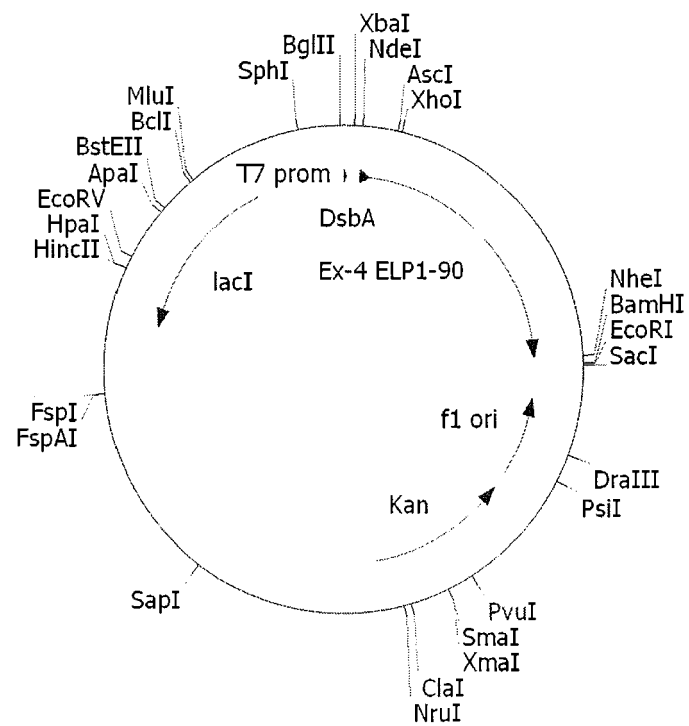


FIGURE 5A

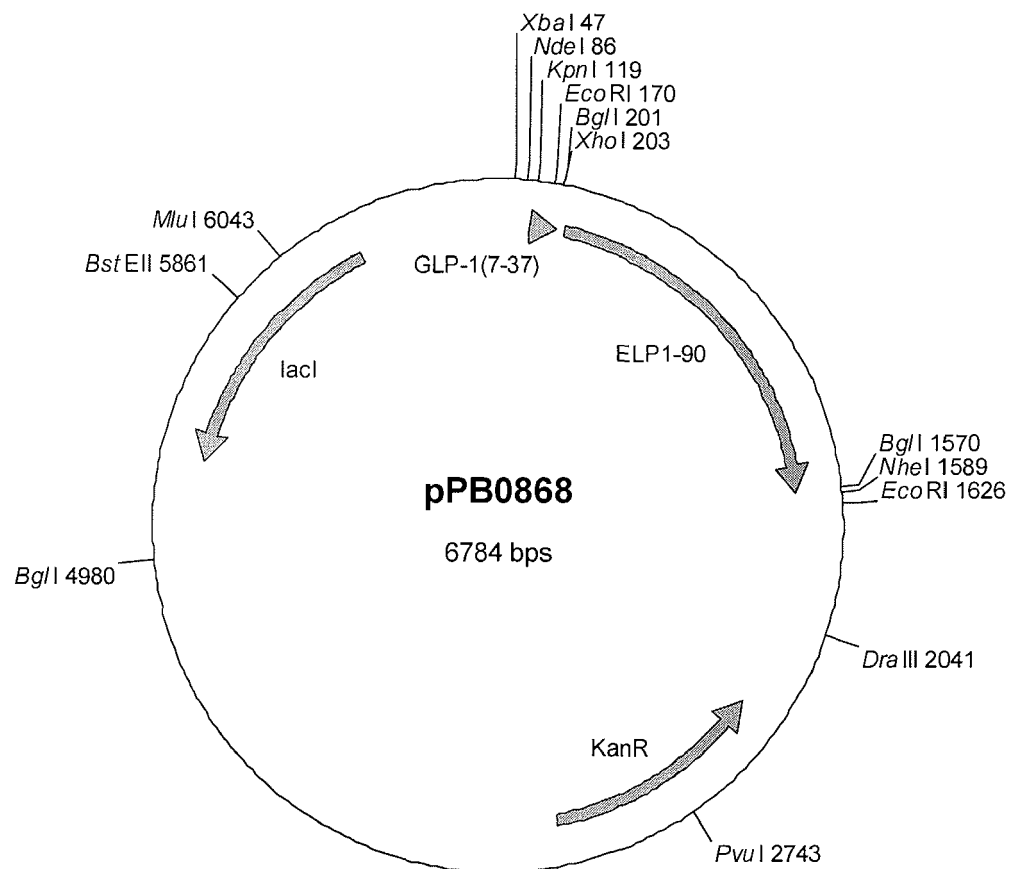


FIGURE 5B

```

                                XbaI
                                --+----
1  taatacgaact cactataggg gaattgtgag cggataacaa ttccctota

                                NdeI
                                ---+----
51 gaaataattt tgtttaactt taagaaggag atatacatat ggagaacotg
                                m e n l

                                KpnI
                                -----+
101 tatttccaac atggcgaagg tacctttaca agcgatgtta gttcatatct
    >>>>.....GLP-1(7-37).....>
    y f q h g e g t f t s d v s s y

151 ggagggccag gcggaagg aattcattgc gtggctggtg aaaggccgcg
    >.....GLP-1(7-37).....>
    l e g q a a k e f i a w l v k g r

                                XhoI
                                -+----
201 gcctcgaggg catgggtggg ccgggcgtgg gtgttcoggg cgtgggtggt
    >>.....ELP1-90.....>
    g l e g m g g p g v g v p g v g v

251 ccgggtggcg gtgtgcoggg cgcaggtggt cctggtgtag gtgtgcoggg
    >.....ELP1-90.....>
    p g g g v p g a g v p g v g v p

301 tgttggtgtg ccgggtggtg gtgtaccagg tggcgggtgtt ccgggtgcag
    >.....ELP1-90.....>
    g v g v p g v g v p g g g v p g a

351 gcgttcoggg tggcgggtgtg ccgggcgtgg gtgttcoggg cgtgggtggt
    >.....ELP1-90.....>
    g v p g g g v p g v g v p g v g v

401 ccgggtggcg gtgtgcoggg cgcaggtggt cctggtgtag gtgtgcoggg
    >.....ELP1-90.....>
    p g g g v p g a g v p g v g v p

451 tgttggtgtg ccgggtggtg gtgtaccagg tggcgggtgtt ccgggtgcag
    >.....ELP1-90.....>
    g v g v p g v g v p g g g v p g a

501 gcgttcoggg tggcgggtgtg ccgggcgtgg gtgttcoggg cgtgggtggt
    >.....ELP1-90.....>
    g v p g g g v p g v g v p g v g v

551 ccgggtggcg gtgtgcoggg cgcaggtggt cctggtgtag gtgtgcoggg
    >.....ELP1-90.....>
    p g g g v p g a g v p g v g v p

601 tgttggtgtg ccgggtggtg gtgtaccagg tggcgggtgtt ccgggtgcag
    >.....ELP1-90.....>
    g v g v p g v g v p g g g v p g a

```

651 gcgttcacggg tggcgggtgtg ccgggcgtgg gtgttcacggg cgtgggtgtt
 >.....ELP1-90.....>
 g v p g g g v p g v g v p g v g v

701 ccgggtggcg gtgtgccggg cgcaggtgtt cctggtgtag gtgtgccggg
 >.....ELP1-90.....>
 p g g g v p g a g v p g v g v p

751 tgttggtgtg ccgggtgttg gtgtaccagg tggcgggtgtt ccgggtgcag
 >.....ELP1-90.....>
 g v g v p g v g v p g g g v p g a

801 gcgttcacggg tggcgggtgtg ccgggcgtgg gtgttcacggg cgtgggtgtt
 >.....ELP1-90.....>
 g v p g g g v p g v g v p g v g v

851 ccgggtggcg gtgtgccggg cgcaggtgtt cctggtgtag gtgtgccggg
 >.....ELP1-90.....>
 p g g g v p g a g v p g v g v p

901 tgttggtgtg ccgggtgttg gtgtaccagg tggcgggtgtt ccgggtgcag
 >.....ELP1-90.....>
 g v g v p g v g v p g g g v p g a

951 gcgttcacggg tggcgggtgtg ccgggcgtgg gtgttcacggg cgtgggtgtt
 >.....ELP1-90.....>
 g v p g g g v p g v g v p g v g v

1001 ccgggtggcg gtgtgccggg cgcaggtgtt cctggtgtag gtgtgccggg
 >.....ELP1-90.....>
 p g g g v p g a g v p g v g v p

1051 tgttggtgtg ccgggtgttg gtgtaccagg tggcgggtgtt ccgggtgcag
 >.....ELP1-90.....>
 g v g v p g v g v p g g g v p g a

1101 gcgttcacggg tggcgggtgtg ccgggcgtgg gtgttcacggg cgtgggtgtt
 >.....ELP1-90.....>
 g v p g g g v p g v g v p g v g v

1151 ccgggtggcg gtgtgccggg cgcaggtgtt cctggtgtag gtgtgccggg
 >.....ELP1-90.....>
 p g g g v p g a g v p g v g v p

