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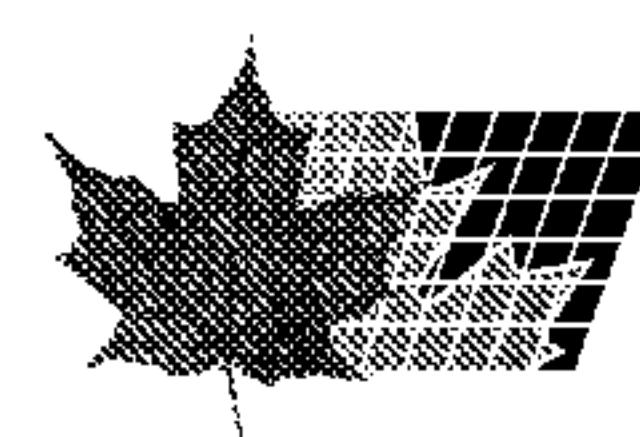
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(54) Titre : TRAITEMENT D'INFECTIONS ET D'AUTRES TROUBLES
(54) Title: TREATMENT OF INFECTIONS AND OTHER DISORDERS

(57) Abrégé/Abstract:

Microbial infections including anthrax infection, and gastrointestinal disorders, are treated or prevented by administration of an actin-sequestering peptide including amino acid sequence LKKTET, such as Thymosin β 4, an isoform of Thymosin β 4, oxidized Thymosin β 4, or T β 4 sulfoxide.



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(54) Title: TREATMENT OF INFECTIONS AND OTHER DISORDERS

(57) Abstract: Microbial infections including anthrax infection, and gastrointestinal disorders, are treated or prevented by administration of an actin-sequestering peptide including amino acid sequence LKKTET, such as Thymosin β 4, an isoform of Thymosin β 4, oxidized Thymosin β 4, or T β 4 sulfoxide.

TREATMENT OF INFECTIONS AND OTHER DISORDERS

BACKGROUND OF THE INVENTION

5 CROSS-REFERENCE TO RELATED APPLICATION

The present application claims the benefit of U.S. Provisional Application Serial Nos. 60/421,038, filed October 25, 2002 and 60/354,250, filed February 6, 2002.

10 1. FIELD OF THE INVENTION

The present invention relates to the field of the treatment of microbial infections and other gastrointestinal disorders.

15 2. DESCRIPTION OF THE BACKGROUND ART

Treatment of microbial infections, bacterial, viral and fungal, can be difficult by conventional methods. These infections may include gastrointestinal infections (E. coli., H. pylori, VRE, etc.) abdominal infections (peritonitis, pancreatitis, gall bladder infections, etc.), surgical infections and osteomyelitis (bone infection).

Another example of microbial infection is anthrax. Anthrax is an infectious agent caused by Bacillus anthracis, a gram positive organism. It is primarily a disease of herbivores. Anthrax can effect many different vertebrates including humans. The symptoms of anthrax vary widely depending on the route of infection. Three forms of the disease commonly occur including a cutaneous form, a gastrointestinal form, and a pulmonary (inhalation) form. The pulmonary form of the disease is typically caused by inhalation of anthrax spores. The symptoms systemic anthrax can be mimicked in a number of animal models by the administration of virulence factors (endotoxins) which are responsible for the major pathologies, morbidity and mortality seen with anthrax. Once large amounts of anthrax toxins are produced within the body by bacteria, administration of antibiotics are usually ineffective. Anthrax induced pathologies mimic septic shock and the sudden death seen with other gram positive and gram negative bacterial infections such as multi-organ failure, edema and ARDS. In both animals and humans the anthrax induced pathologies also include marked elevation of TNF α , IL-1 β , PAF and a number of other inflammatory cytokines. Also seen in anthrax induced septic shock is over production of reactive oxygen intermediates and an increase in arachidonic acid metabolites such as PGE, and thromboxane β_2 and disruption of the actin cytoskeleton.

