USE OF OCTREOTIDE AS A THERAPEUTIC AGENT

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Appl. No.: 12/676,926
PCT Filed: Sep. 9, 2008
PCT No.: PCT/EP2008/007435
§ 371 (c)(1), (2), (4) Date: Mar. 8, 2010

Foreign Application Priority Data
Sep. 11, 2007 (EP) 07017752.2

Publication Classification
Int. Cl.
A61K 38/08 (2006.01)
C07K 7/06 (2006.01)

U.S. Cl. ........................................ 514/16; 530/329

ABSTRACT

The present invention is directed to the use of the peptide compound H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-L-threoninol as a therapeutic agent for the prophylaxis and/or treatment of cancer, autoimmune diseases, fibrotic diseases, inflammatory diseases, neurodegenerative diseases, infectious diseases, lung diseases, heart and vascular diseases and metabolic diseases. Moreover the present invention relates to pharmaceutical compositions preferably in the form of a lyophilisate or liquid buffer solution or artificial mother milk formulation or mother milk substitute containing the peptide H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-L-threoninol optionally together with at least one pharmaceutically acceptable carrier, cryoprotectant, lyoprotectant, excipient and/or diluent.
USE OF OCTREOTIDE AS A THERAPEUTIC AGENT

[0001] The present invention is directed to the use of the peptide compound H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-L-threoninol (Octreotide) as a therapeutic agent for the prophylaxis and/or treatment of cancer, an infectious disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, an autoimmune disease, or a heart and vascular disease.

BACKGROUND OF THE INVENTION

[0002] The identification of a therapeutic compound effective for the prophylaxis and/or treatment of a disease can be based on the activity of the compound in a biological assay. A biological assay that mimics a disease causative mechanism can be used to test the therapeutic activity of a candidate peptide.

[0003] The causative mechanism of many diseases is the over activity of a biological pathway. A peptide that can reduce the activity of the biological pathway can be effective in the prophylaxis and/or treatment of the disease caused by the over activity of the biological pathway. Similarly, the causative mechanism of many diseases is the over production of a biological molecule. A peptide that can reduce the production of the biological molecule or block the activity of the over produced biological molecule can be effective in the prophylaxis and/or treatment of the disease caused by the over production of the biological molecule.

[0004] Conversely, the causative mechanism of many diseases is the under activity of a biological pathway. A peptide that can increase the activity of the biological pathway can be effective in the prophylaxis and/or treatment of the disease caused by the under activity of the biological pathway. Similarly, the causative mechanism of many diseases is the under production of a biological molecule. A peptide that can increase the production of the biological molecule or mimic the biological activity of the under produced biological molecule can be effective in the prophylaxis and/or treatment of the disease caused by the under production of the biological molecule.

[0005] It is the object of the present invention to provide a peptide compound for the prophylaxis and/or treatment of cancer, autoimmune diseases, fibrotic diseases, inflammatory diseases, neurodegenerative diseases, infectious diseases, lung diseases, heart and vascular diseases and metabolic diseases.

[0006] The object of the present invention is solved by the teaching of the independent claims. Further advantageous features, aspects and details of the invention are evident from the dependent claims, the description, and the examples of the present application.

DESCRIPTION OF THE INVENTION

[0007] The present invention relates to the use of the H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-L-threoninol (Octreotide), its use as a therapeutic in medicine and for the prophylaxis and/or treatment of cancer, autoimmune diseases, fibrotic diseases, inflammatory diseases, neurodegenerative diseases, infectious diseases, lung diseases, heart and vascular diseases and metabolic diseases. Also disclosed are pharmaceutical formulations preferably in form of a lyophilized or liquid buffer solution or artificial mother milk formulation containing the inventive peptide. The peptide is especially useful for prophylaxis and/or treatment of Mycobacterium tuberculosis infection, Streptococcus pneumoniae infection, Streptococcus pneumoniae infection related diseases, hemolytic uremic syndrome, pneumonia, meningitis, cystic fibrosis complication, middle ear diseases, infections of the bloodstream and other diseases which are described in the following.

[0008] Cancer, Tumors, Proliferative Diseases, Malignancies and their Metastases


[0010] The peptide of the present invention was tested using the assays described in Examples 1-7, 9-17 for their effect as active therapeutic agents in the prophylaxis and/or treatment of cancer, proliferative diseases, tumors and their metastases.

Infectious Disease

[0011] The immune system in higher vertebrates represents the first line of defense against various antigens that can enter
the vertebrate body, including microorganisms such as bacteria, fungi and viruses that are the causative agents of a variety of diseases.

Despite large immunization programs, viral infections, such as influenza virus, human immunodeficiency virus (“HIV”), herpes simplex virus (“HSV”, type 1 or 2), human papilloma virus (“HPV”, type 16 or 18), human cytomegalovirus (“HCMV”) or human hepatitis B or C virus (“HBV”, Type B; “HCV”, type C) infections, remain a serious source of morbidity and mortality throughout the world and a significant cause of illness and death among people with immune-deficiency associated with aging or different clinical conditions. Although antiviral chemotherapy with compounds such as amantadine and rimantadine have been shown to reduce the duration of symptoms of clinical infections (i.e., influenza infection), major side effects and the emergence of drug-resistant variants have been described. New classes of antiviral agents designed to target particular viral proteins such as influenza neuraminidase are being developed. However, the ability of viruses to mutate the target proteins represents an obstacle for effective treatment with molecules which selectively inhibit the function of specific viral polypeptides. Thus, there is need for new therapeutic strategies to prevent and treat viral infections.

Additionally, there is a need for new therapies for the prevention and treatment of bacterial infections, especially bacterial infections caused by multiple drug resistant bacteria. Currently, bacterial infections are treated with various antibiotics. Although antibiotics have and can be effective in the treatment of various bacterial infections, there are a number of limitations to the effectiveness and safety of antibiotics. For example, some individuals have an allergic reaction to certain antibiotics and other individuals suffer from serious side effects. Moreover, continued use of antibiotics for the treatment of bacterial infections contributes to formation of antibiotic-resistant strains of bacteria.

Examples of infectious diseases are AIDS, alveolar hydatid disease (AHD, echinococcosis), amebiasis (Entamoeba histolytica infection), Angiostrongylus infection, anisakiasis, anthrax, babesiosis (Babesia infection), Balantidium infection (balantidiasis), Baylisascaris infection (racecoon roundworm), bilharzia (schistosomiasis), Blastocystis hominis infection (blastomycosis), borreliosis, botulism, Brainerd diarrhea, brucellosis, bovine spongiform encephalopathy (BSE), candidiasis, capillaritis (Capillaria infection), chronic fatigue syndrome (CFS), Chagas disease (American trypanosomiasis), chickenpox (Varicella-Zoster virus), Chlamydia pneumoniae infection, cholera, Creutzfeldt-Jakob disease (CJD), clonorchiasis (Clonorchis infection), cutaneous larva migrans (CLM) (hookworm infection), coccidioidomycosis, conjunctivitis, Coxackievirus A16 (hand, foot and mouth disease), cryptococcosis, Cryptosporidium infection (cryptosporidiosis), Culex mosquito (West Nile virus vector), cyclosporiasis (Cyclospora infection), cryptosporidiosis (neurocysticercosis), Cytomegalovirus infection, Dengue/Dengue fever, Diethylidium infection (dog and cat fleen tapeworm), Ebola virus hemorrhagic fever, encephalitis, Entamoeba coli infection, Entamoeba dispar infection, Entamoeba hartmanni infection, Entamoeba histolytica infection (amebiasis), Entamoeba polecki infection, enterobiasis (pinworm infection), enterovirus infection (non-polio), Epstein-Barr virus infection, Escherichia coli infection, foodborne infection, foot and mouth disease, fungal dermatitis, gastroenteritis, group A streptococcal disease, group B streptococcal disease, Hansen’s disease (leprosy), Hantavirus pulmonary syndrome, head lice infestation (pediculosis), Helicobacter pylori infection, hematologic disease, Hendra virus infection, hepatitis (HCV, HBV), herpes zoster (shingles), HIV Infection, human ehrlichiosis, human parainfluenza virus infection, influenza, isosporiasis (Isospora infection), Lassa fever, leishmaniasis, Kala-azar (Kala-azar, Leishmania infection), lice (body lice, head lice, pubic lice), Lyme disease, malaria, Marburg hemorrhagic fever, measles, meningitis, mosquito-borne diseases, Mycobacterium avium complex (MAC) infection, Naegleria infection, nosocomial infections, nonpathogenic intestinal ameboid infection, onchocerciasis (river blindness), opisthorchiasis (Opisthorchis infection), parvovirus infection, plague, Pneumocystis carinii pneumonia (PCP), polio, Q fever, rabies, respiratory syncytial virus (RSV) Infection, rheumatic fever, Rift Valley fever, river blindness (onchocerciasis), rotavirus infection, roundworm infection, salmonellosis, salmonella enteritidis, scabies, shigellosis, shingles, sleeping sickness, smallpox, streptococcal Infection, tapeworm infection (Taenia infection), tetanus, toxic shock syndrome, tuberculosis, ulcers (peptic ulcer disease), valley fever, Vibrio parahaemolyticus infection, Vibrio vulnificus infection, viral hemorrhagic fever, warts, waterborne infectious diseases, West Nile virus infection (West Nile encephalitis), whooping cough, yellow fever.

Another aspect of the present invention is directed to the use of the peptide for prophylaxis and/or treatment of prion diseases.

Prions are infectious agents which do not have a nucleic acid genome. It seems that a protein alone is the infectious agent. A prion has been defined as “small proteinaceous infectious particle which resists inactivation by procedures that modify nucleic acids”. The discovery that proteins alone can transmit an infectious disease came as a considerable surprise to the scientific community. Prion diseases are often called “transmissible spongiform encephalopathies”, because of the post mortem appearance of the brain with large vacuoles in the cortex and cerebellum. Probably most mammalian species develop these diseases. Prion diseases are a group of neurodegenerative disorders of humans and animals and the prion diseases can manifest as sporadic, genetic or infectious disorders. Examples of prion diseases acquired by exogenous infection are bovine spongiform encephalitis (BSE) of cattle and the new variant of Creutzfeldt-Jakob disease (vCJD) caused by BSE as well as scrapie of animals. Examples of human prion diseases include kuru, sporadic Creutzfeldt-Jakob disease (sCJD), familial CJD (fCJD), iatrogenic CJD (iCJD), Gerstmann-Sträussler-Scheinker (GSS) disease, fatal familial insomnia (FFI), and especially the new variant CJD (mCJD or vCJD).

The name “prion” is used to describe the causative agents which underlie the transmissible spongiform encephalopathies. A prion is proposed to be a novel infectious particle that differs from viruses and viroids. It is composed solely of one unique protein that resists most inactivation procedures such as heat, radiation, and proteases. The latter characteristic has led to the term prion protein. The prion protein involves a protease-resistant isofrom of the prion protein. The protease-resistant isofrom has been proposed to slowly catalyze the conversion of the normal prion protein into the abnormal form.
The term “isoform” in the context of prions means two proteins with exactly the same amino acid sequence that can fold into molecules with dramatically different tertiary structures. The normal cellular isoform of the prion protein (PrP\textsuperscript{c}) has a high α-helix content, a low β-sheets content, and is sensitive to protease digestion. The abnormal, disease-causing isoform (PrP\textsuperscript{sc}) has a lower α-helix content, a much higher β-sheet content, and is much more resistant to protease digestion.

As used herein the term “prion diseases” refers to transmissible spongiform encephalopathies. Examples for prion diseases comprise scrapie (sheep, goat), transmissible mink encephalopathy (TME; mink), chronic wasting disease (CWD; muledeer, deer, elk), bovine spongiform encephalopathy (BSE; cows, cattle), Creutzfeldt-Jacob Disease (CJD), variant CJD (vCJD), sporadic CJD-Jakob disease (sCJD), familial CJD (fCJD), iatrogenic CJD (iCJD), Gerstmann-Sträussler-Scheinker syndrome (GSS), fatal familial insomnia (FFI), and kuru. Preferred are BSE, vCJD, and CJD.

**Streptococcus pneumoniae** Infection

*Streptococcus pneumoniae*, also known as pneumococcus, is an alpha-hemolytic diplococcus bacterium and a member of the genus *Streptococcus* that is often found in the noses and throats of healthy persons and is spread person-to-person through close contact. Pneumococcus is a common cause of mild illnesses, such as sinus and ear infections, but also causes life-threatening infectious such as pneumonia, meningitis, infections of the bloodstream, acute sinusitis, otitis media, osteomyelitis, septic arthritis, endocarditis, peritonitis, pericarditis, and brain abscess. Many strains are resistant to antibiotics. *Streptococcus pneumoniae* is the most common cause of bacterial meningitis in adults and children, and is one of the top two isolates found in otitis media. Pneumococcal pneumonia is more common in the very young and the very old.

Pneumococcal disease occurs worldwide. Pneumococcal disease is more common in winter months and when respiratory viruses such as influenza are circulating. Outbreaks of pneumococcal disease are not uncommon but can occur in child care centers, nursing homes, or other institutions. In the United States, most deaths from pneumococcal disease occur in older adults, although in developing countries many children die of pneumococcal pneumonia.

Risk for pneumococcal disease is highest in young children, the elderly, and persons of any age who have chronic medical conditions such as heart disease, lung disease, or diabetes, or conditions that suppress the immune system, such as HIV. Smokers and those in close contact with small children are at higher risk. Crowded settings or situations with close, prolonged contact with young children may increase the risk of contracting pneumococcal disease while traveling. Fever and malaise are typical symptoms for all forms of pneumococcal disease and may be the only symptoms in young children with blood infections. Patients with pneumonia usually have cough, often with purulent or blood-tinted sputum, and may have shaking chills, shortness of breath, or pleuritic chest pain. Fever and sputum production may be absent in elderly persons with pneumococcal pneumonia. Pneumococcal meningitis, ear infections or sinus infections resemble these conditions caused by other bacteria.