1201 tgttggtgtg ccgggtgttg gtgtaccagg tggcgggtgtt ccgggtgcag
 >.....ELP1-90.....>
 g v g v p g v g v p g g g v p g a

1251 gcgttcacggg tggcgggtgtg ccgggcgtgg gtgttcacggg cgtgggtgtt
 >.....ELP1-90.....>
 g v p g g g v p g v g v p g v g v

1301 ccgggtggcg gtgtgccggg cgcaggtgtt cctggtgtag gtgtgccggg
 >.....ELP1-90.....>
 p g g g v p g a g v p g v g v p

1351 tgttggtgtg ccgggtgttg gtgtaccagg tggcgggtgtt ccgggtgcag
 >.....ELP1-90.....>
 g v g v p g v g v p g g g v p g a

1401 gcgttccggg tggcgggtgtg ccgggcgtgg gtgttccggg cgtgggtgtt
 >.....ELP1-90.....>
 g v p g g g v p g v g v p g v g v

1451 ccgggtggcg gtgtgccggg cgcaggtgtt cctgggttag gtgtgccggg
 >.....ELP1-90.....>
 p g g g v p g a g v p g v g v p

1501 tgttgggtgtg ccgggtgttg gtgtaccagg tggcgggtgtt ccgggtgcag
 >.....ELP1-90.....>
 g v g v p g v g v p g g g v p g a

NheI

-+-----

1551 gcgttccggg tggcgggtgtg ccgggcgtgg cgtgataagc tagcatgact
 >.....ELP1-90.....>>
 g v p g g g v p g w p - - (SEQ ID NO: 54)

1601 ggtggacagc aaatgggtcg gatccgaatt cgagctccgt cgagcaccac

1651 caccaccacc actgagatcc ggctgctaac aaagcccgaa aggaagctga (SEQ ID NO: 53)

FIGURE 6A

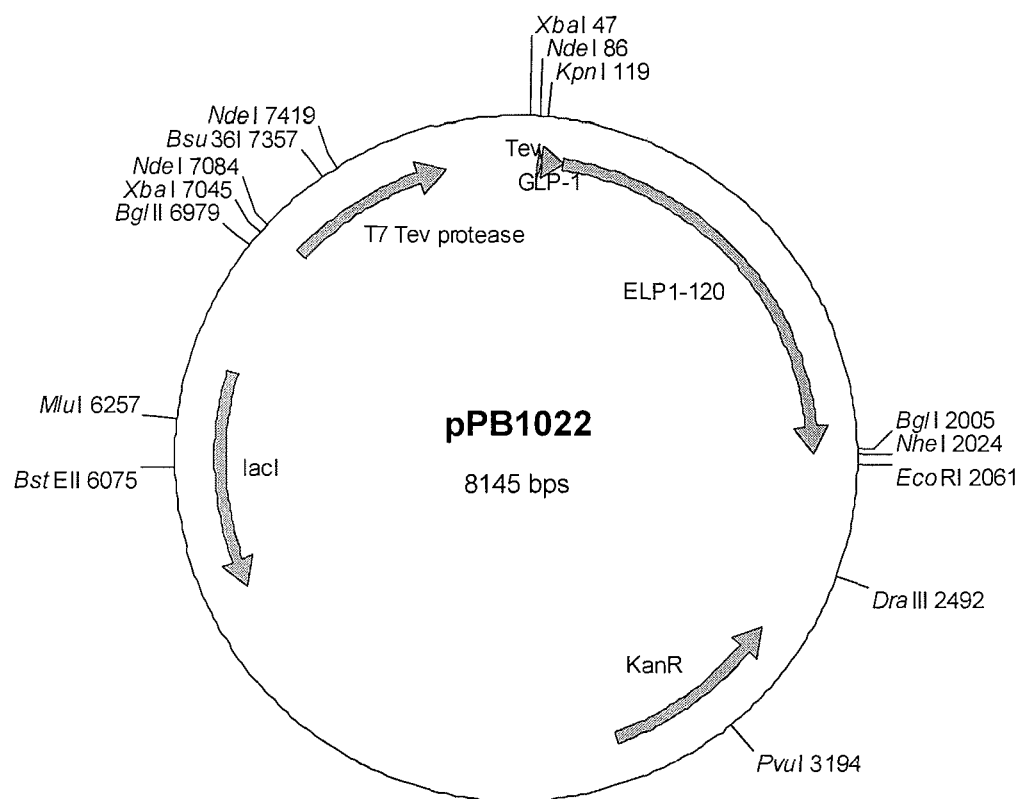


FIGURE 6B

```

                                                    XbaI
                                                    +-+-----
1   taatacgact cactataggg gaattgtgag cggataacaa ttccctcta

                                                    NdeI
                                                    ---+-----
51  gaaataattt tgtttaactt taagaaggag atatacatat ggagaacctg
                                                    >>...Tev....>
                                                    m e n l

                                                    KpnI
                                                    -----+
101 tatttccaac atggcgaagg tacctttaca agcgatgtta gttcatatct
>...Tev.>>
   y f q
   >>.....GLP-1.....>
       h g e g t f t s d v s s y

151 ggagggccag gcggaagg aatttattgc gtggctggtg aaaggccgag
>.....GLP-1.....>
   l e g q a a k e f i a w l v k g r

201 gcgtgccggg cgtgggtgtt ccgggcgtgg gtgttcggg tggcgggtgtg
>> GLP-1
   g
   >>.....ELP1-120.....>
       v p g v g v p g v g v p g g g v

251 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgttg gtgtgccggg
>.....ELP1-120.....>
   p g a g v p g v g v p g v g v p

301 tgttggtgta ccaggtggcg gtgttcggg tgcaggcggt ccgggtggcg
>.....ELP1-120.....>
   g v g v p g g g v p g a g v p g g

351 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcggg tggcgggtgtg
>.....ELP1-120.....>
   g v p g v g v p g v g v p g g g v

401 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgttg gtgtgccggg
>.....ELP1-120.....>
   p g a g v p g v g v p g v g v p

451 tgttggtgta ccaggtggcg gtgttcggg tgcaggcggt ccgggtggcg
>.....ELP1-120.....>
   g v g v p g g g v p g a g v p g g

501 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcggg tggcgggtgtg
>.....ELP1-120.....>
   g v p g v g v p g v g v p g g g v

551 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgttg gtgtgccggg
>.....ELP1-120.....>

```

p g a g v p g v g v p g v g v p
 601 tgttggtgta ccaggtggcg gtgttcggg tgcaggcggt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 651 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 701 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgtt gtgtgccggg
 >.....ELP1-120.....>
 p g a g v p g v g v p g v g v p
 751 tgttggtgta ccaggtggcg gtgttcggg tgcaggcggt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 801 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 851 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgtt gtgtgccggg
 >.....ELP1-120.....>
 p g a g v p g v g v p g v g v p
 901 tgttggtgta ccaggtggcg gtgttcggg tgcaggcggt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 951 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 1001 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgtt gtgtgccggg
 >.....ELP1-120.....>
 p g a g v p g v g v p g v g v p
 1051 tgttggtgta ccaggtggcg gtgttcggg tgcaggcggt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 1101 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 1151 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgtt gtgtgccggg
 >.....ELP1-120.....>
 p g a g v p g v g v p g v g v p
 1201 tgttggtgta ccaggtggcg gtgttcggg tgcaggcggt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 1251 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 1301 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgtt gtgtgccggg
 >.....ELP1-120.....>

p g a g v p g v g v p g v g v p
 1351 tgttggtgta ccaggtggcg gtgttcoggg tgcaggcggtt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 1401 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcoggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 1451 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgttg gtgtgccggg
 >.....ELP1-120.....>
 p g a g v p g v g v p g v g v p
 1501 tgttggtgta ccaggtggcg gtgttcoggg tgcaggcggtt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 1551 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcoggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 1601 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgttg gtgtgccggg
 >.....ELP1-120.....>
 p g a g v p g v g v p g v g v p
 1651 tgttggtgta ccaggtggcg gtgttcoggg tgcaggcggtt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 1701 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcoggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 1751 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgttg gtgtgccggg
 >.....ELP1-120.....>
 p g a g v p g v g v p g v g v p
 1801 tgttggtgta ccaggtggcg gtgttcoggg tgcaggcggtt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 1851 gtgtgccggg cgtgggtgtt ccgggcgtgg gtgttcoggg tggcgggtgtg
 >.....ELP1-120.....>
 g v p g v g v p g v g v p g g g v
 1901 ccgggcgcag gtgttcctgg ttaggtgtg ccgggtgttg gtgtgccggg
 >.....ELP1-120.....>
 p g a g v p g v g v p g v g v p
 1951 tgttggtgta ccaggtggcg gtgttcoggg tgcaggcggtt ccgggtggcg
 >.....ELP1-120.....>
 g v g v p g g g v p g a g v p g g
 BglI NheI
 -----+-----
 2001 gtgtgccggg ctggccgtga taagctagca tgactggtgg acagcaaatg (SEQ ID NO:55)
 >.....ELP1-120.....>>
 g v p g w p - - (SEQ ID NO: 56)

