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Carney

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(54) **THROMBIN PEPTIDE DERIVATIVES**

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(51) **Int. Cl.**
A61K 38/10 (2006.01)

(52) **U.S. Cl.** **514/13**; 530/326

(58) **Field of Classification Search** None
See application file for complete search history.

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,352,664	A *	10/1994	Carney et al.	514/13
5,500,412	A	3/1996	Carney et al.	
5,688,768	A	11/1997	Coughlin et al.	
5,716,789	A	2/1998	Sundelin et al.	
5,759,994	A	6/1998	Coughlin et al.	
5,763,575	A	6/1998	Sundelin et al.	
5,798,248	A	8/1998	Coughlin et al.	
5,912,229	A	6/1999	Thim et al.	
6,191,113	B1	2/2001	Nakahara et al.	
6,630,572	B1 *	10/2003	Carney et al.	530/327
6,855,687	B2	2/2005	Carney et al.	
6,861,407	B2 *	3/2005	Carney	514/13
6,867,190	B2 *	3/2005	Carney	514/13
6,894,027	B2 *	5/2005	Carney et al.	514/13
6,914,050	B2 *	7/2005	Carney et al.	514/13
7,034,001	B2 *	4/2006	Carney	514/13
7,049,294	B2 *	5/2006	Carney	514/13

FOREIGN PATENT DOCUMENTS

WO	WO 88/03151	5/1988
WO	WO 92/14750	9/1992
WO	WO 96/40033	12/1996
WO	WO 99/53943	10/1999
WO	WO 01/49309 A2	7/2001
WO	WO 02/04008	1/2002
WO	WO 02/05836	1/2002

OTHER PUBLICATIONS

Rubas, et al., 1994, Materials Research Society Symposium Proceedings, 331, 179-185.*
Hollenberg, 1996, Journal of Cellular Physiology, 169, 491-496.*
Starling, 1997, J. Exp. Med. vol. 185, No. 8, 1487-1492.*

Goffin, 1992, Molecular Endocrinology, 6, 1381-1392.*

Ali, N.M., et al., "Inhibition of coronary restenosis by antithrombin III in atherosclerotic swine," *Coronary Artery Disease*, 7(11):851-861, (1996).

Aoki, M., et al., "Angiogenesis induced by hepatocyte growth factor in non-infarcted myocardium and infarcted myocardium: up-regulation of essential transcription factor for angiogenesis, etc.," *Gene Therapy*, 7(5):417-427, (2000).

Bi, L.X., et al., "Thrombin Peptide TP508 Regulates BMP-2 and -7 Expression by Human Osteoblasts," *Journal of Bone and Mineral Research* 16(Suppl. 1):S261, (2001).

Carney, D.H. et al., "Initiation of Proliferative Events by Human α -Thrombin Requires Both Receptor Binding and Enzymatic Activity," *J. of Cell. Biochem.* 26:181-195 (1984).

Carney, D.H., et al., "Enhancement of Incisional Wound Healing and Neovascularization in Normal Rats by Thrombin and Synthetic Thrombin Receptor-activating Peptides," *J. Clin. Invest.*, 89:1469-1477 (1992).

Carney, D.H., et al., "Role of High-Affinity Thrombin Receptors in Postclotting Cellular Effects of Thrombin," *Seminars in Thrombosis and Hemostasis*, 18(1):91-102 (1992).

Carney, D.H. et al., "Double-Signal Hypothesis for Thrombin Initiation of Cell Proliferation," *Seminars in Thrombosis and Hemostasis*, 12(3):231 (1986).

Carney, D. H., "Postclotting Cellular Effects of Thrombin Mediated by Interaction with High-Affinity Thrombin Receptors," In *Thrombin Structure and Function*, Berliner, L.J. (ed.) (New York: Plenum Press), pp. 351-396 (1992).

Coleman, C.L., et al., "Systemic Injection of Thrombin Peptide TP508 Mitigates Angioplasty-related Restenosis in Hypercholesterolemic Rabbit Iliac Arteries," Abstract LB14, Presented at the Experimental Biology 2001 Meeting (Orlando, Florida).

Crowther, R.S., et al., "Thrombin Peptide TP508 Significantly Accelerates Repair of Fresh Fractures," *Distributed at Texas Mineralized Tissue Society*, Austin, Texas. Aug. 1998.

Glenn, K.C. et al., "Thrombin Active Site Regions Required for Fibroblast Receptor Binding and Initiation of Cell Division," *J. Biol. Chem.*, 255(6609-6616). 1980.

(Continued)

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(57) **ABSTRACT**

Disclosed are thrombin peptide derivatives comprising a polypeptide having the amino acid sequence SEQ ID NO. 2: Arg-Gly-Asp-Ala-Xaa-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val, or a C-terminal truncated fragment of the polypeptide having at least six amino acids. Xaa is alanine, glycine, serine, or an S-protected cysteine. Zero, one, two, or three amino acids in the polypeptide or polypeptide fragment differ from the corresponding position of SEQ ID NO. 2. Also disclosed are methods of treating a subject in need of treatment with a thrombin receptor agonist. The methods comprise the step of administering an effective amount of the thrombin peptide derivative described above.

OTHER PUBLICATIONS

- Glenn, K.C., et al., "Synthetic Peptides Bind to High-Affinity Thrombin Receptors and Modulate Thrombin Mitogenesis," *Peptide Research*, 1(2):65-73 (1988).
- Yang et al., "Accelerated Repair of Segmental Defects by a Synthetic Thrombin Peptide," Handout that was distributed at the Texas Mineralized Tissue Society Meeting, Sep. 1999.
- Hollenberg, Morley D., et al., "Synergistic Actions of a Thrombin-Derived Synthetic Peptide and a Thrombin Receptor-Activating Peptide in Stimulating Fibroblast Mitogenesis," *J. Cell Physiol.* 169:491-496 (1996).
- Kuraray Co Ltd. (Accession No. AAW83414, Dec. 2, 1998).
- Norfleet, A.M., et al., "Thrombin peptide, TP508, stimulates angiogenic responses in animal models of dermal wound healing, in chick chorioallantoic membranes, and in cultured human aortic and microvascular endothelial cells," *General Pharmacology*, 35:249-254 (2002).
- Norfleet, A.M., et al., "Thrombin peptide TP508 accelerates closure of dermal excisions in animal tissue with surgically induced ischemia," *Wound Repair and Regeneration*, 8(6):517-529 (2000).
- Pernia, S.D., et al., "A Synthetic Peptide Representing the Thrombin Receptor-Binding Domain Enhances Wound Closure in Vivo," *SAAS Bulletin: Biochem. & Biotech.*, 3:8-12 (1990).
- Press Release: Chrysalis BioTechnology to begin clinical trials, (Sep. 24, 1998).
- Press Release: Chrysalis Announces Completion of Pilot Chrysalin® Clinical Trial for Diabetic Ulcers, (Aug. 21, 2001).
- Press Release—Orthologic Licenses Worldwide Orthopedic Soft Tissue Rights for Chrysalin, (Jul. 9, 2001).
- Rouslahti, E. and Pierschbacher, M.D., "Arg-Gly-Asp: A Versatile Cell Recognition Signal," *Cell*, 44: 517-518 (1986).
- Simmons, D.J., et al., "Acceleration of Rat Femoral Fracture Healing by a Synthetic Thrombin Peptide," *Calcium Metabolism: Comparative Endocrinology*. Proc. Satellite Meeting, San Francisco, CA. Nov. 30, 1998. (Eds. C. Dacke, J. Danks, G. Flik and C. Gay). BioScientifica Ltd., Bradley Stoke, Bristol, UK, 1999.
- Wang, H., et al., "Effect of TP508, a thrombin-related peptide, on Cbfa1, VEGF, and collagen type II expression during femoral fracture healing," *Molecular Biology of the Cell* 2:243a, (2000).
- Sower, L.E., et al., "Thrombin Peptide, TP508, Induces Differential Gene Expression in Fibroblasts through a Nonproteolytic Activation Pathway," *Experimental Cell Research*, 247:422-431 (1999).
- Steed, D.L., et al., "Promotion and Acceleration of Diabetic Ulcer Healing by Arginine-Glycine-Aspartic Acid (RGD) Peptide Matrix," *Diabetes Care*, 18(1):39-46 (1995).
- Stiernberg, J., et al., "Acceleration of full-thickness wound healing in normal rats by the synthetic thrombin peptide, TP508," *Wound Repair and Regeneration*, 8(3):204-215 (2000).
- Stiernberg, J., et al., "The Role of Thrombin and Thrombin Receptor Activating Peptide (TRAP-508) in Initiation of Tissue Repair," *Thrombosis and Haemostasis*, 70(1):158-162, (1993).
- Tsopanoglou, N.E., and Maragoudakis, M.E., "On the Mechanism of Thrombin-induced Angiogenesis," *J. Biol. Chem.*, 274(34):23969-23976 (1999).
- Vu, T.-K.H. et al., "Molecular cloning of a functional thrombin receptor reveals a novel proteolytic mechanism of receptor activation," *Cell*, 64:1057-1068 (1991).