**Virulence Factors**

*Streptococcus pneumoniae* expresses different virulence factors on its cell surface and inside the organism. These virulence factors contribute to some of the clinical manifestations during infection with *Streptococcus pneumoniae*:

- **Polysaccharide capsule**—prevents phagocytosis by host immune cells by inhibiting C3b opsonization of the bacterial cells
- **Pneumolysin (Ply)**—a 53-kDa protein that can cause lysis of host cells and activate complement
- **Autolysin (LytA)**—activation of this protein lyses the bacteria releasing its internal contents (i.e., pneumolysin)
- **Hydrogen peroxide**—causes damage to host cells (can cause apoptosis in neuronal cells during meningitis) and has bactericidal effects against competing bacteria (*Haeomophilus influenzae*, *Neisseria meningitidis*, *Staphylococcus aureus*).
- **Pili**—hair-like structures that extend from the surface of many strains of *Streptococcus pneumoniae*. They contribute to colonization of upper respiratory tract and increase the formation of large amounts of TNF by the immune system during sepsis, raising the possibility of septic shock.
- **Choline binding protein A/Pneumococcal surface protein A (CbpA/PspA)**—an adhesin that can interact with carbohydrates on the cell surface of pulmonary epithelial cells and can inhibit complement-mediated opsonization of pneumococci

**Treatment Options**

Historically, treatment relied primarily on β-lactam antibiotics but there has been an increasing prevalence of penicillin resistance, especially in areas of high antibiotic use. A varying proportion of strains may also be resistant to cephalosporins, macrolides (such as erythromycin), tetracycline, clindamycin and the quinolones. Penicillin-resistant strains are more likely to be resistant to other antibiotics. Most isolates remain susceptible to vancomycin, though its use in a β-lactam-susceptible isolate is less desirable because of tissue distribution of the drug and concerns of development of vancomycin resistance, and therefore there is an unmet medical need for developing new types of treatment options.

**Mycobacterium tuberculosis** Infection

Tuberculosis (TB) is an often severe and contagious airborne disease caused by infection with *Mycobacterium tuberculosis* (Mtbo). TB typically affects the lungs but it also may affect any other organ of the body. It is usually treated with a regimen of drugs taken for six months to two years depending on the type of infection. TB infection begins when the mycobacteria reach the pulmonary alveoli, where they invade and replicate within alveolar macrophages. Bacteria are picked up by dendritic cells, which do not allow replication, although these cells can transport the bacilli to local lymph nodes. Further spread is through the bloodstream to the more distant tissues and organs where secondary TB lesions form in lung apices, peripheral lymph nodes, kidneys, brain, and bone.

Tuberculosis is classified as one of the granulomatous inflammatory conditions. Macrophages, T lymphocytes, B lymphocytes and fibroblasts are among the cells that aggregate to form a granuloma, with lymphocytes surrounding the infected macrophages. The granuloma functions not only to prevent dissemination of the mycobacteria, but also provides a local environment for communication of cells of the immune system. Within the granuloma, T lymphocytes (CD4+) secrete cytokines such as interferon gamma, which
activates macrophages to destroy the bacteria with which they are infected. T lymphocytes (CD8+) can also directly kill infected cells. Importantly, bacteria are not always eliminated within the granuloma, but can become dormant, resulting in a latent infection. Another feature of the granulomas of human tuberculosis is the development of cell death, also called necrosis, in the center of tubercles.

If TB bacteria gain entry to the bloodstream from an area of damaged tissue they spread through the body and set up many foci of infection, all appearing as tiny white tubercles in the tissues. This severe form of TB disease is most common in infants and the elderly and is called miliary tuberculosis. Patients with this disseminated TB have a fatality rate of approximately 20%, even with intensive treatment.

In many patients the infection waxes and wanes. Tissue destruction and necrosis are balanced by healing and fibrosis. Affected tissue is replaced by scarring and cavities filled with cheese-like white necrotic material.

Difference Between MtB Infection and TB Disease

In understanding tuberculosis, it’s important to understand the difference between MtB infection and TB disease.

MtB Infection

It is estimated that more than one-third of the world’s population has the TB bacterium in their bodies, which means they have MtB infection. In addition, new infections are occurring at the rate of one per second. Fortunately, only a fraction of people infected with MtB develops active TB disease. Those who do not get sick are known to have non-contagious latent TB, so-called because the bacteria are inactive or “asleep” in the body.

TB bacteria can remain in this dormant state for months, years, and even decades without increasing in number and without making the person sick. Most people with latent MtB infection will test positive on the tuberculin skin test, or their chest X-ray will show signs of latent TB. These findings indicate that they have the TB germ in their bodies, but most infected people will not develop active TB disease, may never get sick, may never show any symptoms, and may never spread the bacteria to others. However, one in ten people infected with TB bacteria, do develop active TB disease.

TB Disease

People with weakened immune systems (individuals with HIV disease, those receiving chemotherapy, or children under five years of age, for example) are at a greater risk for developing TB disease. When they breathe in TB bacteria, the bacteria settle in the lungs and start growing because the individual’s immune system cannot fight the bacteria. In these people, TB disease may develop within days or weeks after the infection. However, in some other people, TB disease may develop months or years after the initial infection, at a time when the immune system becomes weak for other reasons and is no longer able to fight the bacteria. When a person gets active TB, it means the TB bacteria are multiplying and attacking the lung(s), or other parts of the body such as the lymph nodes, bones, kidney, brain, spine, and even the skin. From the lungs, the TB bacteria move through the blood to different parts of the body. Symptoms of active disease include cough, loss of weight and appetite, fever, chills and night sweats as well as symptoms from the specific organ or system that is affected; for example, coughing up blood or pus and in TB of the lungs, or bone pain if the bacteria have invaded the bones. TB disease usually can be cured with prompt and appropriate treatment, but it remains a major cause of death and disability in the world, particularly among persons infected with human immunodeficiency virus (HIV). TB Bacteria are Spread Only from a Person with Active TB Disease

In people who develop active TB of the lungs, the TB skin test will often be positive. In addition, these people will show all the signs and symptoms of TB disease, and can pass the bacteria to others. Statistics show that approximately one-third of the people exposed to pulmonary TB become infected with the bacteria, but only one in ten of these infected people develop active TB disease during their lifetimes.

Among people suffering from TB disease, three out of four have disease affecting the lungs. If not treated immediately, the bacteria have the potential to destroy the lungs and kill the person.

Tuberculosis, which results from an infection with Mycobacterium tuberculosis, can usually be cured with a combination of first-line drugs taken for several months. Shown here are the four drugs in the standard regimen of first-line drugs and their modes of action. Also shown are the dates these four drugs were discovered all more than 40 years ago.

Multidrug-Resistant Tuberculosis (MDR TB)

MDR TB is a form of drug-resistant TB in which the TB bacteria can no longer be killed by at least the two best antibiotics, isoniazid (INH) and rifampin (RIF), commonly used to cure TB. As a result, this form of the disease is more difficult to treat than ordinary TB and requires up to two years of multidrug treatment.

People may get MDR TB in two ways:

1. Directly, if they spend time with an MDR TB patient and breathe in the MDR TB bacteria, and
2. If they already have active TB and do not properly follow their prescribed treatment regimen or TB medication is not reliably available to them. This inconsistent use of TB antibiotics gives the bacteria enough time to evolve and evade the first-line anti-TB medications, and regular TB may then progress to MDR TB, which is more challenging to treat.

Extensively Drug-Resistant Tuberculosis (XDR TB)

XDR TB is a less common form of multidrug-resistant TB in which the TB bacteria have changed enough to circumvent the two best antibiotics, isoniazid (INH) and rifampin (RIF), as well as most of the alternative drugs used against MDR TB. These second-line drugs include any fluoroquinolone, and at least one of the other three injectable anti-TB drugs: amikacin, kanamycin, or capreomycin. As a
result, this form of the disease needs up to two years of extensive drug treatment and is the most challenging to treat.

[0050] People may get XDR TB in two ways:

[0051] directly, if they spend time with an XDR TB patient and breathe in the XDR TB bacteria, and

[0052] if they already have MDR TB or active TB, and do not properly follow their prescribed treatment regimen or TB medication is not reliably available to them. This inconsistent use of TB antibiotics gives the bacteria enough time to evolve and evade most if not all TB drugs, making it extremely difficult or impossible to treat XDR TB.

[0053] XDR TB occurs when a *Mycobacterium tuberculosis* strain is resistant to isoniazid and rifampin, two of the most powerful first-line drugs, as well as key drugs of the second line regimen—any fluoroquinolone and at least one of the three injectable drugs shown above. XDR TB strains may also be resistant to additional drugs, greatly complicating therapy. In the European Community countries, comprising 495 million people, there are estimated 1 300 patients suffering from MDR Tuberculosis. In the USA, comprising 302 million people, there are estimated 200 patients suffering from MDR Tuberculosis.

### TABLE 1

<table>
<thead>
<tr>
<th>PROFILE OF THE COUNTRY AND ITS CONTROL PROGRAMME</th>
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<td>Country examples</td>
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<td>Countrywide: 2001</td>
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Population in year of survey | 284,796,887 | Year N.T.P was established | 1953 |
Notification all cases (rate) | 6/100,000 | Year of Rifampicin introduction | 1971 |
Estimated incidence (all cases) | 5.6/100,000 | Year of Isoniazid introduction | 1952 |
Notification new sputum smear + | 560 | Use of Standardized Regimens | Yes |
Notification new sputum smear + (rate) | 2/100,000 | % Use of Short Course Chemotherapy | Yes | 90% |
Treatment Success | 76% | Use of Directly Observed Therapy | Yes | 49% |
Retreatment cases | NA | During continuation phase | Yes |
Retreatment as % of NTP | NA | % Use of Fixed Dose Combination | Yes |
Estimated HIV positive TB cases | 9.0% | Treatment in private sector | Cat 3 |

**CHARACTERISTICS OF THE SURVEY/SURVEILLANCE PROGRAMME**

| Study Duration | 12 Months |
| Target Area | Countrywide |
| Sampling Method | All cases |
| Culture Media | Various |
| DST Method | Various |
| Supranational Reference Laboratory | Centers for Disease Control and Prevention (CDC), Atlanta, United States of America |

### TABLE 2

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TABLE 3

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<th>NUMBER OF RESISTANT STRAIN ANALYSED AT THE NRC FROM 1993-2004</th>
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<tr>
<td>year</td>
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<tr>
<td></td>
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<td></td>
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<td>(a)</td>
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(a) includes 49 SM resistant strains;
(b) includes 69 SM monoresistant strains

[0054] Currently, the treatment options are overall unsatisfactory and the disease is a ticking bomb.

[0055] The peptide of the present invention was tested using the assays described in Examples 1-7 for their effect as active therapeutic agents in the prophylaxis and/or treatment of infections diseases and disorders.

Autoimmune Disease

[0056] Autoimmune disease refers to any of a group of diseases or disorders in which tissue injury is associated with a humoral and/or cell-mediated immune response to body constituents or, in a broader sense, an immune response to self. The pathological immune response may be systemic or organ specific. That is, for example, the immune response directed to self may affect joints, skin, myelin sheath that protects neurons, kidney, liver, pancreas, thyroid, adrenals, and ovaries.

[0057] In fact, the list of autoimmune diseases is composed of more than eighty disorders. A few autoimmune diseases such as vitiligo, in which patches of skin lose pigmentation, are merely annoying. Most others are debilitating, often progressive with time and eventually fatal. Systemic lupus erythematosus (SLE), for example, is a chronic disease in which 10-15% of patients die within a decade of diagnosis, in all but a few autoimmune diseases, the sex ratio skew towards women. For example, in SLE the ratio of female to male patients is nine to one. In one particular case, Hashimoto’s disease in which the immune system attacks the thyroid gland, the ratio is fifty to one.

[0058] It has long been known that immune complex formation plays a role in the etiology and progression of autoimmune disease. For example, inflammation in patients with arthritis has long been considered to involve phagocytosis by leukocytes of complexes of antigen, antibody and complement-immune complexes. However, only now it is being recognized that inflammation caused by immune complexes in the joints (arthritis), the kidneys (glomerulonephritis), and blood vessels (vasculitis) is a major cause of morbidity in autoimmune diseases. Increased immune complex formation correlates with the presence of antibodies directed to self or so-called autoantibodies, and the presence of the latter can also contribute to tissue inflammation either as part of an immune complex or unbound to antigen (free antibody). In some autoimmune diseases, the presence of free autoantibody contributes significantly to disease pathology. This has been clearly demonstrated for example in SLE (anti-DNA antibodies), immune thrombocytopenia (antibody response directed to platelets), and to a lesser extent rheumatoid arthritis (IgG reactive rheumatoid factor). The important role of immune complexes and free autoantibodies is further demonstrated by the fact that successful treatment of certain autoimmune diseases has been achieved by the removal of immune complexes and free antibody by means of specific immunoabsorption procedures. For example, the use of an apheresis procedure in which immune complexes and antibodies are removed by passage of a patient’s blood through an immunoaffinity column was approved by the U.S. FDA in 1987 for immune thrombocytopenia (ITP) and in 1999 for rheumatoid arthritis. However, currently there is no approved method for the treatment of autoimmune diseases which facilitates the elimination of immune complexes and autoantibodies by administration of a drug.

[0059] Another aspect of the etiology and progression of autoimmune disease is the role of proinflammatory cytokines. Under normal circumstances, proinflammatory cytokines
such as tumor necrosis factor $\alpha$ (TNF$\alpha$) and interleukin-1 (IL-1) play a protective role in the response to infection and cellular stress. However, the pathological consequences which result from chronic and/or excessive production of TNF$\alpha$ and IL-1 are believed to underlie the progression of many autoimmune diseases such as rheumatoid arthritis, Crohn’s disease, inflammatory bowel disease, and psoriasis. Other proinflammatory cytokines include interleukin-6, interleukin-8, interleukin-17, and granulocyte-macrophage colony stimulating factor.

Naturally occurring CD4+CD25+ regulatory T cells (Tregs) play a critical role in the control of periphery tolerance to self-antigens. Interestingly, they also control immune responses to allergens and transplant antigens. Recent studies in animal models have shown that adoptive transfer of CD4+ CD25+ Tregs can prevent or even cure allergic and autoimmune diseases, and appear to induce transplantation tolerance. Thus, adoptive cell therapy using patient-specific CD4+ CD25+ Tregs has emerged as an individualized medicine for the treatment of inflammatory disease including allergy, autoimmune disease and transplant rejection. Furthermore, strategies to activate and expand antigen-specific CD4+ CD25+ Tregs in vivo using pharmacological agents may represent a novel avenue for drug development.

The interaction of leukocytes with the vessel endothelium to facilitate the extravasation into the tissue represents a key process of the body’s defense mechanisms. Excessive recruitment of leukocytes into the inflamed tissue in chronic diseases like autoimmune disorders could be prevented by interfering with the mechanisms of leukocyte extravasation. Significant progress in elucidating the molecular basis of the trafficking of leukocytes from the blood stream to the extravascular tissue has been achieved that enables new strategies for therapeutic approaches. The multistep process of leukocyte rolling, firm adhesion and transmigration through the endothelial wall is facilitated by a dynamic interplay of adhesion receptors on both leukocytes and on endothelial cells as well as chemokines. In preclinical studies using various animal models, promising results have been obtained demonstrating that blocking of adhesion receptors of the selectin and integrin families improved the inflammation process in models of ulcerative colitis, autoimmune encephalomyelitis or contact hypersensitivity. In addition to the targeting of adhesion receptors by antibodies, small molecules that mimic epitopes of adhesion receptor ligands have been developed and successfully applied in animal models. Clinical studies revealed a limited response using antibodies to selectins or leukocyte function-associated antigen 1 (LFA-1) integrins compared with animal models. However, using humanized antibodies to the alpha 4-integrin subunit significant efficacy has been demonstrated in autoimmune diseases like psoriasis, multiple sclerosis and inflammatory bowel disease.