FIGURE 7A

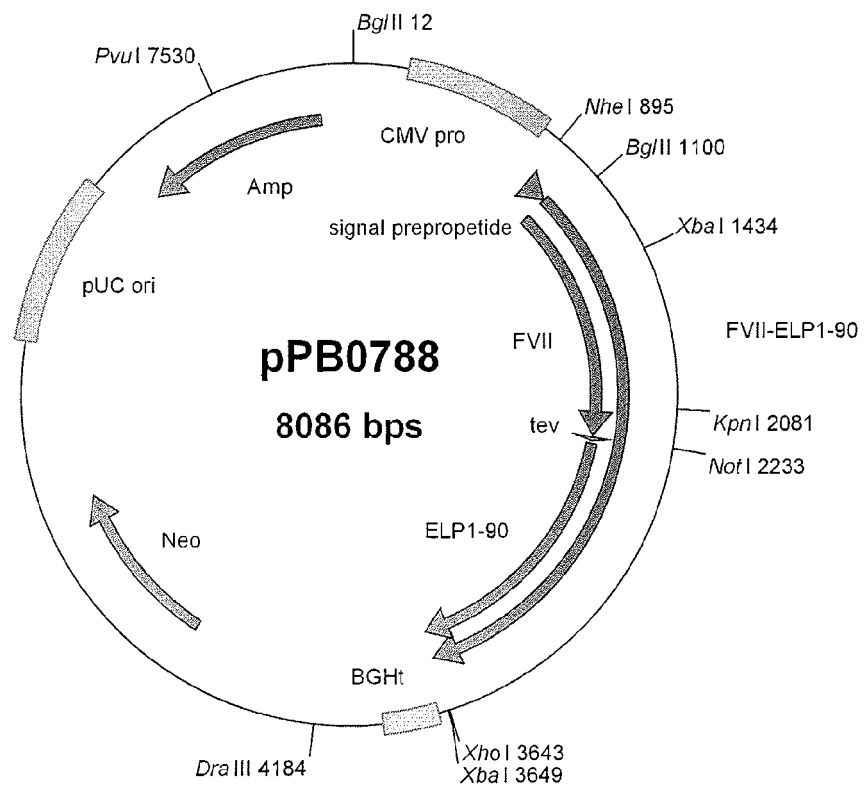


FIGURE 7B

BglII
-+----

1 gacggatcgg gagatctccc gatccctat ggtgcaactct cagtacaatc
51 tgctctgatg ccgcatagtt aagccagtat ctgctccctg cttgtgtgtt
101 ggaggtoget gagtagtgcg cgagcaaaat ttaagctaca acaaggcaag
151 gcttgaccga caattgcatg aagaatctgc ttaggggttag gcgttttgcg
201 ctgcttgcg atgtaogggc cagatatacg cgttgacatt gattattgac
251 tagttattaa tagtaatcaa ttacgggggtc attagttoat agcccatata
301 tggagttccg cgttacataa cttacggtaa atggcccgcc tggtgaccg
351 cccaacgacc ccgcccatt gacgtcaata atgacgtatg tcccatagt
401 aacgccaata gggactttcc attgacgtca atgggtggag tatttaoggt
451 aaactgocca cttggcagta catcaagtgt atcatatgcc aagtaogccc
501 cctattgacg tcaatgacgg taaatggccc gcctggcatt atgccagta
551 catgacctta tgggactttc ctacttggca gtacatctac gtattagtca
601 tcgctattac catggtgatg cggttttggc agtacatcaa tgggcgtgga
651 tagcggtttg actcaogggg atttccaagt ctccacocca ttgacgtcaa
701 tgggagtttg ttttggcacc aaaatcaacg ggactttoca aaatgtogta
751 acaactccgc ccattgacg caaatgggcg gtaggcgtgt acggtgggag
801 gtctatataa gcagagctct ctggctaact agagaacca ctgcttactg

NheI
-+----

851 gcttatogaa attaatacga ctcactatag ggagacocaa gctggctage
901 atgggtctccc aggcctcag gtcctctgc cttctgcttg gcttcaggg
>>.....signal prepropeptide.....>
m v s q a l r l l c l l l g l q
951 ctgcctggct gcagtcttcg taaccagga ggaagccac gcgctctgc
>.....signal prepropeptide.....>
g c l a a v f v t q e e a h g v l
1001 accggcgccg gcgcgccaac gcgttctgg aggagctacg gcgggctcc
>.....>>>.....FVII-ELP1-90.....>
h r r r r a n a f l e e l r p g s
>>.....FVII.....>
1051 ctggagaggg agtgaagga ggagcagtc tccttogagg aggccggga
>.....FVII-ELP1-90.....>
l e r e c k e e q c s f e e a r

```

>.....FVII.....>

BglII
--+----
1101 gatcttcaag gacgcggaga ggacgaagct gttctggatt tcttacagt
>.....FVII-ELP1-90.....>
   e i f k d a e r t k l f w i s y s
>.....FVII.....>

1151 atggggacca gtgtgcctca agtccatgcc agaatggggg ctctgcaag
>.....FVII-ELP1-90.....>
   d g d q c a s s p c q n g g s c k
>.....FVII.....>

1201 gaccagctcc agtcctatat ctgcttctgc ctccctgcct tcgagggcgc
>.....FVII-ELP1-90.....>
   d q l q s y i c f c l p a f e g
>.....FVII.....>

1251 gaactgtgag acgcacaagg atgaccagct gatctgtgtg aacgagaacg
>.....FVII-ELP1-90.....>
   r n c e t h k d d q l i c v n e n
>.....FVII.....>

1301 gcggctgtga gcagtactgc agtgaccaca cgggcaccaa gcgctcctgt
>.....FVII-ELP1-90.....>
   g g c e q y c s d h t g t k r s c
>.....FVII.....>

1351 cgggtgccaag aggggtactc tctgctggca gacggggtgt cctgcacacc
>.....FVII-ELP1-90.....>
   r c h e g y s l l a d g v s c t
>.....FVII.....>

XbaI
--+----
1401 cacagttgaa tatccatgtg gaaaaatacc tattctagaa aaaagaaatg
>.....FVII-ELP1-90.....>
   p t v e y p c g k i p i l e k r n
>.....FVII.....>

1451 ccagcaaacc ccaaggccga attgtggggg gcaaggtgtg ccccaaaggg
>.....FVII-ELP1-90.....>
   a s k p q g r i v g g k v c p k g
>.....FVII.....>

1501 gagtgtocat ggcaggtcct gttgttggtg aatggagctc agttgtgtgg
>.....FVII-ELP1-90.....>
   e c p w q v l l l v n g a q l c
>.....FVII.....>

1551 ggggacctg atcaacacca tctgggtggt ctccgcggcc cactgtttcg
>.....FVII-ELP1-90.....>
   g g t l i n t i w v v s a a h c f
>.....FVII.....>

1601 acaaaatcaa gaactggagg aacctgatcg cgggtctggg cgagcacgac
>.....FVII-ELP1-90.....>
   d k i k n w r n l i a v l g e h d
>.....FVII.....>

```

```

1651 ctcagcagagc acgacgggga tgagcagagc cggcgggtgg cgcagggtcat
>.....FVII-ELP1-90.....>
   l s e h d g d e q s r r v a q v
>.....FVII.....>

1701 catccccagc acgtacgtcc cgggcaccac caaccaogac atcgcgtgc
>.....FVII-ELP1-90.....>
   i i p s t y v p g t t n h d i a l
>.....FVII.....>

1751 tccgcctgca ccagcccggtg gtcctcactg accatgtggt gccctctgc
>.....FVII-ELP1-90.....>
   l r l h q p v v l t d h v v p l c
>.....FVII.....>

1801 ctgccogaac ggacgttctc tgagaggacg ctggccttcg tgcgttctc
>.....FVII-ELP1-90.....>
   l p e r t f s e r t l a f v r f
>.....FVII.....>

1851 attggtcagc ggctggggcc agctgctgga ccgtggcgcc acggccctgg
>.....FVII-ELP1-90.....>
   s l v s g w g q l l d r g a t a l
>.....FVII.....>

1901 agctcatggt cctcaacgtg ccccggtga tgaccagga ctgcctgcag
>.....FVII-ELP1-90.....>
   e l m v l n v p r l m t q d c l q
>.....FVII.....>

1951 cagtcacgga aggtgggaga ctccccaat atcacggagt acatgttctg
>.....FVII-ELP1-90.....>
   q s r k v g d s p n i t e y m f
>.....FVII.....>

2001 tgccggctac tcggatggca gcaaggactc ctgcaagggg gacagtggag
>.....FVII-ELP1-90.....>
   c a g y s d g s k d s c k g d s g
>.....FVII.....>

                                     KpnI
                                     -----+
2051 gccacatgc caccactac cggggcacgt ggtacctgac gggcatcgtc
>.....FVII-ELP1-90.....>
   g p h a t h y r g t w y l t g i v
>.....FVII.....>

2101 agctggggcc agggctgcgc aacgtgggc cactttgggg tgtacaccag
>.....FVII-ELP1-90.....>
   s w g q g c a t v g h f g v y t
>.....FVII.....>

2151 ggtctcccag tacatogagt ggctgcaaaa gtcctatgcgc tcagagccac
>.....FVII-ELP1-90.....>
   r v s q y i e w l q k l m r s e p
>.....FVII.....>

                                     NotI
                                     --+-----
2201 gccaggagt cctcctgcga gcccatcttc ccgcggcgcc tgaaaacctg
>.....FVII-ELP1-90.....>