* cited by examiner

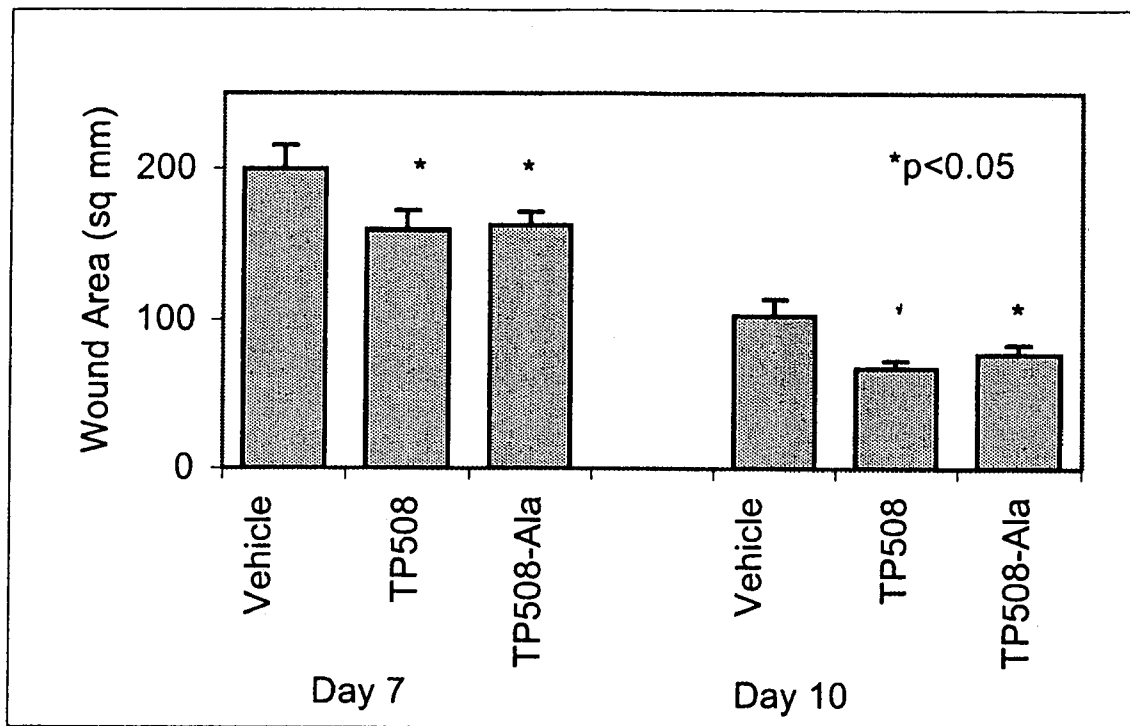


FIG. 1

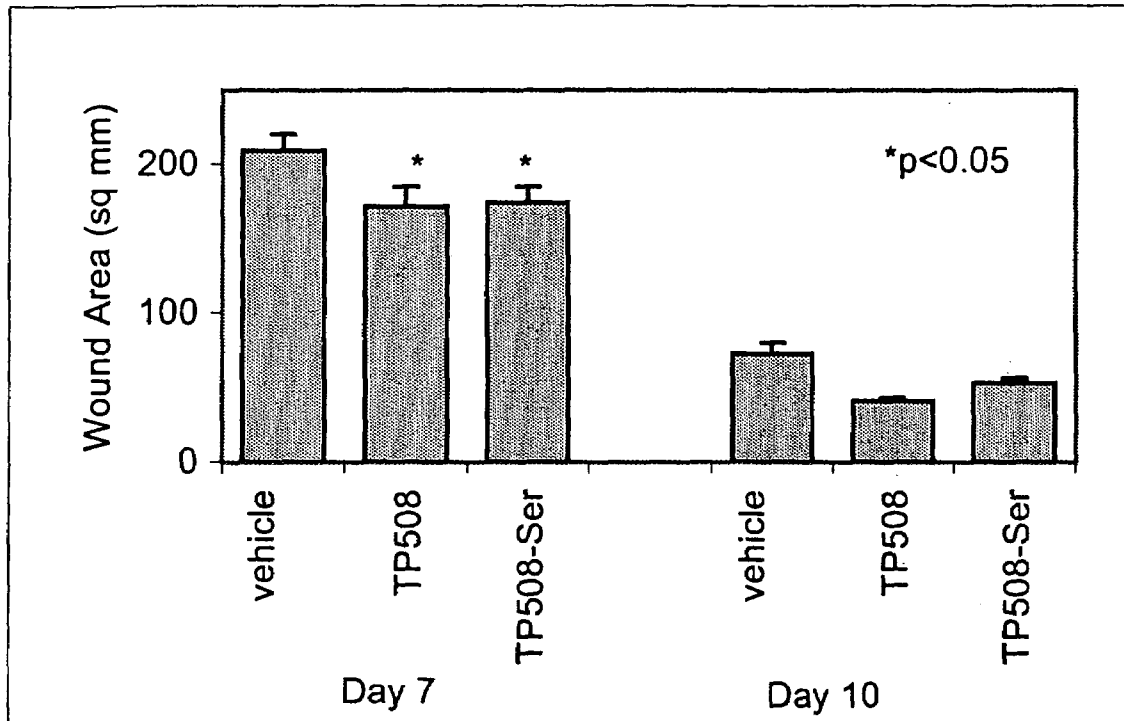


FIG. 2

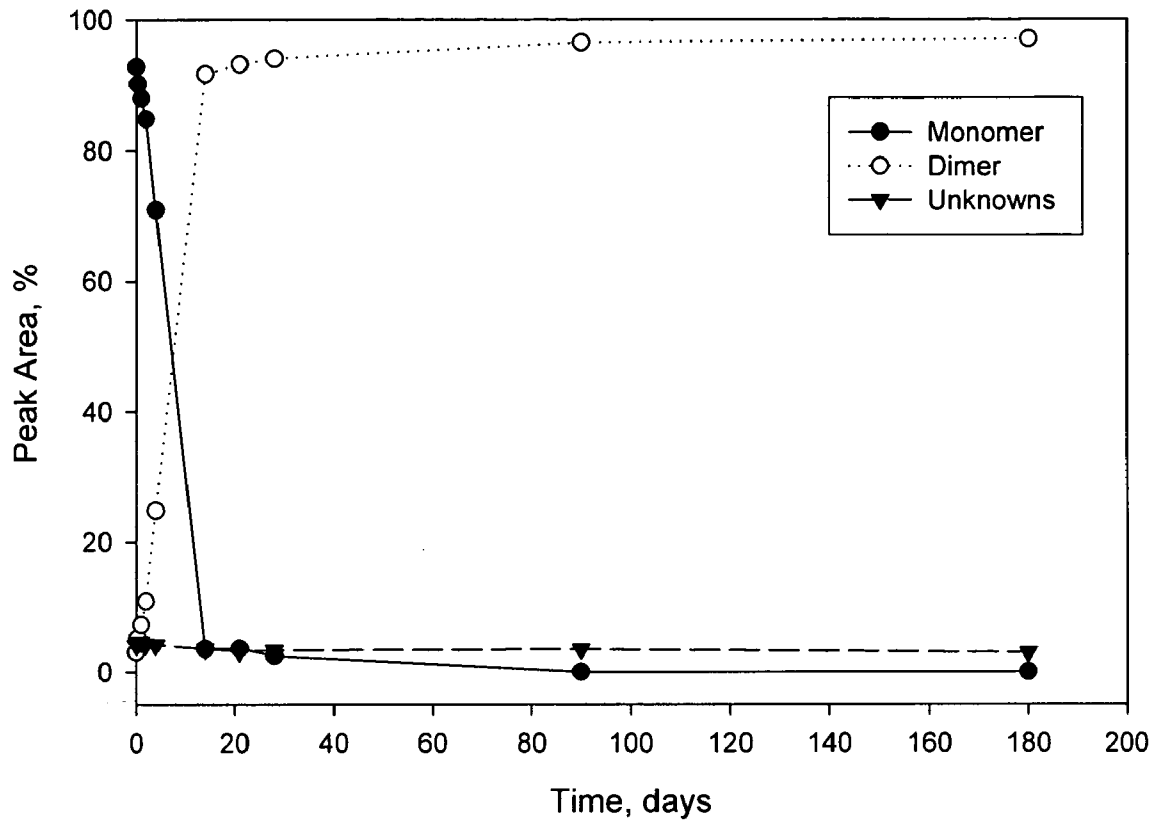


FIG. 3

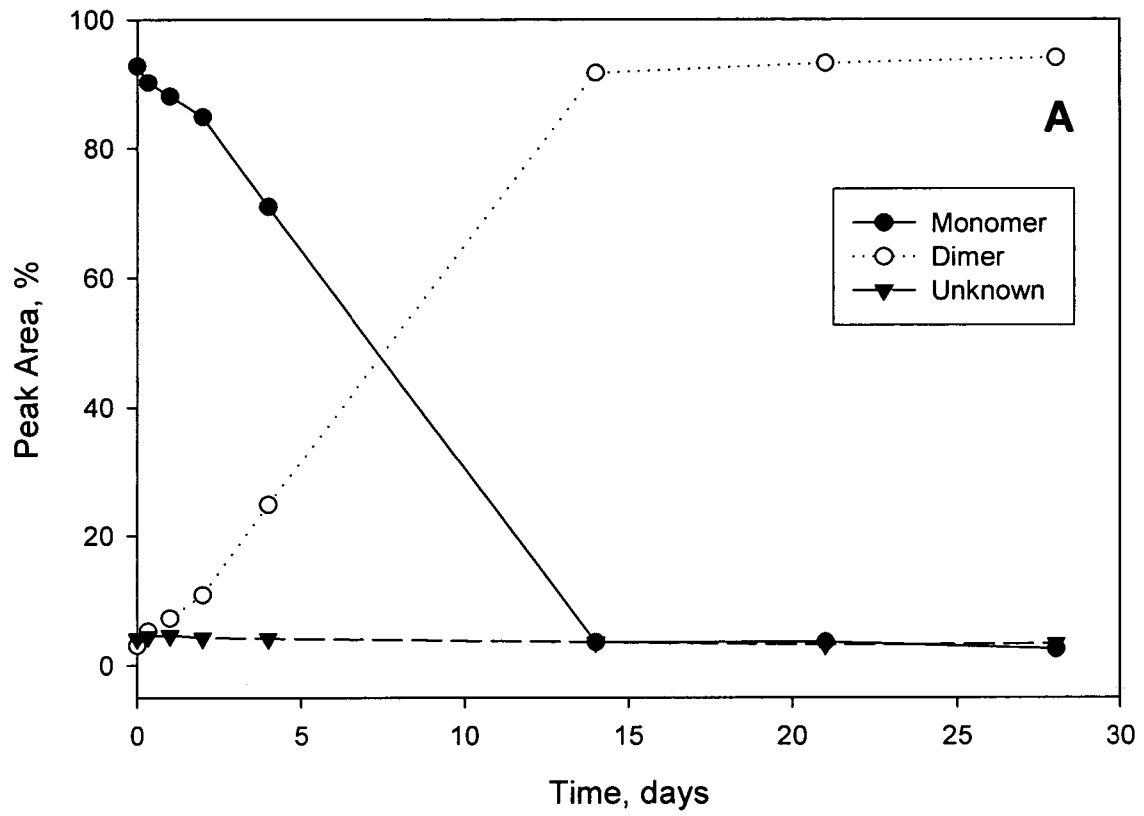


FIG. 4

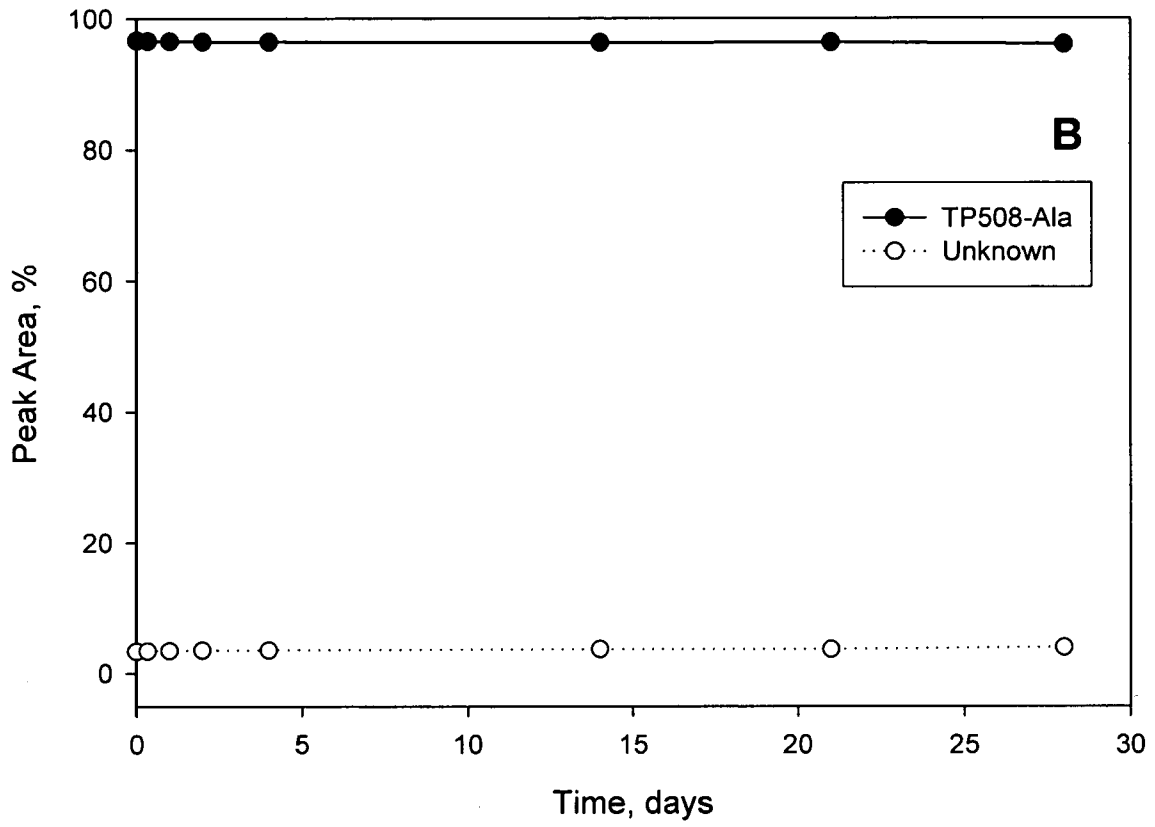


FIG. 5

THROMBIN PEPTIDE DERIVATIVES

RELATED APPLICATIONS

This application is a continuation of International Application No. PCT/US2003/020635, which designated the United States and was filed Jul. 1, 2003, published in English, which claims the benefit of U.S. Provisional Application No. 60/393,580, filed Jul. 2, 2002. The entire teachings of the above applications are incorporated herein by reference.

BACKGROUND OF THE INVENTION

Thrombin, a multi-functional enzyme already known for its blood-clotting activity, has been recently reported to be an important cell-growth factor. For example, thrombin has been shown to promote angiogenesis, the development of new blood vessels, and to stimulate endothelial cell proliferation. These processes are a pivotal part of healing wounds.

Thrombin peptide derivatives are molecules having an amino acid sequence derived at least in part from that of thrombin, and which are active toward certain thrombin receptors. For example, thrombin peptide derivatives from amino acids 508-530 of human pro-thrombin have been described by the present inventors for promoting thrombin receptor mediated cell stimulation and for their use in the treatment of wounds, and stimulation of angiogenesis (see, e.g., U.S. Pat. No. 5,500,412 or 5,352,664, the contents of which are incorporated herein by reference in their entirety). Because of their biological activity, these thrombin peptide derivatives show great potential as pharmaceuticals. TP508 is one such example of a thrombin peptide derivative and has the amino acid sequence of H-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Cys-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val-NH₂ (SEQ ID NO.1).

Strict regulations by the Food and Drug Administration (FDA) require a high degree of purity of biologically active agents when used as pharmaceuticals. It therefore is necessary to obtain active thrombin peptide derivatives that maintain their purity over extended time periods, if these compounds are to be used to treat humans. Unfortunately, the purity of thrombin peptide derivative TP508 diminishes over time because of dimerization resulting from disulfide bond formation. For example, TP508 has a half-life of about 2 to about 4 hours in buffered solutions at neutral pH. Peptide dimers are degradation products and therefore may be considered contaminants of a pharmaceutical composition.

Therefore, there is a need for new peptides with the activity of thrombin peptide derivatives, but which do not form dimers in solution.

SUMMARY OF THE INVENTION

It has now been found that thrombin peptide derivatives in which cysteine is replaced with non-reactive amino acids of similar size are free of dimers in solution and retain their activity toward thrombin receptors. For example, replacing cysteine in SEQ ID NO:1 with alanine (TP508 Cys→Ala) or serine (TP508 Cys→Ser; H-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Ser-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val-NH₂ (SEQ ID NO:14)) results in a thrombin peptide derivative having about the same level of activity toward the thrombin receptor as TP508 (see Examples 1 and 2). Moreover, TP508 Cys→Ala shows no dimerization after 6 months in saline solution (see Example 4). Based on this discovery, the invention provides novel peptides, pharmaceu-