Examples of autoimmune diseases of the eyes are idiopathic opticus-neuritis, ophthalmia sympathica, anterior uveitis and other uveitis forms, retina degeneration, and Mooren’s ulcer.

Examples of autoimmune diseases of the skin are bullous pemphigoides, chronic urticaria (autoimmune subtype), dermatitis herpetiformis (morbus Duhring), epidermolysis bullosa aquisita (EBA), acquired angioedema, herpes gestations, hypercomplementemic urticarial vasculitis syndrome (HUVS), linear IgA-dermatosis, and pemphigus.

Examples of hematological autoimmune diseases are autoimmune hemolytic anemia, autoimmune neutropenia, Evans syndrome, inhibitor hemophilia, idiopathic thrombocytopenic purpura (ITP) and pernicious anemia.

Examples of gynecological autoimmune diseases are habitual abortion and infertility.

Examples of autoimmune diseases of the heart are congenital heart block, idiopathic dilatative cardiomyopathy, peripartum-cardiomyopathy, postcardiotomy syndrome, and postinfarct syndrome (Dressler syndrome).

Examples of autoimmune diseases of the ear, nose and throat are chronic sensorineural hearing loss and morbus Menière.

Examples of autoimmune diseases of the colon are autoimmune enteropathy, colitis ulcerosa, indeterminant colitis, Crohn’s disease and gluten-sensitive enteropathy.

Examples of autoimmune endocrinological autoimmune disorders are autoimmune polyglandular syndrome type 1, autoimmune polyglandular syndrome type 2, diabetes mellitus type 1 (IDDM), Hashimoto-thyroiditis, insulin autoimmune-syndrome (IAS), idiopathic diabetes insipidus, idiopathic hyperparathyroidism, idiopathic Addison’s disease and Graves-Basedow disease.

Examples of autoimmune diseases of the liver are autoimmune hepatitis (AIH type 1, 2 and 3), primary biliary cirrhosis (PBC), and primary sclerosing cholangitis.

Example of autoimmune diseases of the lung is Goodpasture’s syndrome.

An example of an autoimmune disease of the stomach is chronic atrophic (type A) gastritis.

Examples of neurological autoimmune disorders are Guillain-Barré syndrome, IgM gammopathy-associated neuropathy, Lambert-Eaton syndrome, Miller-Fisher syndrome, multiple sclerosis, multifocal motoric neuropathy, myasthenia gravis, paraneoplastic neurological syndrome, Rasmussen’s encephalitis, and stiff-man syndrome.

Examples of autoimmune diseases of the kidney are anti-TBM-nephritis, Goodpasture’s syndrome/anti-GBM-nephritis, IgA-nephropathy, interstitial nephritis, and membrane proliferative glomerulonephritides.

Further diseases that may be caused by an autoimmune reaction are Behcet disease, chronic fatigue immune dysfunction syndrome (CFIDS), Cogan syndrome I, endometriosis, HELLP syndrome, Bechterew’s disease, polymyalgia rheumatica, psoriasis, sarcoidosis and vitiligo.

During the last decade, new biotherapies have been developed for the treatment of systemic autoimmune diseases. The targets of these new treatments are all the steps of the immune response. These new therapies are: B lymphocyte (BL) inhibitors such as anti-CD20 monoclonal antibody, B lymphocyte stimulator (BLyS) antagonists and tolerogens of pathogenic-antibody secreting cells; B inhibitors of the costimulation between antigen-presenting cells and T lymphocyte (TL) like monoclonal anti-CD40 ligand antibody or CTLA4-Ig (abatecept); TL antagonists which can inhibit the proliferation of autoreactive T cells; cytokine antagonists; chemokine and adhesion antagonists which inhibit trafficking of immunocompetent cells to target organs. These new approaches are based on a better understanding of the autoimmune response.

The peptide of the present invention was tested using the assays described in Examples 14-15 for their effect
as active therapeutic agents in the prophylaxis and/or treatment of autoimmune diseases and disorders.

Fibrotic Disease

[0078] Fibrosis or fibrotic associated disorder affects the liver, epidermis, endodermis, muscle, tendon, cartilage, heart, pancreas, lung, uterus, nervous system, testis, ovary, adrenal gland, artery, vein, colon, small intestine, biliary tract, or stomach. In a further embodiment, the fibrosis or fibrosis associated disorder is interstitial lung fibrosis. In another embodiment the fibrosis or fibrosis associated disorder is the result of an infection with schistosoma. In another embodiment the fibrosis or fibrosis associated disorder is the result of wound healing.

[0079] Fibrosis is generally characterized by the pathologic or excessive accumulation of collagenous connective tissue. Fibrotic diseases and disorders include, but are not limited to, collagen disease, interstitial lung disease, human fibrotic lung disease (e.g., obliterative bronchiolitis), idiopathic pulmonary fibrosis, pulmonary fibrosis from a known etiology, tumor stroma in lung disease, systemic sclerosis affecting the lungs, Hamansky-Pudlak syndrome, coal worker’s pneumoconiosis, asbestosis, silicosis, chronic pulmonary hypertension, AIDS-associated pulmonary hypertension, sarcoidosis, and the like), fibrotic vascular disease, tubulointerstitial and glomerular fibrosis, myocardial fibrosis, arterial sclerosis, atherosclerosis, varicose veins, coronary infarcts, cerebral infarcts, myocardial fibrosis, musculoskeletal fibrosis, postsurgical adhesions, human kidney disease (e.g., nephritic syndrome), Alport’s syndrome, HIV-associated nephropathy, polycystic kidney disease, Fabry’s disease, diabetic nephropathy, chronic glomerulonephritis, nephritis associated with systemic lupus, and the like), cutis keloid formation, progressive systemic sclerosis (PSS), primary scarring cholangitis (PSC), liver fibrosis, liver cirrhosis, renal fibrosis, pulmonary fibrosis, cystic fibrosis, chronic graft versus host disease, scleroderma (local and systemic), Grave’s ophthalmopathy, diabetic retinopathy, glaucoma, Peyronie’s disease, penis fibrosis, urethrostenois after a test using a cystoscope, inner accretion after surgery, scarring, myofibrosis, idiopathic retroperitoneal fibrosis, peritoneal fibrosis from a known etiology, drug induced ergotism, fibrosis incident to benign or malignant cancer, fibrosis incident to microbial infection (e.g., viral, bacterial, parasitic, fungal, etc.), Alzheimer’s disease, fibrosis incident to inflammatory bowel disease (including stricture formation in Crohn’s disease and microscopic colitis), fibrosis induced by chemical or environmental insult (e.g., cancer chemotherapy, pesticides, radiation/cancer radiotherapy), and the like.

[0080] Diseases associated with fibrosis include lupus, graft versus host disease, scleroderma, systemic sclerosis, scleroderma-like disorders, sine scleroderma, calcinosis, Raynaud’s esophageal dysfunction, sclerodactyly, telangiectasia, hypersensitivity pneumonitis, collagen vascular disease, asthma, pulmonary arterial hypertension, glomerulonephritis, chronic obstructive pulmonary disease, fibrosis following myocardial infarction, central nervous system fibrosis following a stroke or neuro-degenerative diseases (e.g. Alzheimer’s disease), proliferative vitreoretinopathy (PVR) and arthritis, silicosis, asbestosis induced pulmonary fibrosis, acute lung injury and acute respiratory distress syndrome (including bacterial pneumonia induced, trauma induced, viral pneumonia induced, tuberculosis, ventilator induced, non-pulmonary sepsis induced, and aspiration induced).

Increased Number of Activated Myofibroblasts in Fibrotic Diseases

[0081] The emergence and disappearance of the myofibroblast appears to correlate with the initiation of active fibrosis and its resolution, respectively. In addition, the myofibroblast has many phenotypic features, which embody much of the pathologic alterations in fibrotic tissue, e.g., lung tissue. These features would seem to argue for an important role for the myofibroblast in the pathogenesis of fibrosis, e.g., lung fibrosis. Furthermore, the persistence of the myofibroblast may herald progressive disease, and, conversely, its disappearance may be an indicator of resolution. This in turn suggests that future therapeutic strategies targeting the myofibroblast would be productive.

[0082] Patients usually exhibit evidence of active fibrosis with increased numbers of activated fibroblasts, many of which have the phenotypic characteristics of myofibroblasts. At these sites, increased amounts of extracellular matrix deposition are evident with efficiency of the normal alveolar architecture. Animal model studies show the myofibroblast to be the primary source of type I collagen gene expression in active fibrotic sites. In vitro studies show differentiation of these cells from fibroblasts under the influence of certain cytokines but indicate their susceptibility to nitric oxide mediated apoptosis. In addition to promoting myofibroblast differentiation, transforming growth factor-β1 (TGF-β1) provides protection against apoptosis. Thus, this well-known fibrogenic cytokine is important both for the emergence of the myofibroblast and its survival against apoptotic stimuli. This is consistent with the critical importance of this cytokine in diverse models of fibrosis in various tissues. In view of these properties, the persistence or prolonged survival of the myofibroblast may be the key to understanding why certain forms of lung injury may result in progressive disease, terminating in end stage disease.

[0083] Although pulmonary fibrosis has diverse etiologies, there is a common feature characteristic of this process, namely, the abnormal deposition of extracellular matrix that efficaces the normal lung tissue architecture. A key cellular source of this matrix is the mesenchymal cell population that occupies much of the fibrotic lesion during the active period of fibrosis. This population is heterogeneous with respect to a number of key phenotypes. One of these phenotypes is the myofibroblast, which is commonly identified by its expression in α-smooth muscle actin and by features that are intermediate between the bona fide smooth muscle cell and the fibroblast. The de novo appearance of myofibroblasts at sites of wound healing and tissue repair/fibrosis is associated with the period of active fibrosis and is considered to be involved in wound contraction. Furthermore, the localization of myofibroblasts at sites undergoing active extracellular matrix deposition suggests an important role for these cells in the genesis of the fibrotic lesion.

Increased TGF-β, Family Levels in Fibrotic Diseases

[0084] The transforming growth factor-β1 (TGF-β1) family of proteins has the most potent stimulatory effect on extracellular matrix deposition of any cytokines so far examined. In animal models of pulmonary fibrosis enhanced TGF-β1
gene expression is temporally and spatially related to increased collagen gene expression and protein deposition. TGF-β1 antibodies reduce collagen deposition in murine bleomycin-induced lung fibrosis and human fibrotic lung tissue shows enhanced TGF-β1 gene and protein expression. Several lines of evidence suggest that TGF-β is a central regulator of pulmonary fibrosis. Several animal models over expressing TGF-β showed extensive progressive fibrosis but limited inflammation, indicating that TGF-β may play a predominant role in the progression of pulmonary fibrosis. Therapeutic efforts are therefore focusing on inhibition of TGF-β activity, for instance by anti-TGF-β1-antibodies, or modulators of TGF-β1 such as pirfenidone.

Pirfenidone inhibits TGF-β1 gene expression in vivo resulting in inhibition of TGF-β1-mediated collagen synthesis and appears to slow progression of IPF in patients. Other novel, promising antifibrotic agents include relaxin (inhibits TGF-β-mediated overexpression of collagen and increases collagenases), suramin (inhibits growth factors), prostaglandin E2 (inhibits collagen production) and lovastatin (blocks formation of granulation tissue by induction of fibroblast apoptosis).

Diseases involving the lung associated with increased levels of TGF-β include chronic lung disease of prematurity, idiopathic pulmonary fibrosis, rapid progressive pulmonary fibrosis, giant-cell interstitial pneumonia, acute rejection after lung transplantation, cytomegalovirus pneumonitis after lung transplantation, bronchiolitis obliterans, asbestosis, coal worker’s pneumoconiosis, silicosis, histiocytosis, sarcoidosis, eosinophilic granuloma, scleroderma, systemic lupus erythematosus, lymphangioleiomyomatosis, central fibrosis in pulmonary adenocarcinoma, cystic fibrosis, chronic obstructive lung disease, and asthma.

Increased TNF-α Levels in Fibrotic Diseases

An important role of tumor necrosis factor-α (TNF-α) in interstitial fibrosis has been established using transgenic mice, which either overexpress or display a deficiency of this cytokine. Mice transgenically modified to overexpress TNF-α develop lung fibrosis. In contrast, mice null for TNF-α show marked resistance to bleomycin induced fibrosis. TNF-α can stimulate fibroblast replication and collagen synthesis in vitro, and pulmonary TNF-α gene expression rises after administration of bleomycin in mice. Soluble TNF-α receptors reduce lung fibrosis in murine models and pulmonary overexpression of TNF-α in transgenic mice is characterized by lung fibrosis. In patients with CIA or asbestosis, bronchoalveolar lavage fluid-derived macrophages release increased amounts of TNF-α compared with controls.

Increased TNF-α may induce fibrosis or fibrosis-associated conditions affecting any tissue including, for example, fibrosis of an internal organ, a cutaneous or dermal fibroin disorder, and fibrotic conditions of the eye. Fibrosis of internal organs (e.g., liver, lung, kidney, heart blood vessels, gastrointestinal tract) occurs in disorders such as pulmonary fibrosis, idiopathic fibrosis, autoimmune fibrosis, myelofibrosis, liver cirrhosis, veno-occlusive disease, mesangial proliferative glomerulonephritis, crescentic glomerulonephritis, diabetic nephropathy, renal interstitial fibrosis, renal fibrosis in subjects receiving cyclosporin, allograft rejection, HTV associated nephropathy. Other fibrosis-associated disorders include systemic sclerosis, eosinophilia-myalgia syndrome, and fibrosis-associated CNS disorders such as intraocular fibrosis. Dermal fibrosing disorders include, for example, scleroderma, morphea, keloids, hypertrophic scars, familial cutaneous collagenoma, and connective tissue nevi of the collagen type. Fibrotic conditions of the eye include conditions such as diabetic retinopathy, post-surgical scarring (for example, after glaucoma filtering surgery and after crossed-eyes (strabismus) surgery), and proliferative vitreo-retinopathy. Additional fibrotic conditions that may be treated by the methods of the present invention may result, for example, from rheumatoid arthritis, diseases associated with prolonged joint pain and deteriorated joints; progressive systemic sclerosis, polymyositis, dermatomyositis, eosinophilic fascitis, morphea, Raynaud’s syndrome, and nasal polyposis.