A number of approaches have been reported to delay, prevent and/or treat exposure to anthrax. In the prevention area, a human vaccine is available by the effectiveness of the vaccine is unclear. The best treatment currently available is treatment with specific antibiotics

such as ciprofloxacin or doxycycline. Antibiotics are effective if given at the very early stages of infection and are basically ineffective once the bacteria have had a chance to multiply rapidly producing lethal amounts of the deadly anthrax toxins. Of the three forms of anthrax, the most deadly form is pulmonary (Inhalation) anthrax which has a fatality rate of greater than 75% (even with appropriate antibiotic treatment). Anthrax produces a multi-component toxin that is assembled at the surface of host cells after infection. The lethal action of the anthrax toxins occurs in the cytoplasm of the host cells. The anthrax toxin is only one of many multi-subunit toxins that cause severe illness in humans. A major concern when treating bacterial infections with antibiotics is the appearance of increasing numbers of antibiotic resistant strains. In addition, once the anthrax bacillus has produced large amounts of exotoxins the antibiotics are basically ineffective.

Millions of Americans suffer from other gastrointestinal (GI) disorders such as colitis, ileitis, Crohn's disease, ulcerative colitis, colic, gingivitis, regional enteritis, ulcers, pouchitis, sclerosing, cholangitis, fistulae. The cause of many of these diseases is not known. However, they may have genetic roots or result from exposure to certain chemicals, pathogens, immune dysfunction, or foods during one's lifetime, or result from the normal aging of the human body. GI disorders occur in both men and women and can be acute or chronic, debilitating and life-threatening, and may occur anywhere within the GI tract, including but not limited to the mouth, throat, esophagus, stomach, small and large intestines, colon, and anus. People suffering from GI disorders may have a greatly diminished quality of life and suffer premature death.

A large number of therapeutic approaches to treatment have been reported for gastrointestinal disorders and disease, depending upon the location and the nature of the GI pathology. The treatments vary from surgical intervention, to dietary manipulations, to the use of a variety of drugs and biological agents. These agents include antibiotics, anti-virals, anti-inflammatory drugs, glucocorticoids, immunosuppressive drugs, monoclonal antibodies, antacids, anti-secretory drugs, anti-spasmodics, as well as a large number of others.

Numerous pharmaceutical, nutriceutical or cosmeceutical formulations have been proposed to treat the damage caused by microbial infections and gastrointestinal disorders.

There remains a need in the art for improved methods and compositions for healing or preventing the damage caused by microbial infections and gastrointestinal disorders.

SUMMARY OF THE INVENTION

In accordance with the present invention, treatment of infections and gastrointestinal (GI) disorders, comprises administering to a subject in need of such treatment an effective amount of a composition comprising amino acid sequence LKKTET, or a conservative variant thereof.

DETAILED DESCRIPTION OF THE INVENTION

The present invention is based on a discovery that actin-sequestering peptides such as thymosin β 4 (T β 4) and other actin-sequestering peptides or peptide fragments containing amino acid sequence LKKTET or conservative variants thereof, promote treatment of microbial infections and gastrointestinal disorders. Without being bound to any particular theory, these peptides may have the capacity to promote repair, healing and prevention by having the ability to induce terminal deoxynucleotidyl transferase (a non-template directed DNA polymerase), to decrease the levels of one or more inflammatory cytokines or chemokines, and to act as a chemotactic and/or angiogenic factor for endothelial cells and thus treat damage caused by microbial infections and gastrointestinal disorders.

Thymosin β 4 was initially identified as a protein that is up-regulated during endothelial cell migration and differentiation *in vitro*. Thymosin β 4 was originally isolated from the thymus and is a 43 amino acid, 4.9 kDa ubiquitous polypeptide identified in a variety of tissues. Several roles have been ascribed to this protein including a role in a endothelial cell differentiation and migration, T cell differentiation, actin sequestration and vascularization.