tical compositions comprising these peptides, and methods useful for treating a subject in need of treatment with a thrombin receptor agonist.

One embodiment of the present invention is a thrombin peptide derivative comprising a polypeptide having the amino acid sequence SEQ ID NO:2: Arg-Gly-Asp-Ala-Xaa-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val, or a C-terminal truncated fragment of the polypeptide having at least six amino acids. Zero, one, two, or three amino acids in the peptide or peptide fragment differ from the corresponding position of SEQ ID NO:2, provided that Xaa is alanine, glycine, serine, or an S-protected cysteine. Preferably, the difference is conservative. The thrombin peptide derivatives are optionally amidated at the C-terminus and/or acylated at the N-terminus.

Another embodiment of the invention also relates to pharmaceutical compositions comprising a thrombin receptor agonist or a thrombin peptide derivative described herein and a pharmaceutically acceptable carrier or diluent.

Another embodiment of the invention further relates to methods of treating a subject in need of treatment with a thrombin receptor agonist. The methods comprise the step of administering an effective amount of thrombin peptide derivative described herein.

Advantages of the thrombin peptide derivatives of the present invention include longer storage life in solution compared with TP508. In addition, these thrombin peptide derivatives are less susceptible to oxidation. Therefore, it is possible to deliver precise and reproducible dosages with the disclosed peptides, even after storage in solution for prolonged periods of time. The thrombin peptide derivatives described herein are also inexpensive to produce. The thrombin peptide derivatives can be used in the treatment and/or prevention of diseases and/or conditions in which angiogenesis and cell proliferation would be beneficial. The thrombin peptide derivatives can be used to help treat, for example, wounds such as diabetic ulcers, bone fractures, and cartilage damage. The thrombin peptide derivatives can also be used to prevent restenosis in patients after angioplasty and regenerate blood vessels in cardiac tissue.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a graph showing that the wound healing activity of the thrombin peptide derivative, TP508-Ala, is similar to that of TP508. The graph shows wound area measurements (indicated in mm²) on the dorsum of male Sprague-Dawley rats from post-wounding Day 7 and Day 10. The saline control is indicated as "vehicle," the TP508 control is indicated as "TP508," and the thrombin peptide derivative TP508-Ala is indicated as "TP508-Ala."

FIG. 2 is a graph showing that the wound healing activity of the thrombin peptide derivative, TP508-Ser, is similar to that of TP508. The graph shows wound area measurements (indicated in mm²) on the dorsum of male Sprague-Dawley rats from post-wounding Day 7 and Day 10. The saline control is indicated as "vehicle," the TP508 control is indicated as "TP508," and the thrombin peptide derivative TP508-Ser is indicated as "TP508-Ser."

FIG. 3 is a graph showing the conversion of TP508 to dimer over time. The graph shows the HPLC peak area measurements of TP508-monomer, TP508-dimer and unknowns found in samples of TP508 saline solution (5 mg/mL, incubated at 4° C.), taken at intervals over a time period of 6 months. Peak area is indicated as percent. Time is indicated as days. Monomer is indicated as (●-). Dimer is indicated as (...○...) Unknowns are indicated as (--▼--).