```

```

      r p g v l l r a p f p a a a e n l
>.....FVII.....>>                >>.tev..>

2251  tattttcagg gtggggccgc tgggcccggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
      y f q g g a a g p g v g v p g v
>....tev...>>                >>.....ELP1-90.....>

2301  agttcccgga ggcggagtgc ccggcgcagg agttcctgga gtcggagtgc
>.....FVII-ELP1-90.....>
      g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>

2351  ccggagttgg agtgcccgga gttggagtcc caggaggcgg agtcccggga
>.....FVII-ELP1-90.....>
      p g v g v p g v g v p g g g v p g
>.....ELP1-90.....>

2401  gcaggcgtcc ccggaggcgg agtgcccggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
      a g v p g g g v p g v g v p g v
>.....ELP1-90.....>

2451  agttcccgga ggcggagtgc ccggcgcagg agttcctgga gtcggagtgc
>.....FVII-ELP1-90.....>
      g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>

2501  ccggagttgg agtgcccgga gttggagtcc caggaggcgg agtcccggga
>.....FVII-ELP1-90.....>
      p g v g v p g v g v p g g g v p g
>.....ELP1-90.....>

2551  gcaggcgtcc ccggaggcgg agtgcccggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
      a g v p g g g v p g v g v p g v
>.....ELP1-90.....>

2601  agttcccgga ggcggagtgc ccggcgcagg agttcctgga gtcggagtgc
>.....FVII-ELP1-90.....>
      g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>

2651  ccggagttgg agtgcccgga gttggagtcc caggaggcgg agtcccggga
>.....FVII-ELP1-90.....>
      p g v g v p g v g v p g g g v p g
>.....ELP1-90.....>

2701  gcaggcgtcc ccggaggcgg agtgcccggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
      a g v p g g g v p g v g v p g v
>.....ELP1-90.....>

2751  agttcccgga ggcggagtgc ccggcgcagg agttcctgga gtcggagtgc
>.....FVII-ELP1-90.....>
      g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>

2801  ccggagttgg agtgcccgga gttggagtcc caggaggcgg agtcccggga
>.....FVII-ELP1-90.....>
      p g v g v p g v g v p g g g v p g

```

```

>.....ELP1-90.....>
2851 gcaggcgtcc ccggaggcgg agtgccgggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
    a g v p g g g v p g v g v p g v
>.....ELP1-90.....>
2901 agttcccga ggccgagtgcc ccggcgcagg agttcctgga gtcggagtgcc
>.....FVII-ELP1-90.....>
    g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>
2951 ccggagttgg agtgcccga gttggagttc caggaggcgg agtcccga
>.....FVII-ELP1-90.....>
    p g v g v p g v g v p g g g v p g
>.....ELP1-90.....>
3001 gcaggcgtcc ccggaggcgg agtgccgggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
    a g v p g g g v p g v g v p g v
>.....ELP1-90.....>
3051 agttcccga ggccgagtgcc ccggcgcagg agttcctgga gtcggagtgcc
>.....FVII-ELP1-90.....>
    g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>
3101 ccggagttgg agtgcccga gttggagttc caggaggcgg agtcccga
>.....FVII-ELP1-90.....>
    p g v g v p g v g v p g g g v p g
>.....ELP1-90.....>
3151 gcaggcgtcc ccggaggcgg agtgccgggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
    a g v p g g g v p g v g v p g v
>.....ELP1-90.....>
3201 agttcccga ggccgagtgcc ccggcgcagg agttcctgga gtcggagtgcc
>.....FVII-ELP1-90.....>
    g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>
3251 ccggagttgg agtgcccga gttggagttc caggaggcgg agtcccga
>.....FVII-ELP1-90.....>
    p g v g v p g v g v p g g g v p g
>.....ELP1-90.....>
3301 gcaggcgtcc ccggaggcgg agtgccgggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
    a g v p g g g v p g v g v p g v
>.....ELP1-90.....>
3351 agttcccga ggccgagtgcc ccggcgcagg agttcctgga gtcggagtgcc
>.....FVII-ELP1-90.....>
    g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>
3401 ccggagttgg agtgcccga gttggagttc caggaggcgg agtcccga
>.....FVII-ELP1-90.....>
    p g v g v p g v g v p g g g v p g

```

```

>.....ELP1-90.....>
3451 gcaggcggtcc ccggaggcgg agtgccgggc gtgggagttc ccggcgtggg
>.....FVII-ELP1-90.....>
   a g v p g g g v p g v g v p g v
>.....ELP1-90.....>

3501 agttcccgga ggccggagtgc ccggcgcagg agttcctgga gtccggagtgc
>.....FVII-ELP1-90.....>
   g v p g g g v p g a g v p g v g v
>.....ELP1-90.....>

3551 ccggagttgg agtgcccggg gttggagtcc caggaggcgg agtccccgga
>.....FVII-ELP1-90.....>
   p g v g v p g v g v p g g g v p g
>.....ELP1-90.....>

                                     XbaI
                                     -+-----
                                     XhoI
                                     -+-----
3601 gcaggcggtcc ccggaggcgg agtgccgggc tggccttgat gactcgagtc
>.....FVII-ELP1-90.....>>
   a g v p g g g v p g w p - - (SEQ ID NO: 58)
>.....ELP1-90.....>>

3651 tagagggccccc gtttaaaccg gctgatcagc ctgcactgtg ccttctagtt (SEQ ID NO:57)

```

FIGURE 8A

```

XbaI                                     NdeI
--+-+-----                           -+-----
                                     P15-----
51  ctagaaataa ttttgttttaa ctttaagaag gagatataca tatgtttgtg
    Insulin ELP1-90 >>.....>
                                   m f v
                                   B chain >>...>
----->
101 aaccaacacc tgtgcggctc acacctggtg gaagctctct acctagtgtg
    >.....Insulin ELP1-90.....>
    n q h l c g s h l v e a l y l v
    >.....B chain.....>
151 cggggaacga ggcttcttct acacacccaa gaccgcgcgg gaggcagagg
    >.....Insulin ELP1-90.....>
    c g e r g f f y t p k t r r e a e
    >.....B chain.....>>
                                   C chain >>.....>
201 acctgcaggt ggggcaggtg gagctgggcg ggggccctgg tgcaggcagc
    >.....Insulin ELP1-90.....>
    d l q v g q v e l g g g p g a g s
    >.....C chain.....>
251 ctgcagccct tggccctgga ggggtccctg cagaagcgtg gcattgtgga
    >.....Insulin ELP1-90.....>
    l q p l a l e g s l q k r g i v
    >.....C chain.....>>
                                   >>A chain..>
                                   <-----
301 acaatgctgt accagcatct gctccctcta ccagctggag aactactgca
    cctc ttgatgacgt
    >.....Insulin ELP1-90.....>
    e q c c t s i c s l y q l e n y c
    >.....A chain.....>
    e q c c t s i c s l y q l e n y c
    XhoI
    --+-+-----
351 acctcgaggg catgggtggg cggggcgtgg gtgttcgggg cgtgggtggt
    tggagctccc gtaccc
    -----P16
    >.....Insulin ELP1-90.....>
    n l e g m g g p g v g v p g v g v
    >> A chain
401 cggggtggcg gtgtgccggg cgcaggtggt cctggtgtag gtgtgccggg (SEQ ID NO:31)
    >.....Insulin ELP1-90.....>
    p g g g v p g a g v p g v g v p (SEQ ID NO:32)
P15: SEQ ID NO.: 47
P16: SEQ ID NO.: 48