In accordance with one embodiment, the invention is a method of treatment of damage associated with microbial infections comprising administering to a subject in need of such treatment an effective amount of a composition comprising a microbial infection-inhibiting polypeptide comprising LKKTET, or a conservative variant thereof having microbial infection-inhibiting activity, preferably Thymosin β 4, an isoform of Thymosin β 4, oxidized Thymosin β 4, Thymosin β 4 sulfoxide, or an antagonist of Thymosin β 4.

Compositions which may be used in accordance with the present invention include Thymosin β 4 (T β 4), T β 4 isoforms, oxidized T β 4, Thymosin β 4 sulfoxide, polypeptides or any other actin sequestering or bundling proteins having actin binding domains, or peptide fragments comprising or consisting essentially of the amino acid sequence LKKTET or conservative variants thereof, having microbial infection-inhibiting activity. International Application Serial No. PCT/US99/17282, incorporated herein by reference, discloses isoforms of T β 4 which may be useful in accordance with the present invention as well as amino acid sequence LKKTET and conservative variants thereof having microbial infection-inhibiting activity, which may be utilized with the present invention. International Application Serial No. PCT/GB99/00833 (WO 99/49883), incorporated herein by reference, discloses oxidized Thymosin β 4 which may be utilized in accordance with the present invention. Although the present invention is described primarily hereinafter with respect to T β 4 and T β 4 isoforms, it is to be understood that the following description is intended to be equally applicable to amino acid sequence LKKTET, peptides and fragments comprising or consisting essentially of LKKTET, conservative variants thereof having microbial infection-inhibiting activity, as well as oxidized

Thymosin β 4.

In one embodiment, the invention provides a method for healing damage caused by microbial infection in a subject by contacting an area to be treated with an effective amount of a microbial infection-inhibiting composition which contains $\text{T}\beta 4$ or a $\text{T}\beta 4$ isoform. The contacting may be topically, systemically or enterally. Examples of topical administration include, for example, contacting the skin with a lotion, salve, gel, cream, paste, spray, suspension, dispersion, hydrogel, ointment, or oil comprising $\text{T}\beta 4$, alone or in combination with at least one agent that enhances $\text{T}\beta 4$ penetration, or delays or slows release of $\text{T}\beta 4$ peptides into the area to be treated. Systemic administration includes, for example, intravenous, intraperitoneal, intramuscular or subcutaneous injections, or inhalation, transdermal or oral administration of a composition containing $\text{T}\beta 4$ or a $\text{T}\beta 4$ isoform, etc. Enteral administration may include oral or rectal administration. A subject may be a mammal, preferably human.

$\text{T}\beta 4$, or its analogues, isoforms or derivatives, may be administered in any effective amount. For example, $\text{T}\beta 4$ may be administered in dosages within the range of about 0.1-50 micrograms of $\text{T}\beta 4$, more preferably in amounts of about 1-25 micrograms. A composition in accordance with the present invention can be administered daily, every other day, etc., with a single administration or multiple administrations per day of administration, such as applications 2, 3, 4 or more times per day of administration.

$\text{T}\beta 4$ isoforms have been identified and have about 70%, or about 75%, or about 80% or more homology to the known amino acid sequence of $\text{T}\beta 4$. Such isoforms include, for example, $\text{T}\beta 4^{\text{ala}}$, $\text{T}\beta 9$, $\text{T}\beta 10$, $\text{T}\beta 11$, $\text{T}\beta 12$, $\text{T}\beta 13$, $\text{T}\beta 14$ and $\text{T}\beta 15$. Similar to $\text{T}\beta 4$, the $\text{T}\beta 10$ and $\text{T}\beta 15$ isoforms have been shown to sequester actin. $\text{T}\beta 4$, $\text{T}\beta 10$ and $\text{T}\beta 15$, as well as these other isoforms share an amino acid sequence, LKKTET, that appears to be involved in mediating actin sequestration or binding. Although not wishing to be bound to any particular theory, the activity of $\text{T}\beta 4$ isoforms may be due, in part, to the ability to regulate the polymerization of actin. For example, $\text{T}\beta 4$ can modulate actin polymerization in skin (e.g. β -thymosins appear to depolymerize F-actin by sequestering free G-actin). $\text{T}\beta 4$'s ability to modulate actin polymerization may therefore be due to all, or in part, its ability to bind to or sequester actin via the LKKTET sequence. Thus, as with $\text{T}\beta 4$, other proteins which bind or sequester actin, or modulate actin polymerization, including $\text{T}\beta 4$ isoforms having the amino acid sequence LKKTET, are likely to reduce microbial infection alone or in a combination with $\text{T}\beta 4$, as set forth herein.