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FIG. 4 is a graph showing the conversion of TP508 to dimer over time. The graph shows the HPLC peak area measurements of TP508-monomer, TP508-dimer and unknowns found in samples of TP508 saline solution (5 mg/mL, incubated at 4° C.), taken at intervals over a time period of 1 month. Peak area is indicated as percent. Time is indicated as days. Monomer is indicated as (-●-). Dimer is indicated as (...○...). Unknowns are indicated as (--▼--).

FIG. 5 is a graph showing that TP508-Ala does not convert to dimer. The graph shows the HPLC peak area measurements of TP508-Ala and an unknown found in samples of TP508-Ala saline solution (5 mg/mL, incubated at 4° C.), taken at intervals over a time period of 1 month. Peak area is indicated as Percent. Time is indicated as Days. TP508-Ala is indicated as (-●-). Unknown is indicated as (...○...).

DETAILED DESCRIPTION OF THE INVENTION

Applicants have found that the peptides of the present invention essentially do not dimerize and still have about the same biological activity as the thrombin peptide derivatives in the prior art. To minimize dimerization of the thrombin peptides of the present invention, cysteine residues normally found in thrombin peptide derivatives are replaced with amino acids having similar size and charge properties. Examples of suitable amino acids include alanine, glycine, serine, or an S-protected cysteine. Preferably, cysteine is replaced with alanine.

It will be understood that the thrombin peptide derivatives disclosed herein can have C-terminal amides. A "C-terminal amide" is an amide at the C-terminal amino acid residue in which the alpha carboxylic acid is replaced with an amide. For example, amidated C-terminal amino acid residues have the formula: —NH—CH(R_a)C(O)—NR_bR_c. R_a is an amino acid side chain. An amino acid side chain can be hydrogen, a substituted or unsubstituted C₁-C₁₀ aliphatic group, or a substituted or unsubstituted aromatic group of up to 10 carbon atoms. Preferably R_a is an amino acid side chain corresponding to a side chain in a naturally occurring amino acid. R_b and R_c are independently hydrogen, a C₁-C₁₀ substituted or unsubstituted aliphatic group. Preferably, the C-terminal amide is a carboxamide (—C(O)NH₂). As used herein, "—NH₂" at the C-terminus indicates a C-terminus carboxamide; "—OH" at the C-terminus indicates that the peptide has a free C-terminus; and no designation at the C-terminus indicates that the peptide is amidated at the C-terminus or has a free C-terminus.

It will also be understood that the thrombin peptide derivatives disclosed herein can have an acylated N-terminus. An "acylated N-terminus" is an N-terminal in which the nitrogen of the N-terminal amino acid residue is acylated. For example, acylated N-terminal amino acids residues have the formula: R_dC(O)—NH—CHR_eC(O)—. R_d is hydrogen, a C₁-C₁₀ substituted or unsubstituted aliphatic group, or a substituted or unsubstituted aromatic group of up to 10 carbon atoms. Acetyl is a preferred acyl group. An "—H" at the N-terminus indicates that the N-terminus is unsubstituted; and no designation at the N-terminus indicates that the terminus is acylated or unsubstituted.

Preferably, the N-terminus of a thrombin peptide derivative is free (i.e., unsubstituted) and the C-terminus is free (i.e., unsubstituted) or amidated, preferably a carboxamide (i.e., —C(O)NH₂).

Thrombin peptide derivatives are believed to activate cells by binding to a high-affinity cell-surface thrombin receptor known as the non-proteolytically-activated thrombin receptor (hereinafter "NPAR") (R. Horvat, et. al., *J. Cell Sci.* 108,

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1155-1164 1995). Compounds which stimulate NPAR are said to be thrombin receptor agonists. NPAR activation can be assayed based on the ability of molecules to stimulate cell proliferation when added to fibroblasts in the presence of submitogenic concentrations of thrombin or molecules that activate protein kinase C or compete with ¹²⁵I-thrombin for high affinity binding to thrombin receptors, as disclosed in U.S. Pat. Nos. 5,352,664 and 5,500,412 and in Glenn et al., *J. Peptide Research* 1:65 (1988).

To further identify regions of thrombin involved in high-affinity binding and generation of mitogenic signals, two peptides representing specific domains within p508-530 were tested. The first peptide represented residues 519 to 530 of the B-chain region of human prothrombin, a region of thrombin that is highly conserved among serine proteases. The second peptide represented residues 517 to 520 of prothrombin, a region of thrombin homologous to the fibronectin cell binding domain.

Both of these peptides were able to compete for 30% to 50% of the binding of [¹²⁵I]-alpha-thrombin to ME cells, but both required higher concentrations than was required with the initial peptide p508-530 (Table). For example, 30% inhibition of [¹²⁵I]-alpha-thrombin binding required 33- to 50-fold higher concentrations of p519-530 and p517-520 than p508-530, respectively. Thus, both of these peptides appear to interact with thrombin receptors, but at a lower affinity than p508-530. Because p517-520 is homologous to the fibronectin cell binding domain, a peptide having the sequence Arg-Gly-Ala-Ser (SEQ ID NO:18; the sequence of the fibronectin specific peptide) was also tested for its ability to compete for [¹²⁵I]-alpha-thrombin binding. At a concentration of 1.3 μM, this peptide did not compete with [¹²⁵I]-alpha-thrombin for binding. Thus, the receptor for alpha-thrombin is not the same membrane protein that specifically interacts with fibronectin and causes the apparent growth promoting action of fibronectin. In addition, these results demonstrate the requirement for alanine within the thrombin receptor binding domain, since substitution of alanine with serine eliminated the ability of the synthetic peptide to compete for alpha-thrombin binding.

TABLE

Comparison of Peptide Competition for [¹²⁵ I]-Alpha-Thrombin Binding to ME Cells.			
Peptide	Amino Acid Sequence	Concentration Required for 30% Inhibition	Maximal % Inhibition (and Conc.)
p508-530	AGYKPDEG- KRGDACE- GDSGGPFV (SEQ ID NO:1)	6 nM	78% (40 nM)
p519-530	DACEGDSGGPFV (SEQ ID NO:17)	200 nM	51% (800 nM)
p517-520	RGDA (SEQ ID NO:16)	300 nM	50% (2.7 μM)

The mitogenic activity of the smaller thrombin derivatives, p519-530 and p517-520, was tested. As indicated above, both of these peptides bind to the high-affinity thrombin receptor. In these experiments, increasing concentrations of p519-530 and p517-520 were added to quiescent NIL cells in the presence of 2 and 4 nM alpha-thrombin. As shown in FIG. 6 of U.S. Pat. No. 5,500,412, p519 enhanced DNA synthesis over a range of concentrations while p517-520 did not. In fact, p517-520 actually inhibited DNA synthesis.

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Thrombin peptide derivatives stimulate NPAR and have less than about fifty amino acids, preferably less than about thirty-three amino acids. Thrombin peptide derivatives also have sufficient homology to the fragment of human thrombin corresponding to prothrombin amino acids 508-530: Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Cys-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val (SEQ ID NO:3) so that the polypeptide activates NPAR. The thrombin peptide derivatives described herein typically have at least six amino acids and preferably between about 12 and 33 amino acids, more preferably between about 12 and 23 amino acids.

In a first preferred embodiment, the thrombin peptide derivative comprises a polypeptide having the amino acid sequence of SEQ ID NO:4: Arg-Gly-Asp-Ala-Xaa-X₁-Gly-Asp-Ser-Gly-Gly-Pro-X₂-Val, or a C-terminal truncated fragment thereof having at least six amino acids. More preferably, the thrombin peptide derivative has the amino acid sequence of SEQ ID NO:5: Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val or a fragment thereof comprising amino acids 10-18 of SEQ ID NO:5. Even more preferably, the thrombin peptide derivative has the amino acid sequence SEQ ID NO:6: Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-X₁-Gly-Asp-Ser-Gly-Gly-Pro-X₂-Val, or a fragment thereof comprising amino acids 10-18 of SEQ ID NO:6. Xaa is alanine, glycine, serine or an S-protected cysteine. X₁ is Glu or Gln and X₂ is Phe, Met, Leu, His or Val. Preferably X₁ is Glu, X₂ is Phe, and Xaa is alanine. One example of a thrombin peptide derivative of this type is a polypeptide having the amino acid sequence Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Ala-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val (SEQ ID NO:7). A further example of a thrombin peptide derivative of this type is a polypeptide having the amino acid sequence H-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Ala-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val-NH₂ (SEQ ID NO:8). Zero, one, two or three amino acids in the thrombin peptide derivative differ from the amino acid at the corresponding position of SEQ ID NO. 4, 5, 6, 7 or 8, provided that Xaa is alanine, glycine, serine or an S-protected cysteine. Preferably, the difference is conservative. SEQ ID NO:13 is H-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-X₁-Gly-Asp-Ser-Gly-Gly-Pro-X₂-Val-R1; wherein X₁ is Glu or Gln; X₂ is Phe, Met, Leu, His or Val; and Xaa is alanine, glycine, serine or an S-protected cysteine. and R1 is —OH or —NH₂. SEQ ID NO:15 is H-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val-R1; wherein Xaa is alanine, glycine, serine or an S-protected cysteine; and wherein R1 is —OH or —NH₂.