Increased Matrix Metalloproteases Levels in Fibrotic Diseases

The abnormal extracellular matrix (ECM) remodeling observed in the lungs of patients with interstitial pulmonary fibrosis (IPF) is due, at least in part, to an imbalance between matrix metalloproteases (MMPs) and tissue inhibitor of metalloproteinases (TIMPs). Normal lung fibroblasts do not make MMP-9 in vitro, whereas fibroblasts from IPF lungs strongly express MMP-9. In addition, fibroblasts from patients with IPF express increased levels of all TIMPs. In this setting, TIMPs may play a role in apoptosis in some cell populations. In vitro studies of alveolar macrophages obtained from untreated patients with idiopathic pulmonary fibrosis showed marked increase in MMP-9 secretion compared to macrophages collected from healthy individuals. In animals models of bleomycin-induced pulmonary fibrosis MMPs have been shown to be elevated in bronchoalveolar lavage (BAL) fluid. Indeed, a synthetic inhibitor of MMP, Batimastat, has been shown to significantly reduce bleomycin-induced lung fibrosis, again pointing to the importance of MMPs in the development of this fibrotic disease in the lung. A number of studies have shown that the actions of MMPs can result in the release of growth factors and cytokines. These profibrotic factors require proteolytic processing for their activation or release from extracellular matrix or carrier proteins before they can exert their activity. In fact, the proteolytic activity processing of several key factors involved in the pathogenesis of pulmonary fibrosis such as insulin-like growth factor (IGF), TGF-β, and TNF-α occur through the actions of MMPs, thereby activating or releasing them from inhibitory protein-protein interactions. For example, IGFs in vivo are sequestered by six high affinity IGF binding proteins (IGFBPs1-6), preventing their ability to interact with IGF receptors. Studies examining adults and children IPF and interstitial lung disease show that aside IPF, IGFBP-3 and IGBP-2 levels are increased in IPF BAL fluid. MMPs have recently been shown to regulate the cleavage of IGF binding proteins, thereby liberating the complexed ligand to affect IGF actions in target cells. Observations have also shown that the gelatinases, MMP-9 and MMP-2 may be involved in proteolytic activation of latent TGF-β complexes. Furthermore, the MMP inhibitor Batimastat reduces MMP-9 activity in BAL fluid, which was associated with decreased amount of TGF-β and TNF-α.

Pulmonary fibrosis can be an all too common consequence of an acute inflammatory response of the lung to a host of inciting events. Chronic lung injury due to fibrotic changes can result from an identifiable inflammatory event or an insidious, unknown event. The inflammatory process can include infiltration of various inflammatory cell types, such as
neutrophils and macrophages, the secretion of inflammatory cytokines and chemokines and the secretion of matrix remodeling proteinases.

Increased CCL18 Levels in Fibrotic Diseases

[0090] The expression and regulation of cysteine-cysteine (CC) chemokine ligand 18 (CCL18), a marker of alternative activation, by human alveolar macrophages (AMs) is increased in patients with pulmonary fibrosis and correlates negatively with pulmonary function test parameters. Thus, CCL18 is an ideal diagnostic marker for pulmonary fibrosis.

[0091] The peptide of the present invention was tested using the assays described in Examples 14-15 for their effect as active therapeutic agents in the prophylaxis and/or treatment of fibrotic diseases and disorders.

Inflammatory Disease

[0092] Inflammation is the final common pathway of various insults, such as infection, trauma, and allergies to the human body. It is characterized by activation of the immune system with recruitment of inflammatory cells, production of pro-inflammatory cells and production of pro-inflammatory cytokines. Most inflammatory diseases and disorders are characterized by abnormal accumulation of inflammatory cells including monocytes/macrophages, granulocytes, plasma cells, lymphocytes and platelets. Along with tissue endothelial cells and fibroblasts, these inflammatory cells release a complex array of lipids, growth factors, cytokines and destructive enzymes that cause local tissue damage.

[0093] One form of inflammatory response is neutrophilic inflammation which is characterized by infiltration of the inflamed tissue by neutrophil polymorphonuclear leukocytes (PMN), which are a major component of the host defense. Tissue infection by extracellular bacteria represents the prototype of this inflammatory response. On the other hand, various non-infectious diseases are characterized by extravascular recruitment of neutrophils. This group of inflammatory diseases includes chronic obstructive pulmonary disease, adult respiratory distress syndrome, some types of immune-complex alveolitis, cystic fibrosis, bronchitis, bronchiectasis, emphysema, glomerulonephritis, rheumatoid arthritis, gouty arthritis, ulcerative colitis, certain dermatoses such as psoriasis and vasculitis. In these conditions neutrophils are thought to play a crucial role in the development of tissue injury which, when persistent, can lead to the irreversible destruction of the normal tissue architecture with consequent organ dysfunction. Tissue damage is primarily caused by the activation of neutrophils followed by their release of proteinases and increased production of oxygen species.

[0094] Chronic obstructive pulmonary disease (COPD) is described by the progressive development of airflow limitation that is not fully reversible. Most patients with COPD have three pathological conditions: bronchitis, emphysema and mucus plugging. This disease is characterized by a slowly progressive and irreversible decrease in forced expiratory volume in the first second of expiration (FEV1), with relative preservation of forced vital capacity (FVC). In both asthma and COPD there is significant, but distinct, remodeling of airways. Most of the airflow obstruction is due to two major components, alveolar destruction (emphysema) and small airways obstruction (chronic obstructive bronchitis). COPD is mainly characterized by profound mucus cell hyperplasia. Neutrophil infiltration of the patient’s lungs is a primary characteristic of COPD. Elevated levels of proinflammatory cytokines, like TNF-α, and especially chemokines like interleukin-8 (IL-8) and growth-regulated oncogene-α (GRO-α) play a very important role in pathogenesis of this disease. Platelet thromboxane synthesis is also enhanced in patients with COPD. Most of the tissue damage is caused by activation of neutrophils followed by their release of metalloproteinases, and increased production of oxygen species.

[0095] TNF-α has several biologic activities that are important in homeostasis as well as in pathophysiological conditions. The main sources of TNF-α are monocytes-macrophages, T-lymphocytes and mast cells. The finding that anti-TNF-α antibodies (cA2) are effective in the treatment of patients suffering from rheumatoid arthritis (RA) intensified the interest to find new TNF-α inhibitors as possible potent medicaments for RA. Rheumatoid arthritis is an autoimmune chronic inflammatory disease characterized by irreversible pathological changes of the joints. In addition to RA, TNF-α antagonists are also applicable to several other pathological conditions and diseases such as spondylitis, osteoarthritis, gout and other arthritic conditions, sepsis, septic shock, toxic shock syndrome, atopic dermatitis, contact dermatitis, psoriasis, glomerulonephritis, lupus erythematosus, scleroderma, asthma, cachexia, chronic obstructive lung disease, congestive heart failure, insulin resistance, lung (pulmonary) fibrosis, multiple sclerosis, Crohn’s disease, ulcerative colitis, viral infections and AIDS.

[0096] The term “immunoinflammatory disorder” encompasses a variety of conditions, including autoimmune diseases, proliferative skin diseases, and inflammatory dermatoses. Immunoinflammatory disorders result in the destruction of healthy tissue by an inflammatory process, dysregulation of the immune system, and unwanted proliferation of cells. Examples of immunoinflammatory disorders are acne vulgaris; acute respiratory distress syndrome; Addison’s disease; allergic rhinitis; allergic intraocular inflammatory diseases, antineutrophil cytoplasmic antibody (ANCA)-associated small-vessel vasculitis; ankylosing spondylitis; arthritis, asthma; atherosclerosis; atopic dermatitis; autoimmune hepatitis; autoimmune hemolytic anemia; autoimmune hepatitis; Behcet’s disease; Bell’s palsy; bullous pemphigoid; cerebral ischemia; chronic obstructive pulmonary disease; cirrhosis; Cogan’s syndrome; contact dermatitis; COPD; Crohn’s disease; Cushing’s syndrome; dermatomyositis; diabetes mellitus; discoid lupus erythematosus; eosinophilic fasciitis; erythema nodosum; exfoliative dermatitis; fibromyalgia; focal glomerulosclerosis; focal segmental glomerulosclerosis; giant cell arteritis; gout; gouty arthritis; graft versus host disease; hand eczema; Henoch-Schönlein purpura; herpes gestationis; hirsutism; idiopathic cerealsclerosis; idiopathic pulmonary fibrosis; idiopathic thrombocteytopenic purpura; immune thrombocytoopenic purpura inflammatory bowel or gastrointestinal disorders, inflammatory dermatoses; lichen planus; lupus nephritis; lymphomatous tracheobronchitis; macular edema; multiple sclerosis; myasthenia gravis; myositis; nonspecific fibrosing lung disease; osteoarthritis; pancreatitis; pemphigoid gestations; pemphigus vulgaris; periodontitis; polyarteritis nodosa; polymyalgia rheumatica; purpuric scroti; pruritus; inflammation; psoriasis; psoriatic arthritis; pulmonary histoplasmosis; rheumatoid arthritis; relapsing polychondritis; rosacea caused by sarcoidosis; rosacea caused by sclerodema; rosacea caused by Sweet’s syndrome; rosacea caused by systemic lupus erythematosus; rosacea caused by urticaria; rosacea...
caused by zoster-associated pain; sarcoidosis; scleroderma; segmental glomerulosclerosis; septic shock syndrome; shoulder tendinitis or bursitis; Sjögren’s syndrome; Still’s disease; stroke-induced brain cell death; Sweet’s disease; systemic lupus erythematosus; systemic sclerosis; Takayasu’s arteritis; temporal arteritis; toxic epidermal necrolysis; transplant-rejection and transplant-rejection-related syndromes; tuberculosis; type-1 diabetes; ulcerative colitis; uveitis; vasculitis; and Wegener’s granulomatosis.

As used herein, “non-dermal inflammatory disorders” include, for example, rheumatoid arthritis, inflammatory bowel disease, asthma, and chronic obstructive pulmonary disease. By “dermal inflammatory disorders” or “inflammatory dermatoses” is meant an inflammatory disorder selected from psoriasis, guttate psoriasis, inverse psoriasis, pustular psoriasis, erythrodermic psoriasis, acute febrile neutrophilic dermatosis, eczema, asthenic eczema, dyshidrotic eczema, vesicular palmoplantar eczema, acne vulgaris, atopic dermatitis, contact dermatitis, allergic contact dermatitis, dermatomyositis, exfoliative dermatitis, hand eczema, pompholyx, rosacea, rosacea caused by sarcoidosis, rosacea caused by scleroderma, rosacea caused by Sweet’s syndrome, rosacea caused by systemic lupus erythematosus, rosacea caused by urticaria, rosacea caused by zoster-associated pain, Sweet’s disease, neutrophilic hidradenitis, sterile pseudomonas, drug eruptions, seborrheic dermatitis, pityriasis rosea, cutaneous lupus disease, pruritic urticarial papules and plaques of pregnancy, Stevens-Johnson syndrome and toxic epidermal necrolysis, tattoo reactions, Wells syndrome (eosinophilic cellulitis), reactive arthritis (Reiter’s syndrome), bowel-associated dermatosis-arthritis syndrome, rheumatoid neutrophilic dermatosis, neutrophilic eccrine hidradenitis, neutrophilic dermatosis of the dorsal hands, balanitis circumscription, balanitis, Behçet’s disease, erythema annular centrifugum, erythema dyschromicum perstans, erythema multiforme, granuloma annulare, hand dermatitis, lichen nitidus, lichen planus, lichen sclerosus et atrophicus, lichen simplex chronicus, lichen spinulosus, mnnular dermatitis, pyoderma gangrenosum, sarcoidosis, subcutaneous pustular dermatosis, urticaria, and transient acantholytic dermatitis.

By “proliferative skin disease” is meant a benign or malignant disease that is characterized by accelerated cell division in the epidermis or dermis. Examples of proliferative skin diseases are psoriasis, atopic dermatitis, nonspecific dermatitis, primary irritant contact dermatitis, allergic contact dermatitis, basal and squamous cell carcinomas of the skin, lamellar ichthyosis, epidermolytic hyperkeratosis, premalignant keratoses, acne, and seborrheic dermatitis. As will be appreciated by one skilled in the art, a particular disease, disorder, or condition may be characterized as being both a proliferative skin disease and an inflammatory dermatosis. An example of such a disease is psoriasis.

Symptoms and signs of inflammation associated with specific conditions include:

- **Rheumatoid arthritis:** pain, swelling, warmth and tenderness of the involved joints; generalized and morning stiffness.
- **Insulin-dependent diabetes mellitus-insulin:** this condition can lead to a variety of complications with an inflammatory component, including: retinopathy, neuropathy, nephropathy; coronary artery disease, peripheral vascular disease, and cerebrovascular disease;
- **Autoimmune thyroiditis:** weakness, constipation, shortness of breath, puffiness of the face, hands and feet, peripheral edema, bradycardia;
- **Multiple sclerosis:** spasticity, blurry vision, vertigo, limb weakness, paresthesias;
- **Uveoretinitis:** decreased night vision, loss of peripheral vision;
- **Lupus erythematosus:** joint pain, rash, photosensitivity, fever, muscle pain, puffiness of the hands and feet, abnormal urinalysis (hematuria, cylindruria, proteinuria), glomerulonephritis, cognitive dysfunction, vessel thrombosis, pericarditis;
- **Scleroderma:** Raynaud’s disease; swelling of the hands, arms, legs and face; skin thickening; pain, swelling and stiffness of the fingers and knees, gastrointestinal dysfunction, restrictive lung disease; pericarditis; renal failure;
- **Other arthritic conditions having an inflammatory component such as rheumatoid spondylitis, osteoarthritis, septic arthritis and polymyalgia:** fever, pain, swelling, tenderness;
- **Other inflammatory brain disorders such as meningitis, Alzheimer’s disease, AIDS dementia encephalitis:** photophobia, cognitive dysfunction, memory loss;
- **Other inflammatory eye inflammations such as retinitis:** decreased visual acuity;
- **Other inflammatory skin disorders such as, eczema, other dermatitis (e.g., atopic, contact), psoriasis, burns induced by UV radiation (sun rays and similar UV sources):** erythema, pain, scaling, swelling, tenderness;
- **Inflammatory bowel disease, such as Crohn’s disease, ulcerative colitis:** pain, diarrhea, constipation, rectal bleeding, fever, arthritis;
- **Asthma:** shortness of breath, wheezing;
- **Other allergic disorders such as allergic rhinitis:** sneezing, itching, runny nose;
- **Conditions associated with acute trauma such as cerebral injury following stroke-sensory loss, motor loss, cognitive loss:**
- **Heart tissue injury due to myocardial ischemic:** pain, shortness of breath;
- **Lung injury such as that which occurs in adult respiratory distress syndrome:** shortness of breath, hyperventilation, decreased oxygenation, pulmonary infiltrates;
- **Inflammation accompanying infection, such as sepsis, septic shock, toxic shock syndrome:** fever, respiratory failure, tachycardia, hypotension, leukocytosis;
- **Other inflammatory conditions associated with particular organs or tissues, such as:**
  (i) nephritis (e.g., glomerulonephritis): oliguria, abnormal urinalysis;
  (ii) inflamed appendix: fever, pain, tenderness, leukocytosis;
  (iii) gout: pain, tenderness, swelling and erythema of the involved joint, elevated serum and/or urinary uric acid;
  (iv) inflamed gall bladder: abdominal pain and tenderness, fever, nausea, leukocytosis;
  (v) congestive heart failure: shortness of breath, raales, peripheral edema;
  (vi) Type II diabetes: end organ complications including cardiovascular, ocular, renal, and peripheral vascular disease;
(vii) lung (pulmonary) fibrosis:—hyperventilation, shortness of breath, decreased oxygenation; (viii) vascular disease, such as atherosclerosis and restenosis:—pain, loss of sensation, diminished pulses, loss of function; and (ix) alloimmunity leading to transplant rejection:—pain, tenderness, fever.