```

FIGURE 8B

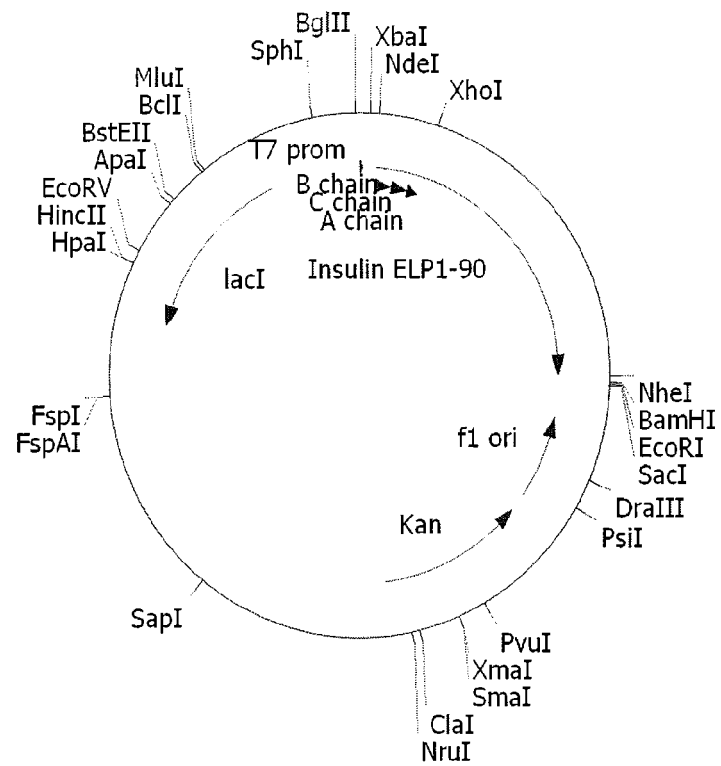


FIGURE 9

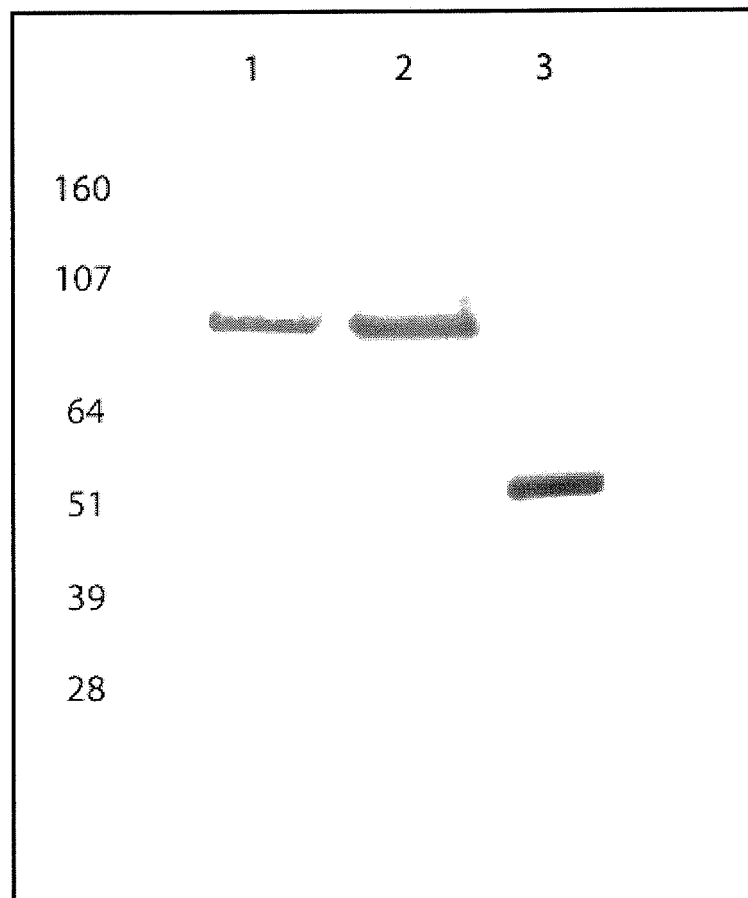


FIGURE 10

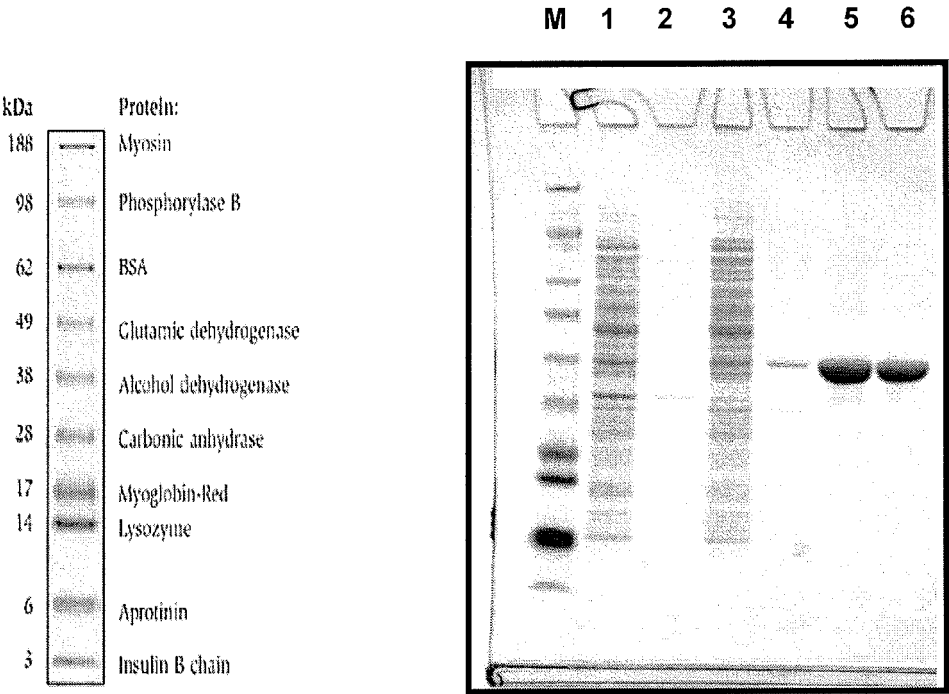


FIGURE 11

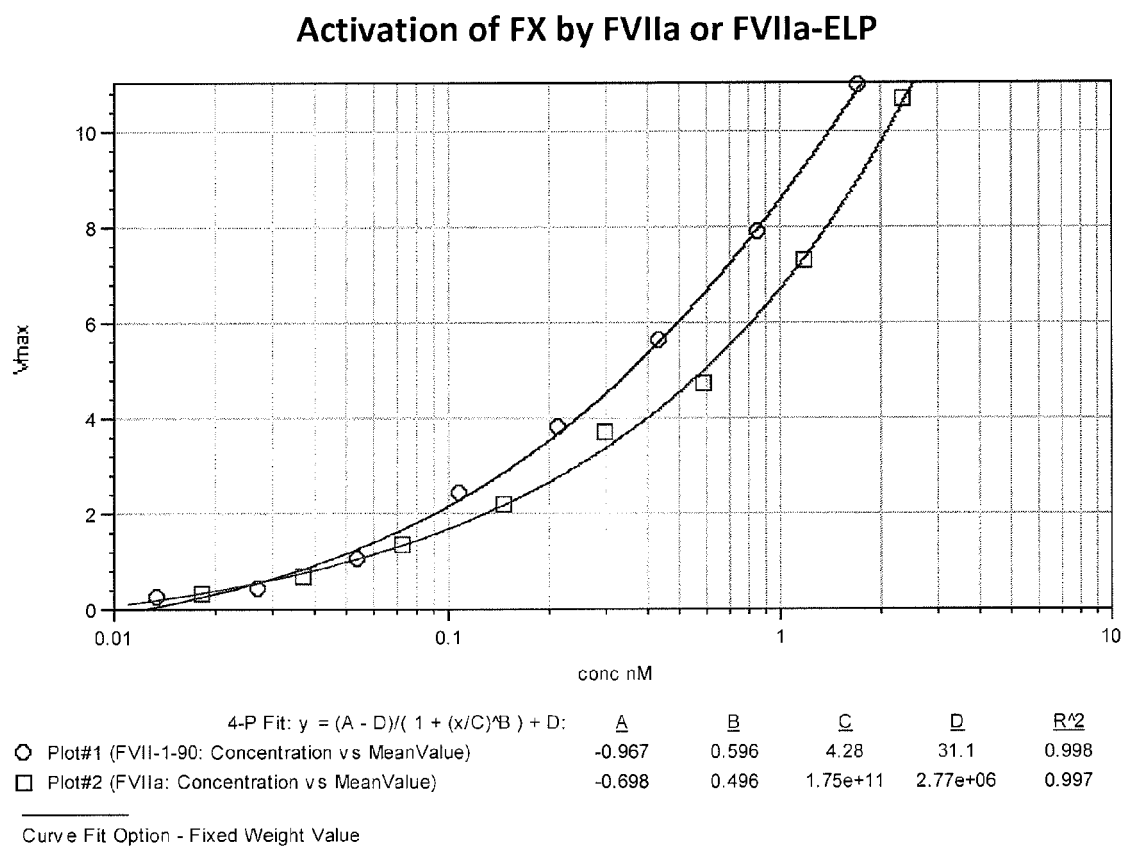


FIGURE 12

FVIIa-ELP has a Long PK in Rats

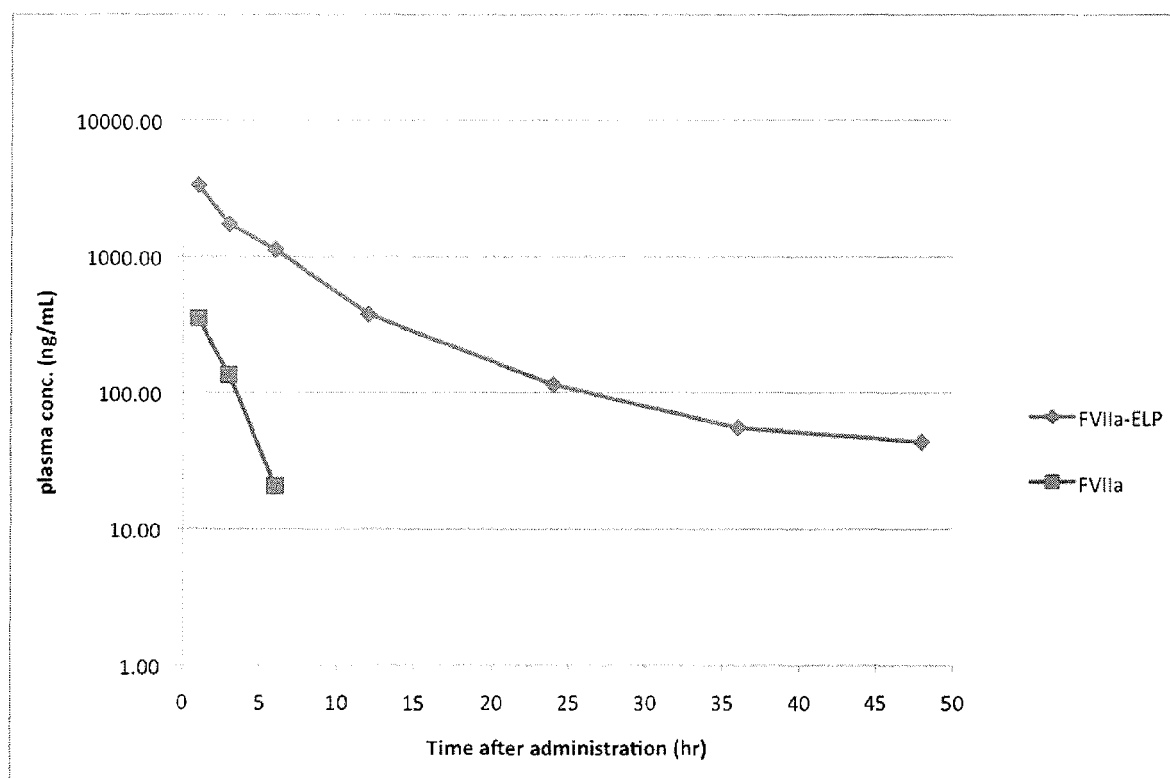
 $T_{1/2}$ FVIIa = 45-60 min $T_{1/2}$ FVIIa-ELP = 690 min