In a second preferred embodiment, the thrombin peptide derivative comprises a polypeptide having the amino acid sequence SEQ ID NO:9: Asp-Asn-Met-Phe-Xbb-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val-Met-Lys-Ser-Pro-Phe, or a fragment thereof comprising amino acids 6-28. More preferably, the thrombin peptide derivative comprises a polypeptide having the amino acid sequence SEQ NO:10: Asp-Asn-Met-Phe-Xbb-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-X₁-Gly-Asp-Ser-Gly-Gly-Pro-X₂-Val-Met-Lys-Ser-Pro-Phe, or a fragment thereof comprising amino acids 6-28. Xaa and Xbb are independently alanine, glycine, serine or an S-protected cysteine. X₁ is Glu or Gln and X₂ is Phe, Met, Leu, His or Val. Preferably X₁ is Glu, X₂ is Phe, and Xaa and Xbb are alanine. One example of a thrombin peptide derivative of this type is a polypeptide having the amino acid sequence Asp-Asn-Met-Phe-Ala-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Ala-

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Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val-Met-Lys-Ser-Pro-Phe (SEQ ID NO:11). A further example of a thrombin peptide derivative of this type is a polypeptide having the amino acid sequence H-Asp-Asn-Met-Phe-Ala-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Ala-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val-Met-Lys-Ser-Pro-Phe-NH₂ (SEQ ID NO:12). Zero, one, two or three amino acids in the thrombin peptide derivative differ from the amino acid at the corresponding position of SEQ ID NO: 9, 10 11 or 12, and Xaa and Xbb are independently alanine, glycine, serine or an S-protected cysteine. Preferably, the difference is conservative.

A “conservative substitution” is the replacement of an amino acid with another amino acid that has the same net electronic charge and approximately the same size and shape. Amino acids with aliphatic or substituted aliphatic amino acid side chains have approximately the same size when the total number carbon and heteroatoms in their side chains differs by no more than about four. They have approximately the same shape when the number of branches in the their side chains differs by no more than one. Amino acids with phenyl or substituted phenyl groups in their side chains are considered to have about the same size and shape. Listed below are five groups of amino acids. Replacing an amino acid in a polypeptide with another amino acid from the same group results in a conservative substitution:

Group I: glycine, alanine, valine, leucine, isoleucine, serine, threonine, cysteine, and non-naturally occurring amino acids with C1-C4 aliphatic or C1-C4 hydroxyl substituted aliphatic side chains (straight chained or monobranched).

Group II: glutamic acid, aspartic acid and non-naturally occurring amino acids with carboxylic acid substituted C1-C4 aliphatic side chains (unbranched or one branch point).

Group III: lysine, ornithine, arginine and non-naturally occurring amino acids with amine or guanidino substituted C1-C4 aliphatic side chains (unbranched or one branch point).

Group IV: glutamine, asparagine and non-naturally occurring amino acids with amide substituted C1-C4 aliphatic side chains (unbranched or one branch point).

Group V: phenylalanine, phenylglycine, tyrosine and tryptophan.

A “highly conservative substitution” is the replacement of an amino acid with another amino acid that has the same functional group in the side chain and nearly the same size and shape. Amino acids with aliphatic or substituted aliphatic amino acid side chains have nearly the same size when the total number carbon and heteroatoms in their side chains differs by no more than two. They have nearly the same shape when they have the same number of branches in the their side chains. Examples of highly conservative substitutions include valine for leucine, threonine for serine, aspartic acid for glutamic acid and phenylglycine for phenylalanine. Examples of substitutions which are not highly conservative include alanine for valine, alanine for serine and aspartic acid for serine.

An “S-protected cysteine” is a cysteine residue in which the reactivity of the thiol moiety, —SH, is blocked with a protecting group. Suitable protecting groups are known in the art and are disclosed, for example, in T. W. Greene and P. G. M. Wuts, *Protective Groups in Organic Synthesis*, 3rd Edition, John Wiley & Sons, (1999), pp. 454-493, the teachings of which are incorporated herein by reference in their entirety. Suitable protecting groups should be non-toxic, stable in pharmaceutical formulations and have minimum additional

functionality to maintain the activity of the thrombin peptide derivative. A free thiol can be protected as a thioether, a thioester, or oxidized to an unsymmetrical disulfide. Preferably the thiol is protected as a thioether. Suitable thioethers include, but are not limited to, S-alkyl thioethers (e.g., C₁-C₅ alkyl), and S-benzyl thioethers (e.g., Cysteine-S—S-t-Bu). Preferably, the protective group is an alkyl thioether. More preferably, the S-protected cysteine is an S-methyl cysteine. Alternatively, the protecting group can be: 1) a cysteine or a cysteine-containing peptide (the “protecting peptide”) attached to the cysteine thiol group of the thrombin peptide derivative by a disulfide bond; or 2) an amino acid or peptide (“protecting peptide”) attached by a thioamide bond between the cysteine thiol group of the thrombin peptide derivative and a carboxylic acid in the protecting peptide (e.g., at the C-terminus or side chain of aspartic acid or glutamic acid). The protecting peptide can be physiologically inert (e.g., a polyglycine or polyalanine of no more than about fifty amino acids optionally interrupted by a cysteine) or can have a desirable biological activity. However, the present invention does not contemplate thrombin peptide derivative dimers wherein the protecting peptide is a second thrombin peptide derivative. Thrombin peptide derivative dimers are disclosed in the co-pending U.S. Provisional Application entitled THROMBIN PEPTIDE DERIVATIVE DIMERS, Provisional Application No. 60/393,579, filed Jul. 2, 2002, the entire teachings of which are incorporated herein by reference.

An “N-terminal truncated fragment” refers to a fragment remaining after removing an amino acid or block of amino acids from the N-terminus, preferably a block of no more than six amino acids, more preferably a block of no more than three amino acids. Optionally, an N-terminal truncated fragment is acylated and/or amidated as described above.

A “C-terminal truncated fragment” refers to a fragment remaining after removing an amino acid or block of amino acids from the C-terminus, preferably a block of no more than six amino acids, more preferably a block of no more than three amino acids. Optionally, a C-terminal truncated fragment is amidated and/or acylated as described above.

A “non-aromatic heterocyclic group”, as used herein, is a non-aromatic carbocyclic ring system that has 3 to 10 atoms and includes at least one heteroatom, such as nitrogen, oxygen, or sulfur. Examples of non-aromatic heterocyclic groups include piperazinyl, piperidinyl, pyrrolidinyl, morpholinyl, thiomorpholinyl.

The term “aryl group”, as used herein, includes both carbocyclic and heterocyclic aromatic ring systems. Examples of aryl groups include phenyl, indolyl, furanyl and imidazolyl.

An “aliphatic group” is a straight chain, branched or cyclic non-aromatic hydrocarbon. An aliphatic group can be completely saturated or contain one or more units of unsaturation (e.g., double and/or triple bonds), but is preferably saturated, i.e., an alkyl group. Typically, a straight chained or branched aliphatic group has from 1 to about 10 carbon atoms, preferably from 1 to about 4, and a cyclic aliphatic group has from 3 to about 10 carbon atoms, preferably from 3 to about 8. Aliphatic groups include, for example, methyl, ethyl, n-propyl, iso-propyl, n-butyl, sec-butyl, tert-butyl, pentyl, cyclopentyl, hexyl, cyclohexyl, octyl and cyclooctyl.

Suitable substituents for an aliphatic group, an aryl group or a non-aromatic heterocyclic group are those which do not significantly lower therapeutic activity of the thrombin peptide derivative, for example, those found on naturally occurring amino acids. Examples include —OH, a halogen (—Br, —Cl, —I and —F), —O(R_e), —O—CO—(R_e), —CN,

—NO₂, —COOH, =O, —NH₂—NH(R_e), —N(R_e)₂, —COO(R_e), —CONH₂, —CONH(R_e), —CON(R_e)₂, —SH, —S(R_e), an aliphatic group, an aryl group and a non-aromatic heterocyclic group. Each R_e is independently an alkyl group or an aryl group. A substituted aliphatic group can have more than one substituent.

A “subject” is preferably a human, but can also be an animal in need of treatment with a thrombin receptor agonist, e.g., companion animals (e.g., dogs, cats, and the like), farm animals (e.g., cows, pigs, horses and the like) and laboratory animals (e.g., rats, mice, guinea pigs and the like).

Subjects “in need of treatment” with a thrombin receptor agonist, are subjects with diseases and/or conditions that can be treated with thrombin receptor agonists and thrombin peptide derivatives to achieve a beneficial therapeutic and/or prophylactic result. A beneficial outcome includes a decrease in the severity of symptoms or delay in the onset of symptoms, increased longevity and/or more rapid or more complete resolution of the disease or condition. For example, a subject in need of treatment requires cell proliferation involving chondrocytes, angiogenesis, bone growth, cardiac repair, wound healing or inhibition of restenosis.

Thrombin peptide derivatives have been shown to stimulate proliferation of endothelial cells, fibroblasts, and keratinocytes (see, e.g., U.S. Pat. No. 5,500,412 or 5,352,664, the contents of which are incorporated herein by reference in their entirety). The disclosed thrombin peptide derivatives can therefore be used to promote healing in acute wounds such as, for example, burns, dermal wounds, surgical wounds and bone fractures. In addition, thrombin peptide derivatives have recently been shown to be particularly effective in promoting the healing of chronic wounds such as, diabetic ulcers, venous ulcers, and pressure sores (see, e.g., WO 03/013569, the contents of which are incorporated herein by reference in their entirety). Thrombin peptide derivatives have also been shown to stimulate the growth of chondrocytes (see, e.g., WO 02/07748, the contents of which are incorporated herein by reference in their entirety). Thus thrombin peptide derivatives, including the compounds of the present invention can be used to stimulate chondrocyte growth and repair in, for example patients with osteoarthritis or joint injuries. Other uses for thrombin peptide derivatives, including those of the present invention, include stimulating bone growth to promote healing of simple fractures, non-union fractures, voids and gaps in bone and bone grafts, preventing restenosis in patients after angioplasty and promoting the regeneration of blood vessels cardiac tissue (see, e.g., WO 02/005836 and WO 02/004008, the contents of which are incorporated herein by reference in their entirety).