Thus, it is specifically contemplated that known $\text{T}\beta 4$ isoforms, such as $\text{T}\beta 4^{\text{ala}}$, $\text{T}\beta 9$, $\text{T}\beta 10$, $\text{T}\beta 11$, $\text{T}\beta 12$, $\text{T}\beta 13$, $\text{T}\beta 14$ and $\text{T}\beta 15$, as well as $\text{T}\beta 4$ isoforms not yet identified, will be useful in the methods of the invention. As such $\text{T}\beta 4$ isoforms are useful in the methods of the invention, including the methods practiced in a subject. The invention therefore further provides

pharmaceutical compositions comprising T β 4, as well as T β 4 isoforms T β 4^{ala}, T β 9, T β 10, T β 11, T β 12, T β 13, T β 14 and T β 15, and a pharmaceutically acceptable carrier.

In addition, other proteins having actin sequestering or binding capability, or that can mobilize actin or modulate actin polymerization, as demonstrated in an appropriate 5 sequestering, binding, mobilization or polymerization assay, or identified by the presence of an amino acid sequence that mediates actin binding, such as LKKTET, for example, can similarly be employed in the methods of the invention. Such proteins include gelsolin, vitamin D binding protein (DBP), profilin, cofilin, adsevertin, propomyosin, fincillin, depactin, DnaseI, vilin, fragmin, severin, capping protein, β -actinin and acumentin, for example. As such methods include those 10 practiced in a subject, the invention further provides pharmaceutical compositions comprising gelsolin, vitamin D binding protein (DBP), profilin, cofilin, depactin, DnaseI, vilin, fragmin, severin, capping protein, β -actinin and acumentin as set forth herein. Thus, the invention includes the use of a microbial infection-inhibiting polypeptide comprising the amino acid 15 sequence LKKTET (which may be within its primary amino acid sequence) and conservative variants thereof.

As used herein, the term "conservative variant" or grammatical variations thereof denotes the replacement of an amino acid residue by another, biologically similar residue. Examples of conservative variations include the replacement of a hydrophobic residue such as isoleucine, valine, leucine or methionine for another, the replacement of a polar residue for 20 another, such as the substitution of arginine for lysine, glutamic for aspartic acids, or glutamine for asparagine, and the like.

T β 4 has been localized to a number of tissue and cell types and thus, agents which stimulate the production of T β 4 can be added to or comprise a composition to effect T β 4 production from a tissue and/or a cell. Such agents include members of the family of growth 25 factors, such as insulin-like growth factor (IGF-1), platelet derived growth factor (PDGF), epidermal growth factor (EGF), transforming growth factor beta (TGF- β), basic fibroblast growth factor (bFGF), thymosin α 1 (T α 1) and vascular endothelial growth factor (VEGF). More preferably, the agent is transforming growth factor beta (TGF- β) or other members of the TGF- β superfamily. T β 4 compositions of the invention may reduce the affects of microbial infection 30 by effectuating growth of the connective tissue through extracellular matrix deposition, cellular migration and vascularization of the skin.