FIGURE 13

High *In vitro* Activity of GLP1-ELP and Exendin-4-ELP Fusion
Proteins

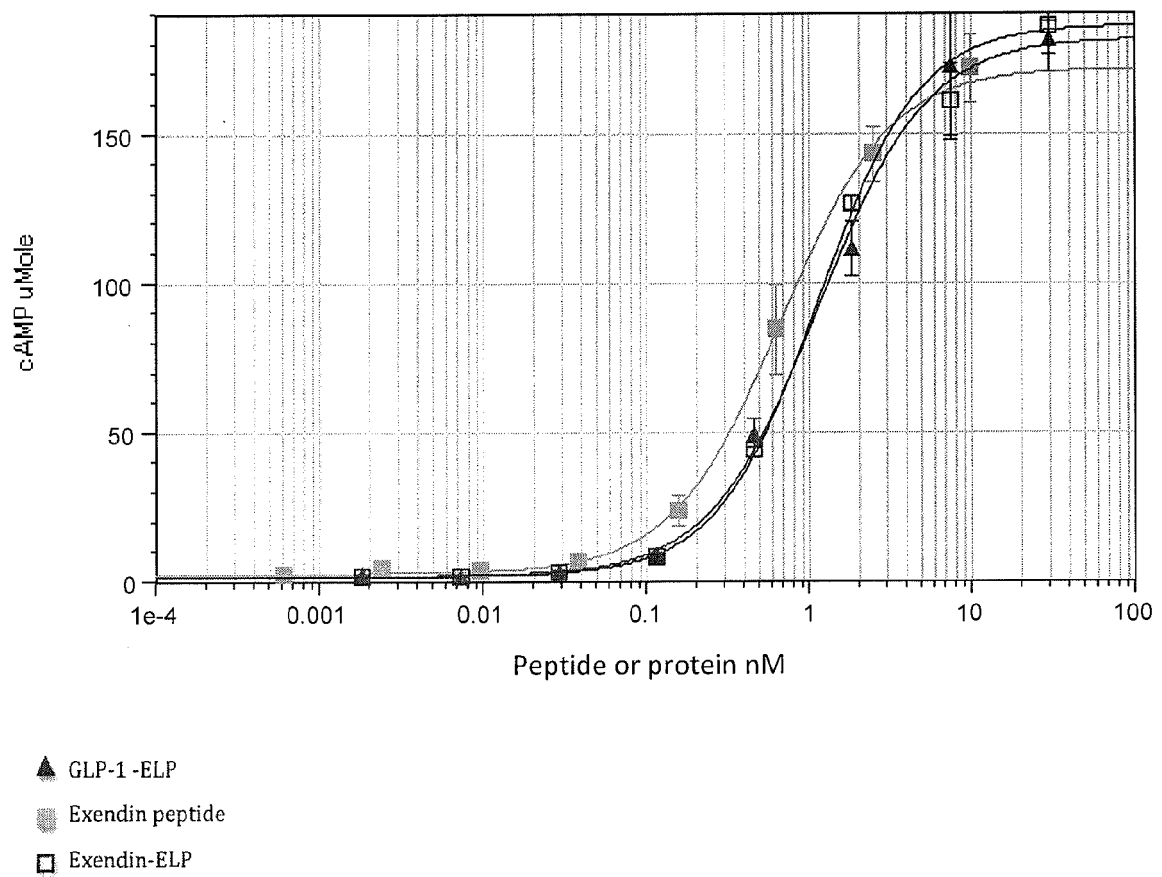


FIGURE 14

Pharmacokinetics of GLP1-ELP in Rats

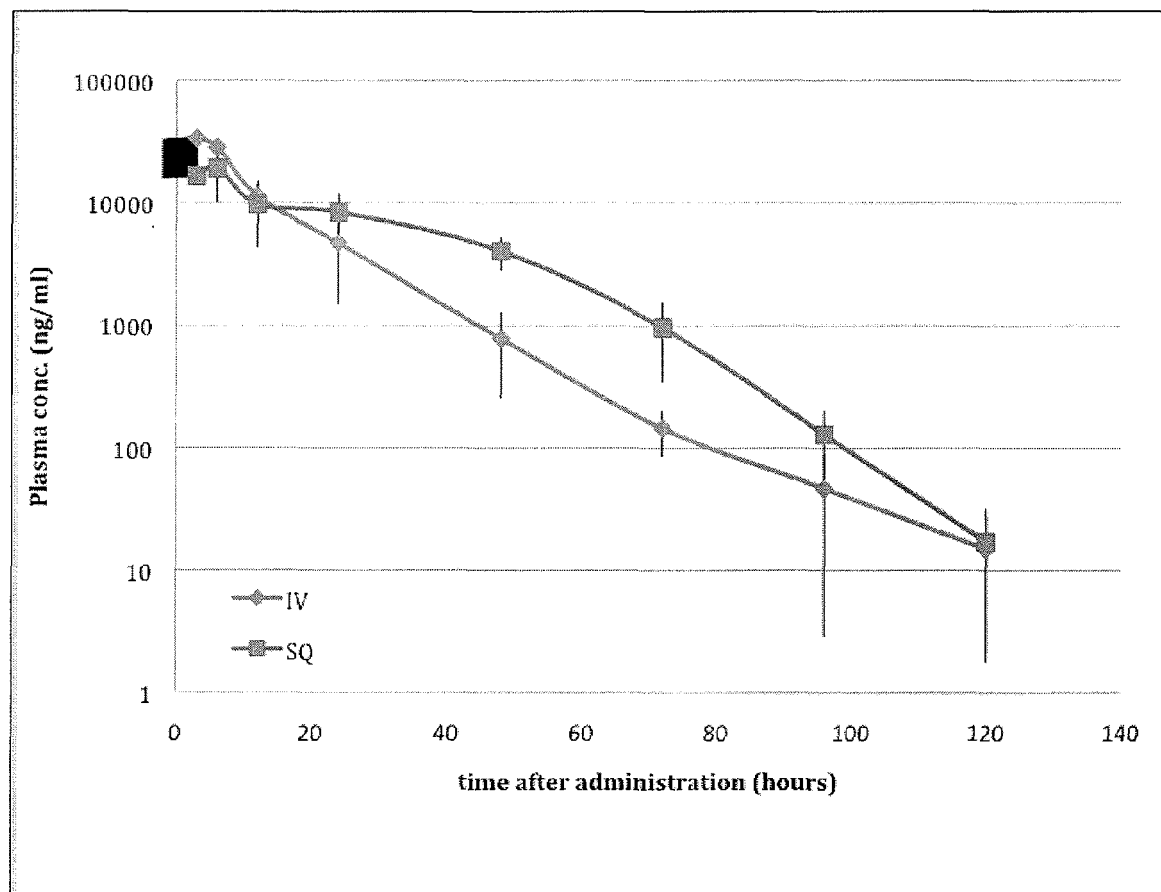
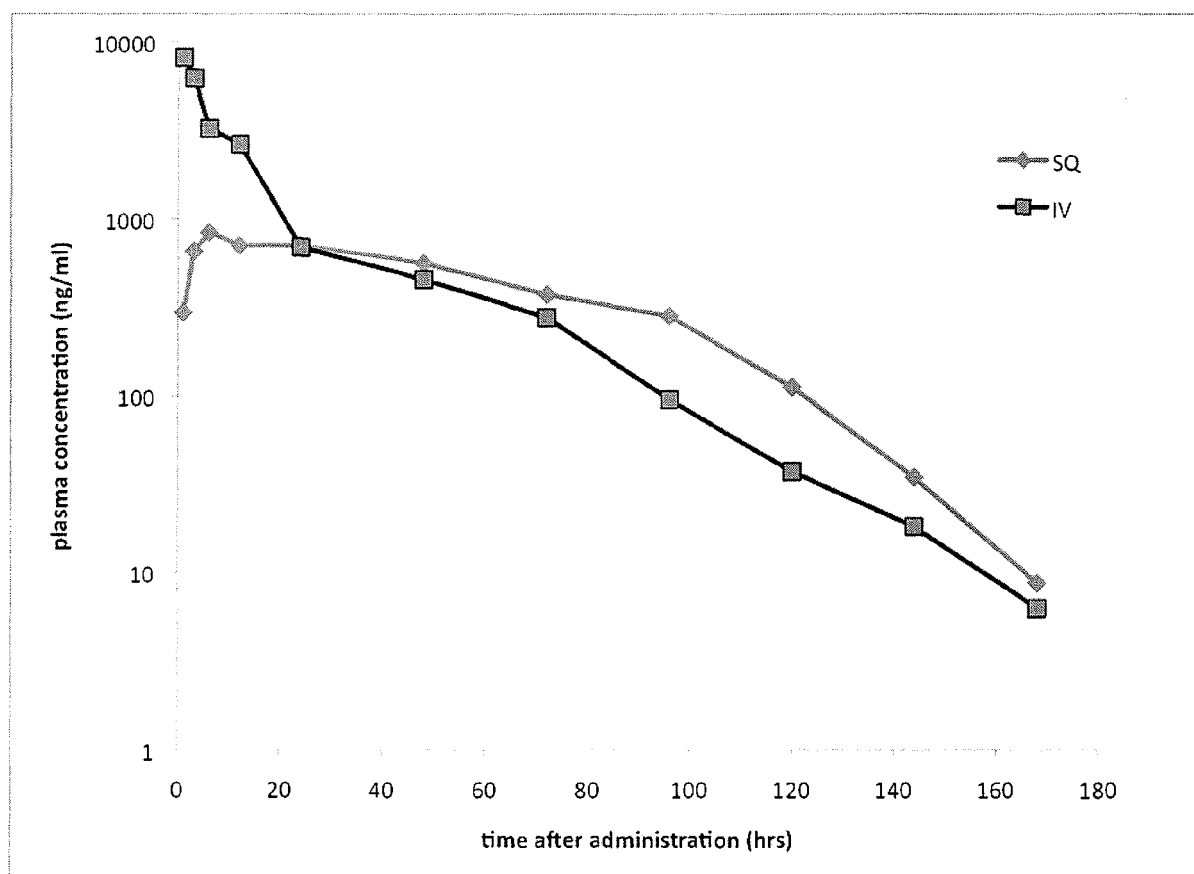
 $T_{1/2}$ IV= 12.9 hours $T_{1/2}$ SQ= 8.6 hours