An “effective amount” is the quantity of thrombin peptide derivative that results in an improved clinical outcome of the condition being treated with the thrombin peptide derivative compared with the absence of treatment. The amount of thrombin peptide derivative administered will depend on the degree, severity, and type of the disease or condition, the amount of therapy desired, and the release characteristics of the pharmaceutical formulation. It will also depend on the subject’s health, size, weight, age, sex and tolerance to drugs. Typically, the agonist is administered for a sufficient period of time to achieve the desired therapeutic effect. Typically between about 1 µg per day and about 1 mg per day of the thrombin peptide derivative (preferably between about 5 µg per day and about 100 µg per day) is administered to the subject in need of treatment.

The thrombin peptide derivative can be administered by any suitable route, locally or systemically, including, for example, by parenteral administration. Parenteral administra-

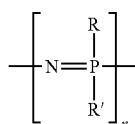
tion can include, for example, intramuscular, intravenous, subcutaneous, or intraperitoneal injection. Topical administration for treating wounds can include, for example, creams, gels, ointments or aerosols. Respiratory administration can include, for example, inhalation or intranasal drops. For certain indications such as stimulating bone growth, cartilage repair, cardiac repair and the treatment of restenosis, it is advantageous to inject or implant the thrombin peptide derivative directly to the treatment site. The thrombin peptide derivative can be advantageously administered in a sustained release formulation.

The thrombin peptide derivative can be administered to the subject in conjunction with an acceptable pharmaceutical carrier as part of a pharmaceutical composition. The formulation of the pharmaceutical composition will vary according to the route of administration selected. Suitable pharmaceutical carriers may contain inert ingredients which do not interact with the compound. The carriers should be biocompatible, i.e., non-toxic, non-inflammatory, non-immunogenic and devoid of other undesired reactions at the administration site. Examples of pharmaceutically acceptable carriers include, for example, saline, aerosols, commercially available inert gels, or liquids supplemented with albumin, methyl cellulose or a collagen matrix. Standard pharmaceutical formulation techniques can be employed, such as those described in Remington's Pharmaceutical Sciences, Mack Publishing Company, Easton, Pa.

For indications such as bone growth, cartilage repair, cardiac repair and inhibition of restenosis, it may be advantageous to administer the thrombin peptide derivative in a sustained release formulation. Polymers are often used to form sustained release formulations. Examples of these polymers include poly α -hydroxy esters such as polylactic acid/polyglycolic acid homopolymers and copolymers, polyphosphazenes (PPHOS), polyanhydrides and poly(propylene fumarates).

Poly(lactic acid)/polyglycolic acid (PLGA) homo and copolymers are well known in the art as sustained release vehicles. The rate of release can be adjusted by the skilled artisan by variation of polylactic acid to polyglycolic acid ratio and the molecular weight of the polymer (see Anderson, et al., *Adv. Drug Deliv. Rev.* 28:5 (1997), the entire teachings of which are incorporated herein by reference). The incorporation of poly(ethylene glycol) into the polymer as a blend to form microparticle carriers allows further alteration of the release profile of the active ingredient (see Cleek et al., *J. Control Release* 48:259 (1997), the entire teachings of which are incorporated herein by reference). Ceramics such as calcium phosphate and hydroxyapatite can also be incorporated into the formulation to improve mechanical qualities.

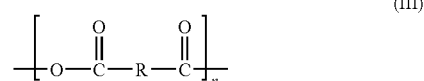
PPHOS polymers contain alternating nitrogen and phosphorous with no carbon in the polymer backbone, as shown below in Structural Formula (II):



The properties of the polymer can be adjusted by suitable variation of side groups R and R' that are bonded to the polymer backbone. For example, the degradation of and drug release by PPHOS can be controlled by varying the amount of

hydrolytically unstable side groups. With greater incorporation of either imidazolyl or ethylglycol substituted PPHOS, for example, an increase in degradation rate is observed (see Laurencin et al., *J. Biomed Mater. Res.* 27:963 (1993), the entire teachings of which are incorporated herein by reference), thereby increasing the rate of drug release.

Polyanhydrides, shown in Structural Formula (III), have well defined degradation and release characteristics that can be controlled by including varying amounts of hydrophobic or hydrophilic monomers such as sebacic acid and 1,3-bis(p-carboxyphenoxy)propane (see Leong et al., *J. Biomed. Mater. Res.* 19:941 (1985), the entire teachings of which are incorporated herein by reference). To improve mechanical strength, anhydrides are often copolymerized with imides to form polyanhydride-co-imides. Examples of polyanhydride-co-imides that are suitable for orthopaedic applications are poly(trimellitylimido-glycine-co-1,6-bis(carboxyphenoxy)hexane and pyromellitylimidoalanine:1,6-bis(p-carboxyphenoxy)hexane copolymers.



Carriers for stimulating bone or cartilage growth advantageously include porous matrices which can then serve as a scaffolding for bone and tissue growth into which bone progenitor cells and osteogenic cells may migrate and attach. Such carriers are said to be osteoconductive. For certain applications, the carrier should preferably have sufficient mechanical strength to maintain its three dimensional structure and help support the immobilization of the bone or tissue segments being united or grafted together.

Examples of suitable osteoconductive carriers include collagen (e.g., bovine collagen), fibrin, calcium phosphate ceramics (e.g., hydroxyapatite and tricalcium phosphate), calcium sulfate, guanidine-extracted allogenic bone and combinations thereof. A number of suitable carriers are commercially available, such as COLLAGRAFT® (Cohension Technologies Inc., Palo Alto, Calif.), which is a mixture of hydroxyapatite, tricalcium phosphate and fibrillar collagen, and PRO OSTEON 500™ (Interpore Cross International, Irvine, Calif.), which is a hydroxyapatite biomatrix formed by the conversion of marine coral calcium carbonate to crystalline hydroxyapatite.

Descriptions of synthetic biodegradable polymers that can serve as osteoconductive carriers with sustained release characteristics, can be found in Behravesch et al., *Clinical Orthopaedics* 367:S118 (1999) and Lichun et al., *Polymeric Delivery Vehicles for Bone Growth Factors* in "Controlled Drug Delivery—Designing Technologies for the Future" Park and Mrsny eds., American Chemical Society, Washington, DC (2000). The entire teachings of these references are incorporated herein by reference. Examples of these polymers include poly α -hydroxy esters such as polylactic acid/polyglycolic acid homopolymers and copolymers, polyphosphazenes (PPHOS), polyanhydrides and poly(propylene fumarates), which are described above in detail.

Implantable pharmaceutical compositions of the present invention are particularly useful because they can be administered at a site in need of bone growth. "Implantation" or "administration at a site" means in sufficient proximity to the site in need of treatment so that bone growth occurs (e.g., more bone growth in the presence of the drug than in its

absence) at the site when the thrombin peptide derivative is released from the pharmaceutical composition. These pharmaceutical compositions can be shaped as desired in anticipation of surgery or shaped by the physician or technician during surgery. It is preferred to shape the matrix to span a tissue defect and to take the desired form of the new tissue. In the case of bone repair of a non-union defect, for example, it is desirable to use dimensions that span the non-union. In bone formation procedures, the material is slowly absorbed by the body and is replaced by bone in the shape of or very nearly the shape of the implant. Alternatively, the pharmaceutical compositions can be administered to the site in the form of microparticles or microspheres. The microparticles are placed in contact or in close proximity to the site in need of osteoinduction either by surgically exposing the site and applying the microparticles on or in close proximity to the site by painting, pipetting, spraying, injecting or the like. Microparticles can also be delivered to the site by endoscopy or by laparoscopy.

Poly(propylene fumarates) (PPF) are highly desirable biocompatible implantable carriers for use in repairing bone defects because they are an injectable, in situ polymerizable, biodegradable material. "Injectable" means that the material can be injected by syringe through a standard needle used for injecting pastes and gels. PPF, combined with a vinyl monomer (N-vinyl pyrrolidinone) and an initiator (benzoyl peroxide), forms an injectable solution that can be polymerized in situ. It is particularly suited for filling skeletal defects of a wide variety of sizes and shapes (see Suggs et al., *Macromolecules* 30:4318 (1997), Peter et al., *J. Biomater. Sci. Poly., Ed.* 10:363 (1999) and Yaszemski et al., *Tissue Eng.* 1:41 (1995), the entire teachings of which are incorporated herein by reference). The addition of solid phase components such as P-tricalcium phosphate and sodium chloride can improve the mechanical properties of PPF polymers (see Peter et al., *J. Biomed. Mater. Res.* 44:314 (1999), the entire teachings of which are incorporated herein by reference).

In yet another alternative, the pharmaceutical composition can be partially enclosed in a supporting physical structure such as a mesh, wire matrix, stainless steel cage, threaded interbody fusion cage and the like before administering to the site in need of bone growth.

Injectable delivery formulations may be administered intravenously or directly at the site in need of treatment. The injectable carrier may be a viscous solution or gel.

Delivery formulations include physiological saline, bacteriostatic saline (saline containing about 0.9% mg/mL benzyl alcohol), phosphate-buffered saline, Hank's solution, Ringer's-lactate, or liquids supplemented with albumin, methyl cellulose, or hyaluronic acid. Injectable matrices include polymers of poly(ethylene oxide) and copolymers of ethylene and propylene oxide (see Cao et al., *J. Biomater. Sci.* 9:475 (1998) and Sims et al., *Plast Reconstr. Surg.* 98:843 (1996), the entire teachings of which are incorporated herein by reference).

Other compositions which are injectable matrices include the solutions of poly(propylene fumarate) copolymers described above and pastes of calcium phosphate ceramics (see Schmitz et al., *J. Oral Maxillofacial Surgery* 57:1122 (1999), the entire teachings of which are incorporated herein by reference). Injectable matrices can be injected directly to the site in need of bone growth and can conveniently be used to fill voids and fuse bones without the need for invasive surgery.