Additionally, other agents may be added to a composition along with T β 4 or a T β 4 isoform. Such agents include angiogenic agents, growth factors, agents that direct 35 differentiation of cells, agents that promote migration of cells and agents that stimulate the provision of extracellular matrix material in the skin. For example, and not by way of limitation, T β 4 or a T β 4 isoform alone or in combination can be added in combination with any one or

more of the following agents: VEGF, KGF, FGF, PDGF, TGF β , IGF-1, IGF-2, IL-1, prothymosin α and thymosin α 1 in an effective amount.

The actual dosage or reagent, formulation or composition that heals damage associated with microbial infection may depend on many factors, including the size and health of a subject. 5 However, persons of ordinary skill in the art can use teachings describing the methods and techniques for determining clinical dosages as disclosed in PCT/US99/17282, *supra*, and the references cited therein, to determine the appropriate dosage to use.

Suitable formulations include T β 4 or a T β 4 isoform at a concentration within the range of about 0.001 - 10% by weight, more preferably within the range of about 0.01 - 0.1% by weight, 10 most preferably about 0.05% by weight.

The therapeutic approaches described herein involve various routes of administration or delivery of reagents or compositions comprising the T β 4 or other compounds of the invention, including any conventional administration techniques (for example, but not limited to, topical administration, local injection, inhalation, systemic or enteral administration), to a subject. The 15 methods and compositions using or containing T β 4 or other compounds of the invention may be formulated into pharmaceutical compositions by admixture with pharmaceutically acceptable non-toxic excipients or carriers.

The invention includes use of antibodies which interact with T β 4 peptide or functional fragments thereof. Antibodies which include pooled monoclonal antibodies with different 20 epitopic specificities, as well as distinct monoclonal antibody preparations are provided.

Monoclonal antibodies are made from antigen containing fragments of the protein by methods well known to those skilled in the art as disclosed in PCT/US99/17282, *supra*. The term antibody as used in this invention is meant to include monoclonal and polyclonal antibodies.

In one embodiment, the invention provides a method for treating bacterial infection 25 comprising administering to a subject in need of such treatment, an effective amount of a composition comprising a bacterial infection-inhibiting polypeptide comprising amino acid sequence LKKTET, or a conservative variant thereof having bacterial infection-inhibiting activity.

In another embodiment, the invention provides a method for treating gastrointestinal infection. Common gastrointestinal infections include, but are not limited to *Helicobacter pylori* 30 (*H. pylori*), *Escherichia coli* (*E. coli*), vancomycin-resistant *Enterococcus faecalis* (VRE), and methicillin-resistant *Staphylococcus aureus* (MRSA). The composition may be delivered systemically by injection, orally, nasally, through suppository or enema, transdermally or any other suitable means.

In another embodiment, the invention provides a method for treating anthrax infection. 35 Damage caused by anthrax infection includes septic shock, sudden death, multi-organ failure, edema, ARDS and inflammatory, degenerative, immunological damage. The composition can

be applied alone or in combination with an antibiotic such as ciprofloxacin, doxycyclin, or penicillin. A therapeutically effective amount of the composition is applied to the site or systemically on a periodic basis during a course of therapy to reduce the mortality and morbidity effects of exposure to biological agents such as anthrax or to prevent such effects. The 5 composition may also be delivered systemically by injection, orally, nasally or any other means to reduce the toxicity of pulmonary or gastrointestinal anthrax.

Another aspect of the invention is treatment of other Gastrointestinal disorders. In accordance with one embodiment, the invention is a method of treatment of damage associated 10 with gastrointestinal disorders comprising administering to a subject in need of such treatment an effective amount of a composition comprising a gastrointestinal disorder-inhibiting polypeptide comprising LKKTET, or a conservative variant thereof having gastrointestinal disorder inhibiting activity. This invention is applicable to inflammatory, ulcerative, degenerative, immunological and other injuries to and disorders of the gastrointestinal tract (from the mouth to the anus). These disorders occur due to genetic abnormalities, food 15 intolerance, chemical exposure, aging, and microbial infections.