A human peptide is “active” in an inflammatory disease if the inhibition is >50% in one of the assays described below. Inhibition (as percentage) was calculated using the following formula: % inhibition = (1-concentration of cytokines in sample/concentration of cytokines in positive control) x 100. The positive control refers to stimulated samples, not treated with substances.

The peptide of the present invention was tested using the assays described in Examples 1-7, 9-17 for their effect as active therapeutic agents in the prophylaxis and/or treatment of inflammatory diseases and disorders.

Neurodegenerative Disease

The present invention also relates generally to the fields of neurology and psychiatry and to methods of protecting the cells of a mammalian central nervous system from damage or injury.

Injuries or trauma of various kinds to the central nervous system (CNS) or the peripheral nervous system (PNS) can produce profound and long-lasting neurological and/or psychiatric symptoms and disorders. One fact that this can take is the progressive death of neurons or other cells of the central nervous system (CNS), i.e., neurodegeneration or neuronal degeneration.

Neuronal degeneration as a result of, for example; Alzheimer’s disease, multiple sclerosis, cerebral-vascular accidents (CVAs)/stroke, traumatic brain injury, spinal cord injuries, degeneration of the optic nerve, e.g., ischemic optic neuropathy or retinal degeneration and other central nervous system disorders is an enormous medical and public health problem by virtue of both its high incidence and the frequency of long-term sequelae. Animal studies and clinical trials have shown that amino acid transmitters (especially glutamate), oxidative stress and inflammatory reactions contribute strongly to cell death in these conditions. Upon injury or upon ischemic insult, damaged neurons release massive amounts of the neurotransmitter glutamate, which is excitotoxic to the surrounding neurons. Glutamate is a negatively charged amino acid that is an excitatory synaptic transmitter in the mammalian nervous system. While the concentration of glutamate can reach the millimolar range in nerve terminals its extracellular concentration is maintained at a low level to prevent neurotoxicity. It has been noted that glutamate can be toxic to neurons if presented at a high concentration. The term “excitotoxicity” has been used to describe the cytotoxic effect that glutamate (and other such excitatory amino acids) can have on neurons when applied at high dosages.

Patients with injury or damage of any kind to the central (CNS) or peripheral (PNS) nervous system including the retina may benefit from neuroprotective methods. This nervous system injury may take the form of an abrupt insult or an acute injury to the nervous system as in, for example, acute neurodegenerative disorders including, but not limited to; acute injury, hypoxia-ischemia or the combination thereof resulting in neuronal cell death or compromise. Acute injury includes, but is not limited to, traumatic brain injury (TBI) including, closed, blunt or penetrating brain trauma, local brain trauma, diffuse brain damage, spinal cord injury, intracranial or intravertebral lesions (including, but not limited to, contusion, penetration, shear, compression or laceration lesions of the spinal cord or whiplash shaken infant syndrome).

In addition, deprivation of oxygen or blood supply in general can cause acute injury as in hypoxia and/or ischemia including, but not limited to, cerebrovascular insufficiency, cerebral ischemia or cerebral infarction (including cerebral ischemia or infarctions originating from embolic occlusion and thrombosis, retinal ischemia (diabetic or otherwise), glaucoma, retinal degeneration, multiple sclerosis, toxic and ischemic optic neuropathy, reperfusion following acute ischemia, perinatal hypoxic-ischemic injury, cardiac arrest or intracranial hemorrhage of any type (including, but not limited to, epidural, subdural, subarachnoid or intracerebral hemorrhage).

Trauma or injury to tissues of the nervous system may also take the form of more chronic and progressive neurodegenerative disorders, such as those associated with progressive neuronal cell death or compromise over a period of time including, but not limited to, Alzheimer’s disease, Pick’s disease, diffuse Lewy body disease, progressive supranuclear palsy (Steel-Richardson syndrome), multisystem degeneration (Shy-Drager syndrome), chronic epileptic conditions associated with neurodegeneration, motor neuron diseases (amyotrophic lateral sclerosis), multiple sclerosis, degenerative ataxias, cortical basal degeneration, ALS-Parkinson’s-dementia complex of Guam, subacute sclerosing panencephalitis, Huntington’s disease, Parkinson’s disease, synucleinopathies (including multiple system atrophy), primary progressive aphasia, striatognital degeneration, Machado-Joseph disease or spinocerebellar ataxia type 3 and olivopontocerebellar degenerations, bulbar and pseudobulbar palsy, spinal and spinobulbar muscular atrophy (Kennedy’s disease), primary lateral sclerosis, familial spastic paraplegia, Werdnig-Hoffmann disease, Kugelberg-Welander disease, Tay-Sach’s disease, Sandhoff disease, familial spastic disease, Wohlfart-Kugelberg-Welander disease, spastic paraparesis, progressive multifocal leukoencephalopathy, familial dysautonomia (Riley-Day syndrome) or prion diseases (including, but not limited to Creutzfeldt-Jakob disease, Gerstmann-Strussler-Scheinker disease, Kuru disease or familial insomnia).

In addition, trauma and progressive injury to the nervous system can take place in various psychiatric disorders, including but not limited to, progressive, deteriorating forms of bipolar disorder or schizoaffective disorder or schizophrenia, impulse control disorders, obsessive compulsive disorder (OCD), behavioral changes in temporal lobe epilepsy and personality disorders.

In one preferred embodiment the compounds of the invention would be used to provide neuroprotection in disorders involving trauma and progressive injury to the nervous system in various psychiatric disorders. These disorders would be selected from the group consisting of; schizoaffective disorder, schizophrenia, impulse control disorders, obsessive compulsive disorder (OCD) and personality disorders.

In addition, trauma and injury may take the form of disorders associated with overt and extensive memory loss including, but not limited to, neurodegenerative disorders associated with age-related dementia, vascular dementia, diffuse white matter disease (Binswanger’s disease), dementia...
of endocrine or metabolic origin, dementia of head trauma and diffuse brain damage, dementia pugilistica or frontal lobe dementia, including but not limited to Pick’s Disease.

[0130] Other disorders associated with neuronal injury include, but are not limited to, disorders associated with chemical, toxic, infectious and radiation injury of the nervous system including the retina, injury during fetal development, prematurity at time of birth, anoxic-ischemia, injury from hepatic, glycemic, uremic, electrolyte and endocrine origin, injury of psychiatric origin (including, but not limited to, psychopathology, depression or anxiety), injury from peripheral diseases and plexopathies (including plexus palsy) or injury from neuropathy (including neuropathy selected from multifocal, sensory, motor, sensory-motor, autonomic, sensory-autonomic or demyelinating mononeuropathy (including, but not limited to, Guillain-Barré syndrome or chronic inflammatory demyelinating polyradiculoneuropathy) or those neuropathies originating from infections, inflammation, immune disorders, drug abuse, pharmacological treatments, toxins, trauma (including, but not limited to compression, crush, laceration or segmentation traumas), metabolic disorders (including, but not limited to, endocrine or paraneoplastic), Charcot-Marie-Tooth disease (including, but not limited to, type 1a, 1b, 2, 4a or 1-X linked), Friedreich’s ataxia, metachromatic leukodystrophy, Refsum’s disease, adenoncyclo-neuropathy, ataxia-telangiectasia, Djerine-Sottas (including, but not limited to, types A or B), Lambert-Eaton syndrome or disorders of the cranial nerves).

[0131] Further indications are cognitive disorders. The term “cognitive disorder” shall refer to anxiety disorders, delirium, dementia, amnestic disorders, disassociative disorders, eating disorders, mood disorders, schizophrenia, psychotic disorders, sexual and gender identity disorders, sleep disorders, somatoform disorders, acute stress disorder, obsessive-compulsive disorder, panic disorder, posttraumatic stress disorder, specific phobia, social phobia, substance withdrawal delirium, Alzheimer’s disease, Creutzfeldt-Jakob disease, head trauma, Huntington’s disease, HIV disease, Parkinson’s disease, Pick’s disease, learning disorders, motor skills disorders, developmental coordination disorder, communication disorders, phonological disorder, perseverative developmental disorders, Asperger’s disorder, autistic disorder, childhood disintegrative disorder, Rett’s disorder, pervasive developmental disorder, attention-deficit/hyperactivity disorder (ADHD), conduct disorder, oppositional defiant disorder, pica, rumination disorder, tic disorders, chronic motor or vocal tic disorder, Tourette’s syndrome, elimination disorders, encopresis, enuresis, selective mutism, separation anxiety disorder, dissociative amnesia, depersonalization disorder, dissociative fugue, dissociative identity disorder, anorexia nervosa, bulimia nervosa, bipolar disorders, schizophrenia, schizoaffective disorder, delusional disorder, psychotic disorder, shared psychotic disorder, delusions, hallucinations, substance-induced psychotic disorder, obsessive disorders, sexual pain disorders, dyspareunia, vaginismus, sexual dysfunction, paraphilias, dysnomias, breathing-related sleep disorder, circadian rhythm sleep disorder, hypersomnia, insomia, narcolepsy, dysomnia, parasomnia, nightmare disorder, sleep terror disorder, sleepwalking disorder, parasomnia, body dysmorphic disorder, conversion disorder, hypochondriasis, pain disorder, somatization disorder, alcohol related disorders, amphetamine related disorders, caffeine related disorders, cannabis related disorders, cocaine related disorders, hallucinogen related disorders, opioid related disorders, nicotine related disorders, phencyclidine-related disorder, abuse, persisting amnestic disorder, intoxication, withdrawal.

[0132] The term “bipolar and clinical disorders” shall refer to adjustment disorders, anxiety disorders, delirium, dementia, amnestic and other cognitive disorders, disorders usually first diagnosed in infancy (e.g.), childhood, or adolescence, dissociative disorders (e.g. dissociative amnesia, depersonalization disorder, dissociative fugue and dissociative identity disorder), eating disorders, factitious disorders, impulse-control disorders, mental disorders due to a general medical condition, mood disorders, other conditions that may be a focus of clinical attention, personality disorders, schizophrenia and other psychotic disorders, sexual and gender identity disorders, sleep disorders, somatoform disorders, substance-related disorders, generalized anxiety disorder (e.g. acute stress disorder, posttraumatic stress disorder), panic disorder, phobia, agoraphobia, obsessive-compulsive disorder, stress, acute stress disorder, anxiety neurosis, nervousness, phobia, posttraumatic stress disorder, posttraumatic stress disorder (PTSD), abuse, obsessive-compulsive disorder (OCD), manic depressive psychosis, specific phobias, social phobia, adjustment disorder with anxious features.

[0133] Examples for disorders usually first diagnosed in infancy, childhood, or adolescence are: mental retardation, learning disorders, mathematics disorder, reading disorder, disorder of written expression, motor skills disorders, developmental coordination disorder, communication disorders, expressive language disorder, phonological disorder, mixed receptive-expressive language disorder, stuttering, pervasive developmental disorders, Asperger’s disorder, autistic disorder, childhood disintegrative disorder, Rett’s disorder, pervasive developmental disorder, attention-deficit/hyperactivity disorder (ADHD), conduct disorder, oppositional defiant disorder, feeding disorder of infancy or early childhood, pica, rumination disorder, tic disorders, chronic motor or vocal tic disorder, Tourette’s syndrome, elimination disorders, encopresis, enuresis, selective mutism, separation anxiety disorder, reactive attachment disorder of infancy or early childhood, stereotypic movement disorder.

[0134] Examples for substance-related disorders are: alcohol related disorders, amphetamine related disorders, caffeine related disorders, cannabis related disorders, cocaine related disorders, hallucinogen related disorders, inhalant related disorders, nicotine related disorders, opioid related disorders, psychotic disorder, psychogenic disorder, phencyclidine-related disorder, abuse, persisting amnestic disorder, anxiety disorder, persisting dementia, dependence, intoxication, intoxication delirium, mood disorder, psychogenic disorder, withdrawal, withdrawal delirium, sexual dysfunction, sleep disorder.

[0135] The term “neuroprotection” as used herein shall mean; inhibiting, preventing, ameliorating or reducing the severity of the dysfunction, degeneration or death of nerve cells, axons or their supporting cells in the central or peripheral nervous system of a mammal, including a human. This includes the treatment or prophylaxis of a neurodegenerative disease; protection against excitotoxicity or ameliorating the cytotoxic effect of a compound (for example, an excitatory amino acid such as glutamate; a toxin; or a prophylactic or therapeutic compound that exerts an immediate or delayed side effect including but not limited to the immediate or delayed induction of apoptosis) in a patient in need thereof.
The term “a patient in need of treatment with a neuroprotective drug” as used herein will refer to any patient who currently has or may develop any of the above syndromes or disorders, or any disorder in which the patient’s present clinical condition or prognosis could benefit from providing neuroprotection to prevent the development, extension, worsening or increased resistance to treatment of any neurological or psychiatric disorder.

The term “treating” or “treatment” as used herein, refers to any indicia of success in the prevention or amelioration of an injury, pathology or condition, including any objective or subjective parameter such as abatement, remission; diminishing of symptoms or making the injury, pathology, or condition more tolerable to the patient; slowing in the rate of degeneration or decline; making the final point of degeneration less debilitating; or improving a subject’s physical or mental well-being. The treatment or amelioration of symptoms can be based on objective or subjective parameters; including the results of a physical examination, neurological examination, and/or psychiatric evaluations.