Methods for encapsulating compositions (such as in a coating of hard gelatin or cyclodextran) are known in the art

(Baker, et al., "Controlled Release of Biological Active Agents", John Wiley and Sons, 1986).

Ointments are typically prepared using an oleaginous base, e.g., containing fixed oils or hydrocarbons, such as white petrolatum or mineral oil, or an absorbent base, e.g., consisting of an absorbent anhydrous substance or substances, for example anhydrous lanolin. Following formation of the base, the active ingredients are added in the desired concentration.

Creams generally comprise an oil phase (internal phase) containing typically fixed oils, hydrocarbons, and the like, such as waxes, petrolatum, mineral oil, and the like, and an aqueous phase (continuous phase), comprising water and any water-soluble substances, such as added salts. The two phases are stabilized by use of an emulsifying agent, for example, a surface active agent, such as sodium lauryl sulfate; hydrophilic colloids, such as acacia colloidal clays, beegum, and the like. Upon formation of the emulsion, the active ingredients are added in the desired concentration.

Gels are comprised of a base selected from an oleaginous base, water, or an emulsion-suspension base, as previously described. To the base is added a gelling agent which forms a matrix in the base, increasing its viscosity to a semisolid consistency. Examples of gelling agents are hydroxypropyl cellulose, acrylic acid polymers, and the like. The active ingredients are added to the formulation at the desired concentration at a point preceding addition of the gelling agent.

Diseases and conditions, treatable with thrombin peptide derivatives, for example, wounds and sites of angioplasty, are often accompanied by symptoms and infirmities such as pain and infection. In certain instances it may be advantageous to co-administer one or more additional pharmacologically active agents along with a thrombin peptide derivative to address such issues. For example, managing pain and inflammation, may require co-administration with analgesic or an anti-inflammatory agents. Managing infection may require co-administration with antimicrobial, antibiotic or disinfectant agents.

Thrombin peptide derivatives can be synthesized by solid phase peptide synthesis (e.g., BOC or FMOC) method, by solution phase synthesis, or by other suitable techniques including combinations of the foregoing methods. The BOC and FMOC methods, which are established and widely used, are described in Merrifield, *J. Am. Chem. Soc.* 88:2149 (1963); Meienhofer, *Hormonal Proteins and Peptides*, C. H. Li, Ed., Academic Press, 1983, pp. 48-267; and Barany and Merrifield, in *The Peptides*, E. Gross and J. Meienhofer, Eds., Academic Press, New York, 1980, pp. 3-285. Methods of solid phase peptide synthesis are described in Merrifield, R. B., *Science*, 232: 341 (1986); Carpino, L. A. and Han, G. Y., *J. Org. Chem.*, 37: 3404 (1972); and Gauspohl, H. et al., *Synthesis*, 5: 315 (1992)). The teachings of these six articles are incorporated herein by reference in their entirety.

The invention is illustrated by the following examples which are not intended to be limiting in any way.

EXEMPLIFICATION

Example 1

Biological Activity of TP508-Ala in Accelerating Wound Closure

Methodology and Study Design

The following experiment was conducted to determine the wound healing activity of the thrombin peptide derivative,

TP508-Ala, in which the cysteine residue at position 521 of TP508 was substituted with alanine.

Two, full-thickness, 2 cm-diameter excisions were created on the dorsum of male Sprague-Dawley rats. Both wounds on a given rat were treated with either saline containing TP508-Ala, saline containing TP508 (positive control) or saline alone (negative control). TP508-Ala and TP508 were administered at a dose of 0.1 μ g per wound. Wound size was determined on post-wounding days 3, 7, and 10 by tracing the perimeter of the wound onto an acetate sheet and using digital analysis to compute the surface area of each wound.

TP508-Ala was compared to the TP508 control and the saline control, yielding a total of three treatment groups. Each group contained 6 rats. The results indicated that the thrombin peptide derivative, TP508-Ala, is biologically active in accelerating wound closure.

Preparation of Treatment Solutions

Approximately 1 mg of lyophilized TP508 was dissolved in 1 mL of saline (sterile 0.9% sodium chloride injectable solution). Saline was used as the vehicle for the experiment. The stock solution of TP508 (1 mg/mL) was further diluted in vehicle to yield a working solution of 2.5 μ g/mL. The working solution was maintained on ice throughout the experiment.

Approximately 1 mg TP508-Ala was dissolved in 1 mL of saline (sterile 0.9% sodium chloride injectable solution). The stock solution of TP508-Ala (1 mg/mL) was further diluted in saline to yield a working solution of 2.5 μ g/mL. The working solution was maintained on ice throughout the experiment.

Wound Treatment

Both wounds on a given animal received the same treatment: a single, topical application of a 40 μ L volume containing saline alone, saline with TP508 (2.5 μ g/mL), or saline with TP508-Ala (2.5 μ g/mL).

Observation and Wound Size Analysis

The rats were observed for ten days following wounding, and no clinical signs of abnormal behavior, infection or toxicity were noted. On post-wounding days 3, 7, and 10, the wounds were evaluated by tracing the wound perimeter onto a flexible acetate sheet, then determining wound area with digital analysis software.

The results are presented in FIG. 1. FIG. 1 shows wound area measurements from post-wounding Days 7 and 10. No differences in wound size between the groups were present on post-wounding Day 3. Each data point represents the mean and standard error of the mean of 12 wounds from 6 rats. Statistical comparisons between groups were made using a repeated measures analysis of variance; Fisher's LSD was used for post hoc testing between groups.

At a dose of 0.1 μ g, TP508-Ala treated wound areas were 18.7% smaller than controls, while TP508 treated wound areas were 20.3% smaller than controls, by post-wounding Day 7. The same trend was observed at Day 10. At a dose of 0.1 μ g, TP508-Ala treated wound areas were 25.1% smaller than controls, while TP508 treated wound areas were 34.5% smaller than controls, by post-wounding Day 10. These data suggest that TP508-Ala is equivalent to TP508 in efficacy and potency in accelerating wound healing.

Example 2

Biological Activity of TP508-Ser in Accelerating Wound Closure

Methodology and Study Design

The following experiment was conducted to determine the wound healing activity of the thrombin peptide derivative, TP508-Ser, in which the cysteine residue at position 521 of TP508 was substituted with serine.

Two, full-thickness, 2 cm-diameter excisions were created on the dorsum of male Sprague-Dawley rats. Both wounds on a given rat were treated with either saline containing TP508-Ser, saline containing TP508 (positive control) or saline alone (negative control). TP508-Ser and TP508 were administered at a dose of 0.1 μ g per wound. Wound size was determined on post-wounding days 3, 7, and 10 by tracing the perimeter of the wound onto an acetate sheet and using digital analysis to compute the surface area of each wound.

TP508-Ser was compared to the TP508 control and the saline control, yielding a total of three treatment groups. Each group contained 8 rats. The results indicated that the thrombin peptide derivative, TP508-Ser, is biologically active in accelerating wound closure.

Preparation of Treatment Solutions

Approximately 1 mg of lyophilized TP508 was dissolved in 1 mL of saline (sterile 0.9% sodium chloride injectable solution). Saline was used as the vehicle for the experiment. The stock solution of TP508 (1 mg/mL) was further diluted in vehicle to yield a working solution of 2.5 μ g/mL. The working solution was maintained on ice throughout the experiment.

Approximately 1 mg TP508-Ser was dissolved in 1 mL of saline (sterile 0.9% sodium chloride injectable solution). The stock solution of TP508-Ser (1 mg/mL) was further diluted in saline to yield a working solution of 2.5 μ g/mL. The working solution was maintained on ice throughout the experiment.

Wound Treatment

Both wounds on a given animal received the same treatment: a single, topical application of a 40 μ L volume containing saline alone, saline with TP508 (2.5 μ g/mL), or saline with TP508-Ser (2.5 μ g/mL).

Observations and Wound Size Analysis

The rats were observed for ten days following wounding, and no clinical signs of abnormal behavior, infection or toxicity were noted. On post-wounding days 3, 7, and 10, the wounds were evaluated by tracing the wound perimeter onto a flexible acetate sheet, then determining wound area with digital analysis software.

The results are presented in FIG. 2. FIG. 2 shows wound area measurements from post-wounding Days 7 and 10. No differences in wound size between the groups were present on post-wounding Day 3. Each data point represents the mean and standard error of the mean of 16 wounds from 8 rats. Statistical comparisons between groups were made using a repeated measures analysis of variance; Fisher's LSD was used for post hoc testing between groups.

At a dose of 0.1 μ g, wound areas were reduced 18.2% by TP508-Ser and 16.7% by TP508 by post-wounding Day 7. The same trend was observed at Day 10 in which, at a dose of 0.1 μ g, wound areas were reduced 43.3% by TP508-Ser and 27.0% by TP508. These data suggest that TP508-Ser is equivalent to TP508 in efficacy and potency in accelerating wound healing.

Example 3

TP508-Dimer Formation

TP508 was dissolved in saline (sterile 0.9% sodium chloride injectable solution) at 5 mg/mL and incubated at 4° C. Over a time period of 6 months, triplicate samples were taken at intervals from the solution. The samples were analyzed by HPLC to separate TP508-monomer, TP508-dimer and unknowns.

The area percent of each HPLC peak was plotted in FIG. 3. The peak area percent corresponds directly to the percent of material in solution. The peak area of TP508-monomer decreases over time whereas the peak area of TP508-dimer

increases over time. No increase in the unknown peaks was observed. The results of FIG. 3 show that TP508 converts to dimer over time.

Example 4

Stability of TP508-Ala

TP508

TP508 was dissolved in saline (sterile 0.9% sodium chloride injectable solution), at 5 mg/mL and incubated at 4° C. Over a time period of 1 month, triplicate samples were taken at intervals from the solution. The samples were analyzed by HPLC to separate TP508-monomer, TP508-dimer and unknowns. The area percent of each HPLC peak was plotted in FIG. 4. The peak area percent corresponds directly to the percent of material in solution. The peak area of TP508-monomer decreases over time whereas the peak area of TP508-dimer increases over time. No increase in the

unknown peaks was observed. The results of FIG. 4 show that TP508 converts to dimer over time.