Gastrointestinal disorders to which the invention is applicable include, but are not limited to, gastrointestinal infections including bacterial, viral and fungal infections, disorders associated with environmental or iatrogenic abrasions, inflammations and other inflammatory disorders, immunological disorders, allergies including food allergies, Crohn's disease, 20 ulcerative colitis, recurrent aphthous stomatitis (recurrent canker sores), ileitis, colic, gingivitis, regional enteritis, ulcers, pouchitis, sclerosing, cholangitis, fistulae and genetic abnormalities. They may result from exposure to certain chemicals, pathogens, immune dysfunction, or foods during one's lifetime, or result from the normal aging of the human body.

In one embodiment, the invention provides a method for healing damage caused by 25 gastrointestinal disorders in a subject by contacting the skin with a gastrointestinal disorder effective amount of a composition which contains T β 4 or a T β 4 isoform. The contacting may be topically, enterally or systemically. Examples of topical administration include, for example, contacting the skin with a lotion, salve, gel, cream, paste, spray, suspension, dispersion, hydrogel, ointment, or oil comprising T β 4, alone or in combination with at least one agent that 30 enhances T β 4 penetration, or delays or slows release of T β 4 peptides into the area to be treated. Systemic administration includes, for example, intravenous, intraperitoneal, intramuscular or subcutaneous injections, or inhalation (orally or nasally), transdermal, suppository, enema or oral administration of a composition containing T β 4 or a T β 4 isoform, etc. A subject may be a mammal, preferably human.

The invention also is directed to a substance for use in manufacture of a medicament 35 for treatment of microbial infections including anthrax, and gastrointestinal disorders, comprising amino acid sequence LKKTET, or a conservative variant thereof.

Example

T β 4 showed antimicrobial activity against Staphylococcus aureus and Escherichia coli at concentrations of 5-20 n mol/ml, and increasingly dose-dependent activity against those microorganisms at concentrations of 50-200 n mol/ml.

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<211> 6

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<213> Homo sapiens

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Leu Lys Lys Thr Glu Thr
1 5

CLAIMS

5 1. A method of treatment of microbial infections comprising administering to a subject in need of such treatment an effective amount of a composition comprising a microbial infection-inhibiting polypeptide comprising amino acid sequence LKKTET, or a conservative variant thereof, having microbial infection-inhibiting activity.

2. The method of claim 1 wherein said polypeptide comprises Thymosin β 4 (T β 4), an N-terminal variant of T β 4, a C-terminal variant of T β 4, an isoform of T β 4, oxidized T β 4 or T β 4 sulfoxide.

3. The method of claim 1 wherein said composition is administered systemically.

10 4. The method of claim 1 wherein said composition is administered topically.

5. The method of claim 1 wherein said composition is administered enterally.

6. The method of claim 1 wherein said microbial infection is a bacterial infection.

7. The method of claim 1 wherein said microbial infection is an anthrax infection.

15 8. The method of claim 1 wherein said microbial infection is a gastrointestinal infection.

9. A method of treatment of gastrointestinal disorders comprising administering to a subject in need of such treatment an effective amount of a composition comprising a gastrointestinal disorder-inhibiting polypeptide comprising amino acid sequence LKKTET, or a conservative variant thereof having gastrointestinal disorder-inhibiting activity.

20 10. The method of claim 9 wherein said gastrointestinal disorder is associated with infection, abrasion, inflammation, allergy, immunological disorder, or genetic abnormality.

11. The method of claim 9 wherein said gastrointestinal disorder is Crohn's disease.

12. The method of claim 9 wherein said gastrointestinal disorder is ulcerative colitis.

25 13. The method of claim 9 wherein said gastrointestinal disorder is recurrent aphthous stomatitis.

14. A substance for use in manufacture of a medicament for treatment of microbial infections, comprising amino acid sequence LKKTET, or a conservative variant thereof, having microbial infection-inhibiting activity.

30 15. A substance for use in manufacture of a medicament for treatment of gastrointestinal disorders, comprising amino acid sequence LKKTET, or a conservative variant thereof, having gastrointestinal disorder-inhibiting activity.