In some embodiments this invention provides methods of neuroprotection. In certain embodiments, these methods comprise administering a therapeutically effective amount of the peptide of the invention to a patient who has not yet developed overt, clinical signs or symptoms of injury or damage to the cells of the nervous system but who may be in a high risk group for the development of neuronal damage because of injury or trauma to the nervous system or because of some known predisposition either biochemical or genetic or the finding of a verified biomarker of one or more of these disorders.

Thus, in some embodiments, the methods and compositions of the present invention are directed toward neuroprotection in a subject who is at risk of developing neuronal damage but who has not yet developed clinical evidence. This patient may simply be at “greater risk” as determined by the recognition of any factor in a subject’s, or their families, medical history, physical exam or testing that is indicative of a greater than average risk for developing neuronal damage. Therefore, this determination that a patient may be at a “greater risk” by any available means can be used to determine whether the patient should be treated with the methods of the present invention.

Accordingly, in an exemplary embodiment, subjects who may benefit from treatment by the methods and peptide of this invention can be identified using accepted screening methods to determine risk factors for neuronal damage. These screening methods include, for example, conventional work-ups to determine risk factors including but not limited to: for example, head trauma, either closed or penetrating, CNS infections, bacterial or viral, cerebrovascular disease including but not limited to stroke, brain tumors, brain edema, cysticercosis, porphyria, metabolic encephalopathy, drug withdrawal including but not limited to sedative-hypnotic or alcohol withdrawal, abnormal perinatal history including anoxia at birth or birth injury of any kind, cerebral palsy, learning disabilities, hyperactivity, history of febrile convulsions as a child, history of status epilepticus, family history of epilepsy or any seizure related disorder, inflammatory disease of the brain including lupus, drug intoxication either direct or by placental transfer, including but not limited to cocaine poisoning, parental consanguinity, and treatment with medications that are toxic to the nervous system including psychotropic medications.

The determination of which patients may benefit from treatment with a neuroprotective drug in patients who have no clinical signs or symptoms may be based on a variety of “surrogate markers” or “biomarkers”.

As used herein, the terms “surrogate marker” and “biomarker” are used interchangeably and refer to any anatomical, biochemical, structural, electrical, genetic or chemical indicator or marker that can be reliably correlated with the present existence or future development of neuronal damage. In some instances, brain-imaging techniques, such as computer tomography (CT), magnetic resonance imaging (MRI) or positron emission tomography (PET), can be used to determine whether a subject is at risk for neuronal damage. Suitable biomarkers for the methods of this invention include, but are not limited to: the determination by MRI, CT or other imaging techniques, of sclerosis, atrophy or volume loss in the hippocampus or overt mesial temporal sclerosis (MTS) or similar relevant anatomical pathology; the detection in the patient’s blood, serum or tissues of a molecular species such as a protein or other biochemical biomarker, e.g., elevated levels of ciliary neurotrophic factor (CNTF) or elevated serum levels of a neuronal degradation product; or other evidence from surrogate markers or biomarkers that the patient is in need of treatment with a neuroprotective drug.

It is expected that many more such biomarkers utilizing a wide variety of detection techniques will be developed in the future. It is intended that any such marker or indicator of the existence or possible future development of neuronal damage, as the latter term is used herein, may be used in the methods of this invention for determining the need for treatment with the compounds and methods of this invention.

A determination that a subject has, or may be at risk for developing, neuronal damage would also include, for example, a medical evaluation that includes a thorough history, a physical examination, and a series of relevant bloods tests. It can also include an electroencephalogram (EEG), CT, MRI or PET scan. A determination of an increased risk of developing neuronal damage or injury may also be made by means of genetic testing, including gene expression profiling or proteomic techniques. For psychiatric disorders that may be stabilized or improved by a neuroprotective drug, e.g., bipolar disorder, schizoaffective disorder, schizophrenia, impulse control disorders, etc. the above tests may also include a present state exam and a detailed history of the course of the patient’s symptoms such as mood disorder symptoms and psychotic symptoms over time and in relation to other treatments the patient may have received over time, e.g., a life chart. These and other specialized and routine methods allow the clinician to select patients in need of therapy using the methods and formulations of this invention. In some embodiments of the present invention peptide suitable for use in the practice of this invention will be administered either singly or concomitantly with at least one or more other compounds or therapeutic agents, e.g., with other neuroprotective drugs or antiepileptic drugs, anticonvulsant drugs. In these embodiments, the present invention provides methods to treat or prevent neuronal injury in a patient. The method includes the step of; administering to a patient in need of treatment, an effective amount of the peptide disclosed herein in combination with an effective amount of one or more other compounds or therapeutic agents that have the ability to provide neuroprotection or to treat or prevent seizures or epileptoge-
nosis or the ability to augment the neuroprotective effects of the compounds of the invention.

[0145] As used herein the term “combination administration” of a compound, therapeutic agent or known drug with the peptide of the present invention means administration of the drug and the one or more compounds at such time that both the known drug and the peptide will have a therapeutic effect. In some cases this therapeutic effect will be synergistic. Such concomitant administration can involve concurrent (i.e. at the same time), prior, or subsequent administration of the drug with respect to the administration of the peptide of the present invention. A person of ordinary skill in the art would have no difficulty determining the appropriate timing, sequence and dosages of administration for particular drugs and peptide of the present invention.

[0146] The said one or more other compounds or therapeutic agents may be selected from compounds that have one or more of the following properties: antioxidant activity; NMDA receptor antagonist activity, augmentation of endogenous GABA inhibition; NO synthase inhibitor activity; iron binding ability, e.g., an iron chelator; calcium binding ability, e.g., a Ca (II) chelator; zinc binding ability, e.g., a Zn (II) chelator; the ability to effectively block sodium or calcium ion channels, or to open potassium or chloride ion channels in the CNS of a patient.

[0147] The peptide of the present invention was tested using the assays described in Examples 1-7, 9-17 for their effect as active therapeutic agents in the prophylaxis and/or treatment of neurodegenerative diseases and disorders.

**Heart and Vascular Disease**

[0148] Heart disease is a general term used to describe many different heart conditions. For example, coronary artery disease, which is the most common heart disease, is characterized by constriction or narrowing of the arteries supplying the heart with oxygen-rich blood, and can lead to myocardial infarction, which is the death of a portion of the heart muscle. Heart failure is a condition resulting from the inability of the heart to pump an adequate amount of blood through the body. Heart failure is not a sudden, abrupt stop of heart activity, but rather, typically develops slowly over many years, as the heart gradually loses its ability to pump blood efficiently. Risk factors for heart failure include coronary artery disease, hypertension, valvular heart disease, cardiomyopathy, disease of the heart muscle, obesity, diabetes, and/or a family history of heart failure.

[0149] Examples of cardiovascular diseases and disorders are: anemia, unstable angina, angina pectoris, angioneurotic edema, aortic valve stenosis, aortic aneurysm, arrhythmia, arrhythmogenic right ventricular dysplasia, arteriosclerosis, arteriovenous malformations, atrial fibrillation, Behcet syndrome, bradycardia, cardiac tamponade, cardiomegaly, congestive cardiomopathy, hypertrophic cardiomyopathy, restrictive cardiomyopathy, cardiomyopathy, cerebral hemorrhage, Churg-Strauss syndrome, diabetes, Ebstein’s Anomaly, Eisenmenger complex, cholesterol embolism, bacterial endocarditis, fibromuscular dysplasia, congenital heart defects, heart diseases, congestive heart failure, heart valve diseases, heart attack, epidural hematoma, hematoma, subdural, Hippel-Lindau syndrome, hyperemia, hypertension, pulmonary hypertension, cardiac hypertrophy, left ventricular hypertrophy, right ventricular hypertrophy, hypoplastic left heart syndrome, hypertension, intermittent claudication, ischemic heart disease, Klippel-Trenaunay-Weber syndrome, lateral medullary syndrome, long QT syndrome mitral valve prolapse, moyamoya disease, mucocutaneous lymph node syndrome, myocardial infarction, myocardial ischemia, myocarditis, pericarditis, peripheral vascular diseases, phlebitis, polyarteritis nodosa, pulmonary atresia, Raynaud disease, Sneddon syndrome, superior vena cava syndrome, syndrome X, tachycardia, Takayasu’s arteritis, hereditary hemorrhagic telangiectasia, telangiectasia, temporal arteritis, tetralogy of Fallot, thromboangiitis obliterans, thrombosis, thromboembolism, tricuspid atresia, varicose veins, vascular diseases, vasculitis, vasospasm, ventricular fibrillation, Williams syndrome, peripheral vascular disease, varicose veins and leg ulcers, deep vein thrombosis, Wolff-Parkinson-White syndrome.

[0150] Vascular diseases are often the result of decreased perfusion in the vascular system or physical or biochemical injury to the blood vessel.

[0151] Peripheral vascular disease (PVD) is defined as a disease of blood vessels often encountered as narrowing of the vessels of the limbs. There are two main types of these disorders, functional disease which doesn’t involve defects in the blood vessels but rather arises from stimuli such as cold, stress, or smoking, and organic disease which arises from structural defects in the vasculature such as atherosclerotic lesions, local inflammation, or traumatic injury. This can lead to occlusion of the vessel, aberrant blood flow, and ultimately to tissue ischemia.

[0152] One of the more clinically significant forms of PVD is peripheral artery disease (PAD). PAD is often treated by angioplasty and implantation of a stent or by artery bypass surgery. Clinical presentation depends on the location of the occluded vessel. For example, narrowing of the artery that supplies blood to the intestine can result in severe postprandial pain in the lower abdomen resulting from the inability of the occluded vessel to meet the increased oxygen demand arising from digestive and absorptive processes. In severe forms the ischemia can lead to intestinal necrosis. Similarly, PAD in the leg can lead to intermittent pain, usually in the calf, that comes and goes with activity. This disorder is known as intermittent claudication (IC) and can progress to persistent pain while resting, ischemic ulceration, and even amputation.

[0153] Peripheral vascular disease is also manifested in atherosclerotic stenosis of the renal artery, which can lead to renal ischemia and kidney dysfunction.

[0154] One disease in which vascular diseases and their complications are very common is diabetes mellitus. Diabetes mellitus causes a variety of physiological and anatomical irregularities, the most prominent of which is the inability of the body to utilize glucose normally, which results in hyperglycemia. Chronic diabetes can lead to complications of the vascular system which include atherosclerosis, abnormalities involving large and medium size blood vessels (macroangiopathy) and abnormalities involving small blood vessels (microangiopathy) such as arteriolas and capillaries.

[0155] Patients with diabetes mellitus are at increased risk of developing one or more foot ulcers as a result of established long-term complications of the disease, which include impaired nerve function (neuropathy) and/or ischemia. Local tissue ischemia is a key contributing factor to diabetic foot ulceration.

[0156] In addition to large vessel disease, patients with diabetes suffer further threat to their skin perfusion in at least two additional ways. First, by involvement of the non-condui
arteries, which are detrimentally affected by the process of atherosclerosis, and secondly, and perhaps more importantly, by impairment of the microcirculatory control mechanisms (small vessel disease). Normally, when a body part suffers some form of trauma, the body part will, as part of the body’s healing mechanism, experience an increased blood flow. When small vessel disease and ischemia are both present, as in the case of many diabetics, this natural increased blood flow response is significantly reduced. This fact, together with the tendency of diabetics to form blood clots (thrombosis) in the microcirculatory system during low levels of blood flow, is believed to be an important factor in ulcer pathogenesis.

Neuropathy is a general term which describes a disease process which leads to the dysfunction of the nervous system, and is one of the major complications of diabetes mellitus, with no well-established therapies for either its symptomatic treatment or for prevention of progressive decline in nerve function.

The thickening and leakage of capillaries caused by diabetes primarily affect the eyes (retinopathy) and kidneys (nephropathy). The thickening and leakage of capillaries caused by diabetes are also associated with skin disorders and disorders of the nervous system (neuropathy).

The eye diseases associated with diabetes are non-proliferative diabetic retinopathy, proliferative diabetic retinopathy, diabetic maculopathy, glaucoma, cataracts, and the like.

Angiogenesis is a physiological process involving the growth of new blood vessels from pre-existing vessels. Angiogenesis is a normal process in growth and development, as well as in wound healing. However, this is also a fundamental step in the transition of tumors from a dormant state to a malignant state. Angiogenesis occurs in several well-characterized stages. First, biological signals known as angiogenic growth factors activate receptors present on endothelial cells present in pre-existing blood vessels. Second, the activated endothelial cells begin to release enzymes called proteases that degrade the basement membrane in order to allow endothelial cells to escape from the original (parent) vessel walls. The endothelial cells then proliferate into the surrounding matrix and form solid sprouts connecting neighboring vessels. As sprouts extend toward the source of the angiogenic stimulus, endothelial cells migrate, using adhesion molecules, called integrins. These sprouts then form loops to become a fully-fledged vessel lumen as cells migrate to the site of angiogenesis. Sprouting occurs at a rate of several millimeters per day and enables new vessels to grow across gaps in the vasculature.

Therapeutic angiogenesis is the application of specific compounds which may inhibit or induce the creation of new blood vessels in the body in order to combat disease. The presence of blood vessels where there should be none may affect the mechanical properties of a tissue, increasing the likelihood of failure. The absence of blood vessels in a repairing or otherwise metabolically active tissue may retard repair or some other function. Several diseases are the result of failure or insufficient blood vessel formation and may be treated by a local expansion of blood vessels, thus bringing new nutrients to the site, facilitating repair. Other diseases may be created by a local expansion of blood vessels, interfering with normal physiological processes.

Angiogenesis represents an excellent therapeutic target for the treatment of, for example, cardiovascular dis-
eases. It is a potent, physiological process that underlies the natural manner in which the human body responds to a diminution of blood supply to vital organs, namely the production of new collateral vessels to overcome the ischemic insult. [0167] The modern clinical application of the principle "angiogenesis" can be divided into two main areas:

[0168] 1. Anti-angiogenic therapies
[0169] 2. Pro-angiogenic therapies.

[0170] Whereas anti-angiogenic therapies are trying to fight:

[0171] Any type of cancer and malignancies and their metastases in numerous organs, like hemangiomas (because tumors, in general, are nutrition- and oxygen-dependent, thus being in need of adequate blood supply).