FIGURE 15

Long Half Life of GLP-1 ELP in Rabbits

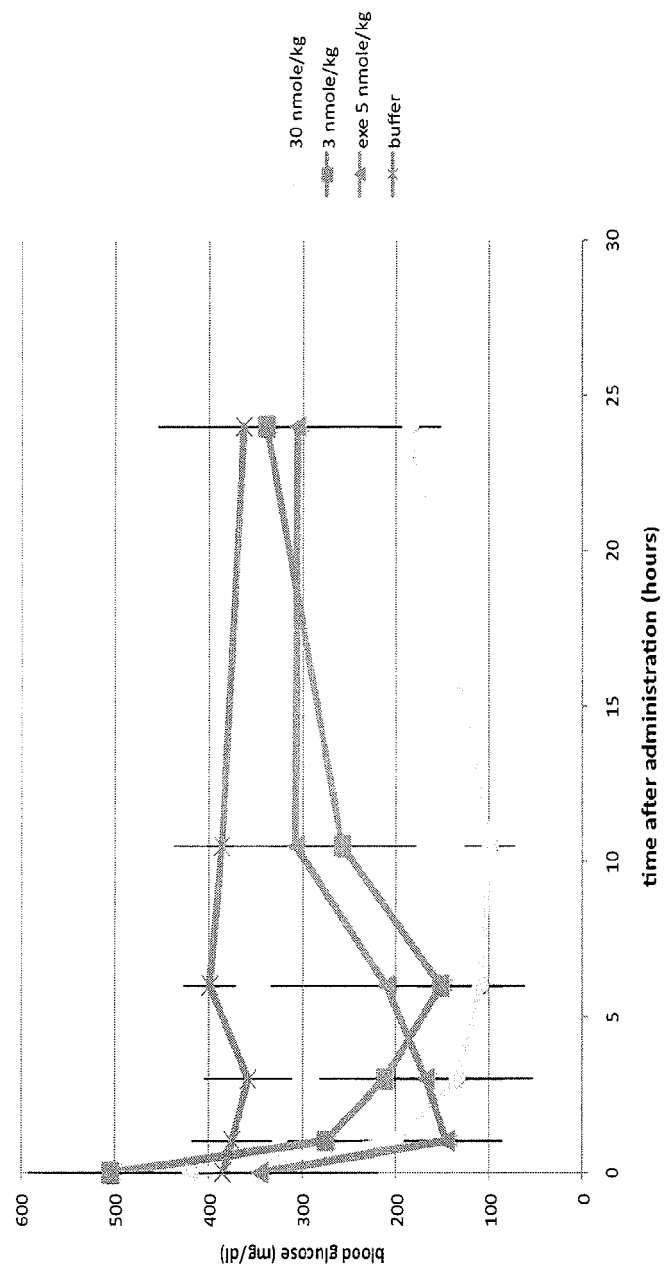


$T_{1/2}$ IV= 12.920 hours

$T_{1/2}$ SQ= 8.624 hours

FIGURE 16

Sustained Glycemic Control of GLP1-ELP in diabetic Mice





EUROPEAN SEARCH REPORT

Application Number
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DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (IPC)
Y	US 2007/009602 A1 (SETTON LORI A [US] ET AL) 11 January 2007 (2007-01-11) * the whole document *	1,4,5,14	INV. A61K38/00 A61K38/28 C07K1/00
Y	WO 2008/030968 A2 (PHASE BIOSCIENCE INC [US]; CHILKOTI ASHUTOSH [US]) 13 March 2008 (2008-03-13) * abstract * * paragraphs [0033], [0034], [0042], [0055], [0068], [0080] - [0082], [0094], [0095], [0103] *	1,4,5,14	A61P3/04 A61P3/08
Y	US 2005/026826 A1 (HOENIG MARGARETHE [US]) 3 February 2005 (2005-02-03) * abstract * * sequences 6,16 * * figure 1 * * claim 33 *	1,4,5,14	
Y	GARY WALSH: "Therapeutic insulins and their large-scale manufacture", APPLIED MICROBIOLOGY AND BIOTECHNOLOGY, SPRINGER, BERLIN, DE, vol. 67, no. 2, 1 April 2005 (2005-04-01), pages 151-159, XP019331799, ISSN: 1432-0614, DOI: 10.1007/S00253-004-1809-X * the whole document *	1,4,5,14	TECHNICAL FIELDS SEARCHED (IPC) C07K A61K A61P
A	WO 2007/100535 A2 (MERCK & CO INC [US]; ANGELETTI P IST RICHERCHE BIO [IT]; ROY RANABIR S) 7 September 2007 (2007-09-07) * abstract * * page 1, column 2 * * page 9, column 3 *	1,4,5,14	
The present search report has been drawn up for all claims			
Place of search Munich		Date of completion of the search 17 June 2015	Examiner Weisser, Dagmar
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	

EPO FORM 1503 03 82 (P04C01)



EUROPEAN SEARCH REPORT

Application Number
EP 15 16 3709

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DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (IPC)
A	HUI HONGXIANG ET AL: "Structure and function studies of glucagon-like peptide-1 (GLP-1): the designing of a novel pharmacological agent for the treatment of diabetes", DIABETES/METABOLISM RESEARCH AND REVIEWS, WILEY, LONDON, GB, vol. 21, no. 4, 1 July 2005 (2005-07-01), pages 313-331, XP002559636, ISSN: 1520-7552, DOI: 10.1002/DMRR.553 [retrieved on 2005-04-26] * abstract * * page 313, columns 2,3 * * page 314, column 5 * * page 317, columns 3,5; table 1 * * page 318; table 2 * * page 319, column 6 * * page 320, columns 1,3; table 3 * * page 321, columns 4,5 * * page 322, column 2 * -----	1,4,5,14	
			TECHNICAL FIELDS SEARCHED (IPC)
The present search report has been drawn up for all claims			
Place of search Munich		Date of completion of the search 17 June 2015	Examiner Weisser, Dagmar
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