TP508-Ala

TP508-Ala was dissolved in saline (sterile 0.9% sodium chloride injectable solution) at 5 mg/mL and incubated at 4° C. Over a time period of 1 month, triplicate samples were taken at intervals from the solution. The samples were analyzed by HPLC to separate TP508-Ala and unknown. The area percent of each HPLC peak was plotted in FIG. 5. The peak area percent corresponds directly to the percent of material in solution. The peak area of TP508-Ala showed no decrease over time. No increase in the unknown peak was observed. The results show that TP508-Ala does not convert to dimer.

While this invention has been particularly shown and described with references to preferred embodiments thereof, it will be understood by those skilled in the art that various changes in form and details may be made therein without departing from the scope of the invention encompassed by the appended claims.

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 <223> OTHER INFORMATION: Thrombin Peptide Derivative

<400> SEQUENCE: 3

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Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Cys Glu Gly
 1 5 10 15

Asp Ser Gly Gly Pro Phe Val
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<210> SEQ ID NO 4
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 <222> LOCATION: 6
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 <222> LOCATION: 13
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<400> SEQUENCE: 4

Arg Gly Asp Ala Xaa Xaa Gly Asp Ser Gly Gly Pro Xaa Val
 1 5 10

<210> SEQ ID NO 5
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 <220> FEATURE:
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<400> SEQUENCE: 5

Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Xaa Glu Gly
 1 5 10 15

Asp Ser Gly Gly Pro Phe Val
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<210> SEQ ID NO 6
 <211> LENGTH: 23
 <212> TYPE: PRT
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 <223> OTHER INFORMATION: Xaa = Phe, Met, Leu, His or Val
 <220> FEATURE:
 <223> OTHER INFORMATION: Thrombin Peptide Derivative

<400> SEQUENCE: 6

Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Xaa Xaa Gly
 1 5 10 15

Asp Ser Gly Gly Pro Xaa Val
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<210> SEQ ID NO 7
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<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Thrombin Peptide Derivative

<400> SEQUENCE: 7

Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Ala Glu Gly
 1           5           10          15
Asp Ser Gly Gly Pro Phe Val
 20

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<210> SEQ ID NO 8
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<220> FEATURE:
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<223> OTHER INFORMATION: Thrombin Peptide Derivative

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Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Ala Glu Gly
 1           5           10          15
Asp Ser Gly Gly Pro Phe Val
 20

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<223> OTHER INFORMATION: Xaa = Ala, Gly, Ser or S-protected Cys
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<400> SEQUENCE: 9

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Asp Asn Met Phe Xaa Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly
 1           5           10          15
Asp Ala Xaa Glu Gly Asp Ser Gly Gly Pro Phe Val Met Lys Ser Pro
 20          25          30
Phe

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<222> LOCATION: 27
<223> OTHER INFORMATION: Phe, Met, Leu, His or Val
<220> FEATURE:
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Asp Asn Met Phe Xaa Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly
 1           5           10           15

Asp Ala Xaa Xaa Gly Asp Ser Gly Gly Pro Xaa Val Met Lys Ser Pro
 20           25           30

Phe

<210> SEQ ID NO 11
<211> LENGTH: 33
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Asp Asn Met Phe Ala Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly
 1           5           10           15

Asp Ala Ala Glu Gly Asp Ser Gly Gly Pro Phe Val Met Lys Ser Pro
 20           25           30

Phe

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<400> SEQUENCE: 12

Asp Asn Met Phe Ala Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly
 1           5           10           15

Asp Ala Ala Glu Gly Asp Ser Gly Gly Pro Phe Val Met Lys Ser Pro
 20           25           30

Phe

<210> SEQ ID NO 13
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<221> NAME/KEY: VARIANT
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<223> OTHER INFORMATION: Xaa = Phe, Met, Leu, His or Val
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<223> OTHER INFORMATION: valine is unsubstituted or amidated as -NH2
<223> OTHER INFORMATION: Thrombin Peptide Derivative

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<400> SEQUENCE: 13

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Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Xaa Xaa Gly
  1             5             10             15

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Asp Ser Gly Gly Pro Xaa Val
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<400> SEQUENCE: 14

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Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Ser Glu Gly
  1             5             10             15

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Asp Ser Gly Gly Pro Phe Val
      20

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<210> SEQ ID NO 15
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<223> OTHER INFORMATION: Xaa = Ala, Gly, Ser or S-protected Cys
<220> FEATURE:
<221> NAME/KEY: VARIANT
<222> LOCATION: 23
<223> OTHER INFORMATION: valine is unsubstituted or amidated as -NH2
<220> FEATURE:
<223> OTHER INFORMATION: Thrombin Peptide Derivative

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<400> SEQUENCE: 15

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Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Xaa Glu Gly
  1             5             10             15

```

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Asp Ser Gly Gly Pro Phe Val
      20

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<223> OTHER INFORMATION: Thrombin Peptide

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Arg Gly Asp Ala
1

<210> SEQ ID NO 17

<211> LENGTH: 12

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<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Thrombin Peptide Derivative

<400> SEQUENCE: 17

Asp Ala Cys Glu Gly Asp Ser Gly Gly Pro Phe Val
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<210> SEQ ID NO 18

<211> LENGTH: 4

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Fibronectin Specific Peptide

<400> SEQUENCE: 18

Arg Gly Ala Ser
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<210> SEQ ID NO 19

<211> LENGTH: 23

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Thrombin Peptide Derivative

<400> SEQUENCE: 19

Ala Gly Tyr Lys Pro Asp Glu Gly Lys Arg Gly Asp Ala Ser Glu Gly
1 5 10 15

Asp Ser Gly Gly Pro Phe Val
20

What is claimed is:

1. A polypeptide, wherein the amino acid sequence of said polypeptide is Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val (SEQ ID NO: 5) or a fragment of said polypeptide, wherein said fragment comprises the amino acid sequence Arg-Gly-Asp-Ala-Xaa-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val (SEQ ID NO: 2), wherein said polypeptide or said fragment comprises an N-terminus which is unsubstituted and a C-terminus which is unsubstituted or which comprises an amide represented by $-C(O)NH_2$ and wherein Xaa is an S-protected cysteine, the thiol of which is blocked with a protecting group, provided that the protecting group is not a second polypeptide comprising SEQ ID NO: 2.

2. A polypeptide, wherein the amino acid sequence of said polypeptide is Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-Xi-Gly-Asp-Ser-Gly-Gly-Pro-X₂-Val (SEQ ID NO: 6) or a fragment of said polypeptide, wherein said fragment of said polypeptide comprises Arg-Gly-Asp-Ala-Xaa-X₁-Gly-Asp-Ser-Gly-Gly-Pro-X₂-Val (SEQ ID NO: 4); wherein Xaa is an S-protected cysteine, the thiol of

45 which is blocked with a protecting group, provided that the protecting group is not a second polypeptide comprising SEQ ID NO:4; and wherein X₁ is Glu or Gln; and X₂ is Phe, Met, Leu, His or Val; and wherein the polypeptide or said fragment comprises an N-terminus which is unsubstituted, and a C-terminus which is unsubstituted or which comprises an amide represented by $-C(O)NH_2$.

3. The polypeptide of claim 2, wherein X₁ is Glu and X₂ is Phe.

4. The polypeptide H-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-X₁-Gly-Asp-Ser-Gly-Gly-Pro-X₂-Val-R1 (SEQ ID NO: 13), wherein X₁ is Glu or Gln and X₂ is Phe, Met, Leu, His or Val; wherein Xaa is an S-protected cysteine protected as a thioether or thioester; and wherein R1 is $-OH$ or $-NH_2$.

5. The polypeptide H-Ala-Gly-Tyr-Lys-Pro-Asp-Glu-Gly-Lys-Arg-Gly-Asp-Ala-Xaa-Glu-Gly-Asp-Ser-Gly-Gly-Pro-Phe-Val-R1 (SEQ ID NO: 15); wherein Xaa is an S-protected cysteine protected as a thioether or thioester; and wherein R1 is $-OH$ or $-NH_2$.

* * * * *

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 7,713,934 B2
APPLICATION NO. : 11/026521
DATED : May 11, 2010
INVENTOR(S) : Darrell H. Carney

Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

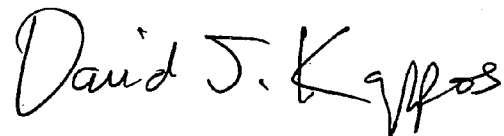
In claim 2, Column 25, Line 60, change “Lvs” to “Lys”;

In claim 2, Column 25, Line 61, change “Gly” to “Gly” in two instances; and

In claim 2, Column 25, Line 61, change “Xi” to “X₁”.

Signed and Sealed this

Twenty-eighth Day of September, 2010

A handwritten signature in black ink that reads "David J. Kappos". The signature is written in a cursive style with a large, stylized 'D' and 'K'.

David J. Kappos
Director of the United States Patent and Trademark Office

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

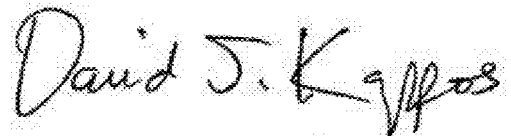
PATENT NO. : 7,713,934 B2
APPLICATION NO. : 11/026521
DATED : May 11, 2010
INVENTOR(S) : Darrell H. Carney

Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Title Page, Item (73) Assignee should be listed as Orthologic Corp., Tempe, AZ.

Signed and Sealed this
Fourteenth Day of June, 2011

A handwritten signature in black ink that reads "David J. Kappos". The signature is written in a cursive style with a large initial "D".

David J. Kappos
Director of the United States Patent and Trademark Office