EPO FORM 1503 03 82 (P04C01)

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ON EUROPEAN PATENT APPLICATION NO.**

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This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report.
The members are as contained in the European Patent Office EDP file on
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17-06-2015

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2007009602 A1	11-01-2007	CA 2613355 A1	04-01-2007
		CN 101500606 A	05-08-2009
		EP 1896072 A2	12-03-2008
		EP 2664340 A2	20-11-2013
		JP 5253159 B2	31-07-2013
		JP 2009501703 A	22-01-2009
		KR 20080045118 A	22-05-2008
		US 2007009602 A1	11-01-2007
		US 2011236384 A1	29-09-2011
		US 2014364371 A1	11-12-2014
WO 2008030968 A2	13-03-2008	WO 2007002362 A2	04-01-2007
		AU 2007292295 A1	13-03-2008
		CA 2663047 A1	13-03-2008
		CN 101578373 A	11-11-2009
		CN 103230598 A	07-08-2013
		EP 2059606 A2	20-05-2009
		JP 5395664 B2	22-01-2014
		JP 2010502734 A	28-01-2010
		JP 2014051510 A	20-03-2014
		US 2011039776 A1	17-02-2011
US 2005026826 A1	03-02-2005	US 2013143802 A1	06-06-2013
		US 2013178416 A1	11-07-2013
		WO 2008030968 A2	13-03-2008
		NONE	
WO 2007100535 A2	07-09-2007	AU 2007221366 A1	07-09-2007
		CA 2638800 A1	07-09-2007
		CN 101389648 A	18-03-2009
		EP 1991574 A2	19-11-2008
		EP 2471810 A1	04-07-2012
		EP 2471811 A1	04-07-2012
		JP 5297817 B2	25-09-2013
		JP 2009527558 A	30-07-2009
		US 2010144617 A1	10-06-2010
		US 2011183902 A1	28-07-2011
WO 2007100535 A2	07-09-2007	WO 2007100535 A2	07-09-2007

EPO FORM P0459

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REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- US 61076221 B [0001]
- US 2007077767 W [0002] [0047]
- WO 2008030968 A [0002] [0047]
- US 2006048572 W [0002]
- WO 2007073486 A [0002]
- US 7238667 B [0005]
- US 7176278 B [0005]
- US 5766883 A [0005]
- US 9605186 W [0036] [0045]
- US 20070041951 A [0056]
- US 5614492 A [0056]
- US 5545618 A [0056]
- EP 0658568 A1 [0056]
- WO 9325579 A [0056]
- WO 98008871 A [0059] [0061]
- WO 00034331 A [0059] [0064]
- WO 00069911 A [0059] [0066]
- WO 01004156 A [0059] [0068]
- WO 03018516 A [0059] [0070]
- WO 6630438 A [0093]
- WO 08019368 A [0093]
- WO 02077218 A [0111] [0112]
- WO 05074975 A [0111] [0112]
- US 7235638 B [0112]
- US 7176288 B [0113]
- US 6905683 B [0113]

Non-patent literature cited in the description

- KREYMANN et al. *Lancet*, 1987, vol. 2 (8571), 1300-4 [0052]
- WETTERGEN et al. *Dig Dis Sci*, 1993, vol. 38, 665-73 [0052]
- D'ALESSIO et al. *J. Clin Invest*, 1994, vol. 93, 2293-6 [0052]
- GUTNIAK et al. *Diabetes Care*, 1994, vol. 17, 1039-44 [0053]
- WILLIAMS et al. *J. Clin Endo Metab*, 1996, vol. 81, 327-32 [0053]
- HOLZ et al. *Nature*, 1993, vol. 361, 362-5 [0053]
- ORSKOV. *Diabetologia*, 1992, vol. 35, 701-11 [0054]
- WEIR et al. *Diabetes*, 1989, vol. 38, 338-342 [0054]
- GUTNIAK, M. *N. Engl J Med*, 1992, vol. 226, 1316-22 [0054]
- NAUCK et al. *Diabetologia*, 1993, vol. 36, 741-4 [0054]
- CREUTZFELDT et al. *Diabetes Care*, 1996, vol. 19, 580-6 [0054]
- DEACON et al. *Diabetes*, 1995, vol. 44, 1126-31 [0055]
- HARDER et al. *Diabetes Care*, 2004, vol. 27, 1915-21 [0060]
- DEGN et al. *Diabetes*, 2004, vol. 53, 1187-94 [0060]
- 68th American Diabetes Association Meeting, 07 June 2008 [0063]
- KIM et al. *Diabetes*, vol. 52, 751-9 [0065]
- RATNER et al. 65th American Diabetes Association Meeting, 10 June 2005 [0065]
- RATNER et al. 68th American Diabetes Association Meeting [0067]
- JACKSON et al. 65th American Diabetes Association Meeting, 10 June 2005 [0069]
- SAHA et al. *J. Pharm. Exp. Ther.*, 2006, vol. 316, 1159-64 [0069]
- TATUSOVA et al. Blast 2 sequences - a new tool for comparing protein and nucleotide sequences. *FEMS Microbiol Lett.*, 1999, vol. 174, 247-250 [0071] [0095]
- SIEGEL et al. *Regul Pept*, 1999, vol. 79 (2-3), 93-102 [0076]
- FEHMANN et al. *Endocrinology*, 1992, vol. 130, 159-166 [0076]
- BUTEAU et al. *Diabetes*, 2003, vol. 52, 124-32 [0077]
- BULOTTA et al. *C Biochem Biophys*, 2004, vol. 40 (3), 65-78 [0077]
- OWENS et al. *Lancet*, 2001, vol. 358, 739-46 [0086]
- VAJO et al. *Endocr Rev*, 2001, vol. 22, 706-17 [0086]
- DUTTARROY et al. *Diabetes*, 2005, vol. 54, 251-8 [0094]
- JEHLLE et al. *Diabetologia*, 1996, vol. 39, 421-432 [0098]
- HAGEN et al. *PNAS USA*, 1986, vol. 83, 2412-6 [0102]
- MORRISSEY et al. *Cell*, 1987, vol. 50, 129-35 [0104]
- MULLER et al. *Biochem.*, 1994, vol. 33, 10864-70 [0104]
- DRAKE et al. *J. Cell Biol.*, 1989, vol. 109, 389 [0105]
- HEDNER. *Seminars Hematol.*, 2001, vol. 38 (12), 43-7 [0108]
- HEDNER. *Seminars Hematol.*, 2004, vol. 41 (1), 35-9 [0108]

- **ROBERTS et al.** *Blood*, 2004, vol. 104, 3858-64 [0108] [0109]
- **TRANHOLM et al.** *Blood*, 2003, vol. 102 (10), 3615-20 [0111]
- **CHAIING et al.** *Blood*, 1994, vol. 83 (12), 3524-35 [0117]
- **NILSSON et al.** *Ann Rev Biophys Bio Structure*, 2005, vol. 34, 91-118 [0128]
- **GREG T. HERMANSON.** *Bioconjugate Techniques*. Academic Press, Inc, 1995 [0129]
- **MEYER ; CHILKOTI.** *Nat. Biotechnol.*, 1999, vol. 17 (11), 1112-5 [0151] [0153]
- *Current Protocols in Molecular Biology*. 1995 [0151]
- **BARBEE, R.W. et al.** *Am. J. Physio.*, 1992, vol. 263 (3), R728-R733 [0188]
- **DONG WOO LIM ; KIMBERLY TRABBIC-CARLSON ; J. ANDREW MACKAY ; ASHUTOSH CHILKOTI.** Improved Non-chromatographic Purification of a Recombinant Protein by Cationic Elastin-like Polypeptides. *Biomacromolecules*, 2007, vol. 8, 1417-1424 [0194]
- **WANG L ; BROCK A ; HERBERICH B ; SCHULTZ PG.** Expanding the genetic code of Escherichia coli. *Science*, 2001, vol. 292, 498-500 [0197]
- **GE, X. ; YANG, D.S.C. ; TRABBIC-CARLSON, K. ; KIM, B. ; CHILKOTI, A. ; FILIPE, C.D.M.** Self-Cleavable Stimulus Responsive Tags for Protein Purification without Chromatography. *J. Am. Chem. Soc.*, 2005, vol. 127, 11228-11229 [0199]
- **R. M. BELAGAJE ; S. G. REAMS ; S. C. LY ; W. F. PROUTY.** Increased production of low molecular weight recombinant proteins in Escherichia coli. *Protein Sci.*, 1997, vol. 6, 1953-1962 [0211]

摘要

本發明提供包含彈性蛋白樣肽 (ELPs) 和治療性蛋白質的治療劑和組合物。該治療性蛋白質是GLP-1受體激動劑、胰島素或因子VII/VIIa，包括功能類似物。本發明還提供編碼多核苷酸，以及製備和使用所述治療劑的方法。就其用作治療劑的用途而言，本發明治療劑具有改進性質，特別包括體內半壽期、清除率和/或持久性、溶解度和生物利用度中的一種或多種改進性質。