[0172] Infectious diseases,

[0173] Vasculitis and excessive angiogenesis in autoimmune disorders such as systemic sclerosis (Scleroderma), multiple sclerosis, Sjogren's disease,

[0174] Vascular malformations in blood and lymph vessels like DiGeorge syndrome, hereditary hemorrhagic telangiectasia, cavernous hemangioma, cutaneous hemangioma, lymphatic malformations, transplant arteriopathy, athroclerosis, vascular anastomoses,

[0175] Adipose tissue in obesity,

[0176] Chronic allograft rejections,

[0177] Skin diseases like psoriasis, warts, allergic dermatitis, scar keloids, pyogenic granulomas, blistersing disease, Kaposis sarcoma in AIDS patients, systemic sclerosis (Scleroderma),

[0178] Eye diseases like persistent hyperplastic vitreous syndrome, diabetic retinopathy, retinopathy of prematurity, choroidal neovascularization,

[0179] Lung diseases like any type of pulmonary hypertension, asthma, nasal polyps, rhinitis, chronic airway inflammation and obstruction (COPD), cystic fibrosis, acute lung injury, bronchiolitis obliterans organizing pneumonia,

[0180] Gastrointestinal tract diseases like inflammatory bowel disease, periodontal disease, ascites, perihepatic adhesions, liver cirrhosis,

[0181] Reproductive system diseases like endometriosis, uterine bleeding, ovarian cysts, ovarian hyperstimulation,

[0182] Bone and joint diseases like arthritis and synovitis, osteomyelitis, osteophyte formation, HIV-induced bone marrow angiogenesis,

[0183] Kidney diseases like early diabetic nephropathy

[0184] the pro-angiogenic therapies are important in the search of new treatment options for diseases characterized or caused by insufficient angiogenesis or vessel regression:

[0185] Nervous system diseases like Alzheimer's disease, amyotrophic lateral sclerosis, diabetic neuropathy, stroke,

[0186] Blood and lymph vessels diseases like diabetic angiopathy, impaired reendothelinization in restenosis, lymphedema,

[0187] Gastrointestinal ulcerations, oral ulcers, mucosal ischemia in Crohn's disease

[0188] Skin diseases like lupus,

[0189] Reproductive system diseases like preeclampsia, menorrhagia,

[0190] Lung diseases like neonatal respiratory distress syndrome, pulmonary fibrosis, emphysema,

[0191] Kidney diseases like nephropathy, glomerulosclerosis, tubulointerstitial fibrosis,

[0192] Bone diseases like osteoporosis, impaired bone fracture healing,

[0193] Heart diseases like ischaemic heart disease, cardiac failure,

[0194] Any type of wound healing disorders.

[0195] Angiogenesis research is also a cutting edge field in cancer research, and traditional therapies, such as radiation therapy, may work in part by targeting the genomically stable endothelial cell compartment, rather than the genomically unstable tumor cell compartment. New blood vessel formation is a relatively fragile process, subject to disruptive interference at several levels. In short, the therapy is the selection agent which is being used to kill the tumor compartment. Tumor cells evolve resistance rapidly due to rapid generation time (days) and genomic instability (variation), whereas endothelial cells are a good target because of a long generation time (months) and genomic stability (low variation). Angiogenesis-based tumor therapy relies on natural and synthetic angiogenesis inhibitors like angiotatin, endostatin and tumstatin. These are proteins that mainly originate as specific fragments pre-existing structural proteins like collagen or plasmogen.

[0196] Recently, the 1st FDA-approved therapy targeted at angiogenesis in cancer came on the market in the US. This is a monoclonal antibody directed against an isofom of VEGF, and the therapy has been approved for use in colorectal cancer in combination with established chemotherapy. Therefore there is a wide medical need for additional medications in the field of angiogenesis.

[0197] In addition, in terms of tissue engineering, medications that influence angiogenesis in vascular grafts are needed. More than 450,000 vascular grafts were used in coronary bypass surgeries annually. Other uses for vascular grafts include treatments for blood vessel aneurysms and fistulas, as well as replacements for diseased arteries in other locations in the body. When possible, the best choice for a replacement vessel is an autograft, where sections of the patient's healthy blood vessels (usually veins) are harvested and implanted in the required location. Many patients, however, especially those with pre-existing vascular disease or patients that have already had autograft procedures, do not have blood vessels that are healthy enough to adequately serve as replacements. In these cases, the most common form of treatment has been the use of synthetic polymeric materials, like ePTEF (extended polytetrafluoroethylene) and Dacron (poly[ethylene terephthalate]), to form either permanent or resorbable replacements for the damaged vessels. In cases where the graft can be of a large diameter (greater than 5-6 mm), the synthetic material has been effective. However, in situations where a smaller vessel diameter is required, the synthetic materials cannot be used due to high rates of stenosis and thrombus formation. One possible solution is to use natural materials like collagen, either modified or combined with a synthetic material, to form a graft that more closely mimics the body's natural function and has low thrombogenicity and low incidence of stenosis. Failure of the autograft is usually due to some form of occlusion that results from luminal narrowing. Damage of the vessel during removal and reimplantation may cause the recruitment of factors or cells that adhere to the autograft wall and decrease the diameter of the lumen. The restricted flow then increases the thrombogenicity, making full occlusion even more likely. Other problems
are preparation and preservation of the autograft, procedures that can result in vessel damage or diminished in vivo performance. Finally, due to increased and/or different mechanical forces, endothelial cells can shrink, diminishing barrier performance, and degrade, also resulting in increased thrombogenicity. In order to reduce thrombus formation anticoagulation drugs are necessary. The use of these drugs often results in undesirable systemic side effects and can be very problematic. Therefore synthetic materials are poor choices for materials for small diameter vascular grafts.

[0198] By incorporating biological materials into a synthetic vascular graft the host response can be modulated to help insure that the graft will not fail. The use of collagen as a material for a synthetic vascular graft is quite promising because it is biodegradable and has good mechanical properties. Since collagen is biodegradable, as the device degrades tissue can grow into the device. This is advantageous because ideally as the collagen implant degrades the newly formed tissue will replace it, which results in a gradual transfer of stress from the implanted device to the newly formed tissue. If a collagen vascular implant material was seeded with endothelial cells so that they coat the lumen, the surface would theoretically be more biocompatible. Recently, endothelial cells have been cultured on the collagen small diameter vascular grafts.

[0199] Therefore by incorporating biodegradable peptides into the collagen vascular implant material, endothelial cells can be seeded on the top of the material to create a luminal surface that is comprised of endothelial cells to more closely mimic the natural biological environment. Migration of endothelial cells on biomaterials is very important for the development of implantable devices. These cell property control the rates of reendothelialization and angiogenesis that are important for the success of the implant.

[0200] Angiogenesis is a complex, multi-stage process by which new blood vessels are formed from pre-existing vasculature. Two critical steps in this process are endothelial cell migration and assembly into new tubules. Over the last decade, diverse arrays of molecular regulators that participate in the process of angiogenesis have been identified. The receptor tyrosine kinases, for example, are one such family of angiogenesis regulators that play a prominent role in endothelial cell assembly and migration.

Rare or Orphan diseases

[0201] Another aspect of the present invention is directed to the use of the peptide as a therapeutic agent for the prophylaxis and/or treatment of the following orphan diseases as well as for the prophylaxis and/or treatment of an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, an infectious disease, or a heart and vascular disease in patients suffering from one or more of the following Rare or Orphan Diseases:

drome, Davies disease, Davis lafer syndrome, De Barsy syndrome, De Haewere-Leroy-Adriaenssens syndrome, De Santis-Cacchiione syndrome, De Smet-Fabry-Fryns syndrome, De Vail disease, De la Chapelpe dysplasia, De morssier syndrome, Deafness-small bowel diverticulitis-neuropathy, Del bambri dillon syndrome, Degos disease, Dejerine-Sottas syndrome, Dekaban-Arima syndrome, Delayed graft function after organ transplantation, Dellemann-Oorthuys syndrome, Dementia associated with a metabolic disease, Dementia associated with a neurodegenerative disease, Dementia associated with an infectious disease, Dementia associated with hepatic and renal failure, Demodicidiosis, Dentritic cell sarcoma, Dentritic cell tumor, Dengue, Dennis Cohen syndrome, Dennis fairhurst Moore syndrome, Dense (delta) granule disease, Dent disease, Dentin dysplasia, Denys-Drash syndrome, Der Kaloustian-Jaridi-Khoury syndrome, Der kaloustian micatosh silver syndrome, Dercum’s disease, Dermatofibrosarcoma protuberans, Dermatologic allergic disease, Dermatostomatitis Stevens Johnson type, Desbuquois syndrome, Desminopathy, Desmo disease, Desmosterolosis, Devic’s disease, Devriendt legius iryns syndrome, Devriendt vandenbergh iryns syndrome, DiGeorge syndrome, Diabetes, Dialysis-related arthropathy, Diaphanopondylodysostosis, Diaphragmatic agenesia, Diaphragmatic muscular atrophy, Diffuse alveolar haemorrhage, Diffuse large B cell lymphoma, Diffuse leiomymatosis-Alport syndrome X-linked, Diffuse neonatal hamangiomatosis, Dihydropyrimidinuria, Dilated cardiomyopathy with ataxia, Dinoco-Salii-Patel syndrome, Dinno sheeter weisskopf syndrome, Diomedi bernardi placti syndrome, Dionisi-Vici-Sabotta-Gambarruda syndrome, Diphtheria, Diprosopiasis, Discoid lupus erythematosus, Discrete fibromuscular subaortic stenosis, Distichiasis-congeni
Mast cell sarcoma, Mastocytosis, Mastroiacovo de rosa satta syndrome, Mathieu de broca bony syndrome, Matsoukas liarios giannika syndrome, Matthew-Wood syndrome, Mature B-cell tumour, Mature T-cell and NK-cell tumour, May-Hegglin thrombocytopenia, Mayer-Rokitansky-Kuster-Hauser syndrome, Mazabraud syndrome, McArdle disease, McCabe’s disease, McCune-Albright syndrome, McDonough syndrome, McDowell syndrome, McGrath syndrome, McKusick-Kaufman syndrome, McLeod syndrome, McPherson-Hall syndrome, Mcclister cran syndrome, Mccallum macadam johnston syndrome, Megillivray syndrom, McInan-Dekaban syndrome, McPherson clemens syndrome, Mreceham winn culler syndrome, Meadows’s syndrome, Meckle like syndrom, Meckel syndrome, Meckel-gruber syndrome, Meconium aspiration syndrome, Medaliea dennis donnai syndrome, Medialinal (thymic) large b-cell lymphoma, Medialinal diffuse large-cell lymphoma with sclerosis, Mediastinal fibrosis, Medrano ronaldan syndrome, Medullar disease, Medullary cystic kidney disease, Medulloblastoma, Megacalyxosis, Megadenodium and/or megacystis, Megaloblastic anaeemia, Megabane-Loiselet syndrome, Mehes syndrome, Mehta-Levis-Patton syndrome, Meier blumberg imahorn syndrome, Meier-Gorlin syndrome, Meige disease, Meinecke pepper syndrome, Meinecke syndrome, Melanoma, Meleda disease, Melhem fahl syndrome, Meliodiosis, Melkerson rosenthal syndrome, Melnick-Needles syndrome, Melorheostosis, Membranoproliferative glomerulonephritis, Membranous glomerulopathy, Menetri’s disease, Menengioma, Meningitis, Menkes syndrome, Mental retardation, Meretoja syndrome, Merkel cell carcinoma (MCC), Merlob gruenbaum reiser syndrome, Mesangial sclerosis, Mesodermic dysplasia, Mesothelioma, Mesulam syndrome, Metabolic intoxication disease, Metabolic liver disease, Metaphysseal dysplasia, Michels syndrome, Mickeyson syndrome, Micro syndrome, Microcephaly, Microcoria, Microcystic infiltrating lymphatic malformation, Microcytic anaeemia, Microphthalmia, Microscopic colitis Microtia, Microvillous inclusion disease, Mid-aortic dysplastic syndrome, Midas syndrome, Middle aortic syndrome, Midline heart, Mietens syndrome, Mievis verellen dumoulin syndrome, Mikati najar sahil syndrome, Mikulicz disease, Mild campomelic dysplasia, Miller syndrome, Miller-Dieker syndrome, Miller-Fisher syndrome (MFS), Mills syndrome, Milroy disease, Minimal change nephrotic syndrome (MCNS), Minkowski-Chauffard disease, Mirhosseini-Holmes-Walton syndrome, Mitral valve prolapse disease, Mitra syndrome, Mixed connective tissue disease, Mixed phenotype acute leukaemia, Mixed serebrosis bone dystrophy, Miyoshi myopathy, Ms syndrome, Moderate and severe traumatic brain injury, Moebius syndrome, Moeran vandenberghie fryns syndrome, Moersch-Woltman syndrome, Moeschler clarren syndrome, Mohr syndrome, Mohr-Traubejaer syndrome, Mollica pavone antener syndrome, Moloney syndrome, Mono syndrome, Monilethrix, Mononen-Karnes-Senac syndrome, Monostotic fibrous dysplasia, Montefiore syndrome, Moore-Federman syndrome, Morav-Mehes syndrome, Morgagni-Stewart-Morel syndrome, Morillo cucci passage syndrome, Morning glory syndrome, Morquio disease, Morris syndrome, Morse ramsley sargent syndrome, Morvan syndrome, Moschcowitz disease, Mounier-Kuhn syndrome, Mousa-Al Din-Al Nassar syndrome, Movement disease, Mowat-Wilson syndrome, Moya-moya disease, Moynahan syndrome, Mpo deficiency, Msbd syndrome, Mseleni joint disease (MJD), Mucha Habermann Disease, Muckle-Wells syndrome, Mucoepithelial dysplasia, Mucoepidermoid, Muco- polysaccharidosis, Mucomyosis, Mucosal pemphigoid, Mucosulphatidosis, Muenke syndrome, Muir-Torre syndrome, Mullerian aplasia, Multicentric Castleman disease (MCD), Multicentric giant lymph node hyperplasia, Multicentric osteosclerosis, Multifocal acquired demyelinating sensory and motor neuropathy, Multifocal pattern dystrophy simulating fundus flavimaculosus, Multiglandular hyperplasia, Multiminicore disease (MmD), Multinodular goiter cystic kidney polyductyly), Multiple carboxylase deficiency, Multiple contracture syndrome, Multiple cutaneous and uterine leiomyomases, Multiple endocrine neoplasia, Multiple episphyal dysplasia, Multiple fibrofolliculoma, Multiple hamartoma syndrome, Multiple keratoacanthoma, Multiple pterygium syndrome, Multiple sclerosis, Multiple sulfatase deficiency, Multiple system atrophy, Multiple ventricular septal defects, Mulwihill-Smith syndrome, MURCS association, Murray-Puretic-Drescher syndrome, Muscular chanelopathy, Muscular dystrophy, Muscular fibrosis multifocal obstructed vessels, Mutchinick syndrome, Myalgia eosinophilia associated with tryptophan, Myasthenia gravis, Myasthenic syndromes, Myectomy, Mycoplasma encephalitis, Mycrosis fungoides, Myelinolastic diffuse sclerosis, Myelinosis centralis diffusa, Myelocerebellar disorder, Myelodysplastic or myeloproliferative disease, Myelofibrosis with myeloid metaplasia, Myeloid sarcoma, Myeloma, Myhre syndrome, Myiasis, Myelocytic dystonia, Myoclonic epilepsy, Myodysplasia, Myofibrillar myopathy, Myoglobinuria, Myopathy and diabetes mellitus, Myopathy, Myopia, Myositis ossificans progressiva, Myotilinopathy, Myotonia congenita, Myotonic disease, Myotubular myopathy, Myxofibrosarcoma, Myxoid liposarcoma, Myxoid malignant fibrous histiocytoma, Myomma with fibrous dysplasia, Möbius syndrome, N syndrome, NACG, NAGS deficiency, NAMe syndrome, NAO syndrome, NARP syndrome, NASH syndrome, NBS, NCL, NCMD, NF1, NFJ syndrome, NIH, NHPP, NISCH syndrome, NOMID syndrome, NPLCA, NSIP, NTD, Naegeli syndrome, Naegeli-Franceschetti-Jadassohn syndrome, Nager syndrome, Naugib syndrome, Nail anomaly, Nail dysplasia, Naito-Oyanagi disease, Nakagawa’s angioblastoma, Nakajo nishimura syndrome, Nakajo syndrome, Nakamura osme syndrome, Nancre-Horan syndrome, Narcolepsy without cataplexy, Narcolepsy-Cataplexy, Nasodigitoacoustic syndrome, Nasopharyngeal cancer, Nasit-Hakola disease, Nathalie syndrome, Navajo brainstem syndrome, Naxos disease, Necrotising hypophysitis, Necrotizing myelitis, Nema-line myopathy, Neonatal Onset Multisystem Inflammatory Disease, Neonatal death immune deficiency, Neonatal hemochromatosis, Neonatal neutropenia, Neonatal respiratory distress syndrome, Nephroblastoma, Nephrogenic fibrosing dermopathy, Nephrogenic systemic fibrosis, Nephrolithiasis, Nephronophthisis-hepatic fibrosis, Nephropathy, Nephrotic syndrome with diffuse mesangial sclerosis, Nephrotic syndrome, Nervous system tumour, Netherton disease, Neu-Laxova syndrome, Neuhauer daily magnelli syndrome, Neuhauer eicher opitz syndrome, Neuhauer’s anomaly, Neural crest tumour, Neuroacanthocytosis, Neuroaxonal dystrophy, Neuroblastoma, Neurocutaneous melanosis, Neurodegeneration due to 3-hydroxyisobutyryl-CoA hydrolase deficiency, Neurodegeneration with brain iron accumulation (NBIA), Neurodegenerative disease, Neuroectodermal syndrome, Neuroepithelio, Neurofibromatosis, Neuroliptonatosis, Neuromuscular junction disease, Neuromyelitis
actinic dermatitis, cutaneous porphyrias, actinic prurigo and solar urticaria), Uremic pruritus, Tricyclic antidepressants poisoning, Traumatic spinal cord injury, Renal cell carcinoma, Superficial bladder cancer, Staphylococcus aureus bacteremia, Spinal cord injury, Spinia bifida, Soft tissue sarcoma, Small cell lung cancer. Sickle cell disease, Severe myoclonic epilepsy in infancy, Severe combined immunodeficiency (SCID), Severe closed traumatic brain injury, Retinopathy of prematurity, Retinitis pigmentosa, Respiratory distress syndrome in premature neonates of less than 32 weeks of gestational age, Recurrent hepatitis C virus induced liver disease in liver transplant recipients, Radiation proctitis, Pseudomonas aeruginosa lung infection in cystic fibrosis, Progressive myoclonic epilepsies, Primary malignant bone tumors, Primary apnoea of premature newborns, Post-transplant lymphoproliferative disorders, Post-neonatal intracerebral haemorrhage, Post transplantation graft dysfunction, Polycythemia vera, Peritumoral oedema derived from brain tumors, Peripheral T-cell lymphoma (nodal, other extranodal and leukemic/ disseminated), Ductus arteriosus in premature neonates of less than 34 weeks of gestational age, Partial deep dermal and full thickness burns, Paroxysmal nocturnal haemoglobinuria, Pancreatic cancer, Painful HIV-associated neuropathy, Ovarian cancer, Osteosarcoma, Orthostatic hypotension in patients with pure autonomic failure, Orthostatic hypotension in patients with multiple system atrophy, Ornithine-transcarbamylase deficiency, Oral mucositis in head and neck cancer patients undergoing radiation therapy, Oesophageal cancer, Non-traumatic osteonecrosis, Non-keto- tic hyperglycaemia, Non-infectious uveitis affecting the posterior segment of the eye, Non-24-hour sleep-wake disorders in blind people with no light perception, Neuroblastoma, Neovascular glioma, Nephritic syndrome, Myelodysplastic syndromes, Myasthenia gravis, Moderate and severe traumatic brain injury, Metachromatic leukodystrophy, Medul- lary thyroid carcinoma, Mastocytosis, Mantle cell lymphoma, Malignant melanoma, Malignant gastrointestinal stromal tumors, Malabsorption due to exocrine pancreatic enzyme insufficiency, Low flow priapism, Lipoprotein lipase deficiency, Ligneous conjunctivitis, Leber’s hereditary optic neuropathy, Leber’s congenital amaurosis, Late onset sepsis in premature infants of less than or equal to 32 weeks gestational age, Juvenile myelomonocytic leukaemia, Japanese encephalitis, Intestinal graft-versus-host disease, Indolent non-Hodgkin’s lymphoma, Inborn errors in primary bile acid synthesis, Hyperphenylalaninemia, Hyperexosinophilic syndrome, Glioma, High-grade dysplasia in Barrett’s oesophagus, Herpes simplex virus stromal keratitis, Hereditary factor XIII deficiency, Hepatocellular carcinoma, Hepatitis B re-infection following liver transplantation, Hepatic veno-occlusive disease, Gram negative bacterial lung infection in cystic fibrosis, Gastric cancer, Gamma sarcoglycanopathy, Follicular lymphoma, Familial adenomatous polyposis, Emphysema secondary to congenital alpha-1 antitrypsin deficiency, Duchenne muscular dystrophy, Diffuse large B cell lymphoma, Diffuse alveolar haemorrhage, Diarrhoea associated with intestinal microsporidial infection, Cutaneous T-cell lymphoma, Cutaneous forms of lupus erythematosus, Cushing’s syndrome secondary to ectopic ACTH secretion, Corneal graft rejection, Congenital venous malformations, Congenital lymphatic malformations, Congenital alpha-1 antitrypsin deficiency, Congenital adrenal hyperplasia, Chronic pain, Cocaine poisoning, Chronic myeloid leukaemia, Chronic lymphocytic leukaemia, Chronic iron over- load requiring chelation therapy, Chronic idiopathic myelofibrosis, Chronic eosinophilic leukaemia and the hyperesinophilic syndrome, Cholangiocarcinoma, Charcot-Marie-Tooth disease type 1A, Cardiogenic shock, Bronchopulmonary dysplasia in premature neonates of less than 30 weeks of gestational age, B-cell chronic lymphocytic leukaemia, Autoimmune uveitis, Atypical Haemolytic Uraemic Syndrome (aHUS) associated with an inherited abnormality of the complement system, Aspiration pneumonia requiring intubation and mechanical ventilation, Aneurysm subarachnoid haemorrhage, Anaplastic thyroid cancer, Anal fistula, Acute sensorineural hearing loss (acute acoustic trauma, sudden deafness and surgery induced acoustic trauma), Acute peripheral arterial occlusion, Acute intermittent porphyria, Active phase of Peyronie’s disease, Acanthamoeba keratitis, A-mannosidosis, 5q spinal muscular atrophy, Cavopulmonary Anastomosis, Ataxic Septal Defects (ASD), Partial Anomalous Pulmonary Venous Return, Persistent Common Atrial Ventricular Canal Endocardial Cushion Defect. Ostium Primum, Single Atrium, Patent Ductus Arteriosus (PDA), Total Anomalous Pulmonary Venous Return, Ventricular Septal Defects (VSD), Pulmonary Valve Stenosis, Pulmonary Artery Stenosis and Stenosis of Pulmonary Artery Branches, Pulmonary Atresia with Intact Ventricular Septum, Congenital Mitral Valve Disease, Aortic Valvar Stenosis and Congenital Aortic Valvar Regurgitation, Supravalvular Aortic Stenosis, Transposition of the Great Arteries, Double Outlet Right Ventricle, Corrected Transposition of the Great Arteries, Truncus Arteriosus, Aorto Pulmonary Window, Tricuspid Atresia, Ebstein Anomaly, Malformations of the Vena Cava, Coarctation of the Aorta, Atria of Aortic Valve, Anomalies of the Aortic Arch, Anomalous Origin of the Right Subclavian Artery with Coarctation of the Aorta, Idiopathic Dilatation of the Pulmonary Artery, Left Pulmonary Artery Arising from Right Pulmonary Artery, Dextrocardia-Situs Inversus Totais, Association of Heart Malformations with Asplenia, Malformations of the Vena Cava, Congenital Coronary Artery Arte rio-Venous Fistula, Abnormal Origin of the Coronary Articies, Aneurysm of the Sinus of Valsalva (Aortic Sinus Aneurysm), Endocardial Fibroelastosis, Idiopathic Hypertrophic Subaortic Stenosis (IHSS), Mitral Valve Prolapse-Barlow’s Syndrome, Hypoplastic Left Heart.

Pharmaceutical Compositions

[0203] Still another aspect of the present invention relates to the use of the peptide according to claim 1 as an active ingredient, together with at least one pharmaceutically acceptable carrier, excipient and/or diluents for the manufacture of a pharmaceutical composition for the treatment and/or prophylaxis of cancer, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, an infectious disease, a lung disease, a heart and vascular disease or a metabolic disease or any other disease disclosed herein.

[0204] Such pharmaceutical compositions comprise the peptide as an active ingredient, together with at least one pharmaceutically acceptable carrier, excipient, binders, disintegrates, glidants, diluents, lubricants, coloring agents, sweetening agents, flavoring agents, preservatives or the like. The pharmaceutical compositions of the present invention can be prepared in a conventional solid or liquid carrier or diluents and a conventional pharmaceutically-made adjuvant at suitable dosage level in a known way.
Preferably the peptide is suitable for intravenous administration or suitable for oral administration or suitable for administration by inhalation.

Administration forms include, for example, pills, tablets, film tablets, coated tablets, capsules, liposomal formulations, micro- and nano-formulations, powders and deposits. Furthermore, the present invention also includes pharmaceutical preparations for parenteral application, including dermal, intradermal, intragastral, intracutan, intravasal, intravenous, intramuscular, intraperitoneal, intrasal, intravaginal, intrabuccal, percutan, rectal, subcutaneous, supravaginal, topical, or transdermal application, which preparations in addition to typical vehicles and/or diluents contain the peptide according to the present invention.

The present invention also includes the mammalian milk, artificial mammalian milk as well as mammalian milk substitutes as a formulation for oral administration of the peptide to newborns, toddlers, and infants, either as pharmaceutical preparations, and/or as dietary food supplements.

The peptide of the invention can also be administered in form of its pharmaceutically active salts. Suitable pharmaceutically active salts comprise acid addition salts and alkali or earth alkali salts. For instance, sodium, potassium, lithium, magnesium or calcium salts can be obtained.

The peptide of the invention forms pharmaceutically acceptable salts with organic and inorganic acids. Examples of suitable acids for such acid addition salt formation are hydrochloric acid, hydrobromic acid, sulfuric acid, phosphoric acid, acetic acid, citric acid, oxalic acid, malonic acid, salicylic acid, p-aminosalicylic acid, maleic acid, succinic acid, ascorbic acid, maleic acid, sulfonic acid, phosphonic acid, perchloric acid, nitric acid, formic acid, propionic acid, gluconic acid, laetic acid, tartaric acid, hydroxymaleic acid, pyruvic acid, phenylacetic acid, benzoic acid, p-amino benzoic acid, p-hydroxy benzoic acid, methanesulfonic acid, ethanesulfonic acid, nitrous acid, hydroxyethanesulfonic acid, ethylenesulfonic acid, p-tolu enesulfonic acid, naphthalenesulfonic acid, sulfamic acid, cam phersulfonic acid, chinit acid, mandelic acid, o-methyl mandelic acid, hydrogen-benzensulfonic acid, picric acid, adipic acid, o-o-tolyluric acid, tartronic acid, a-toluic acid, (o, m, p)-toluic acid, naphthylamine sulfonic acid, and other mineral or carboxylic acids well known to those skilled in the art. The salts are prepared by contacting the free base form with a sufficient amount of the desired acid to produce a salt in the conventional manner.

The pharmaceutical compositions according to the present invention will typically be administered together with suitable carrier materials selected with respect to the intended form of administration, i.e. for oral administration in the form of tablets, capsules (either solid filled, semi-solid filled or liquid filled), powders for constitution, aerosol preparations consist with conventional pharmaceutical practices. Other suitable formulations are gels, elixirs, dispersible granules, syrups, suspensions, creams, lotions, emulsions, suspensions, dispersions, and the like. Suitable dosage forms for sustained release include tablets having layers of varying disintegration rates or controlled release polymeric matrices impregnated with the active components and shaped in tablet form or capsules containing such impregnated or encapsulated polymeric matrices. The pharmaceutical compositions may be comprised of 5 to 95% by weight of the peptide.

As pharmaceutically acceptable carrier, excipient and/or diluents can be used lactose, starch, sucrose, cellulose, magnesium stearate, dicalcium phosphate, calcium sulfate, talc, mannitol, ethyl alcohol (liquid filled capsules).

Suitable binders include starch, gelatin, natural sugars, corn sweeteners, and synthetic gums such as acacia, sodium alginate, carboxymethyl-cellulose, polyethylene glycol and waxes. Among the lubricants that may be mentioned for use in these dosage forms, boric acid, sodium benzene, sodium acetate, sodium chloride, and the like. Disintegrants include starch, methylcellulose, guar gum and the like. Sweetening and flavoring agents and preservatives may also be included where appropriate. Some of the terms noted above, namely disintegrants, diluents, lubricants, binders and the like, are discussed in more detail below.

Additionally, the compositions of the present invention may be formulated in sustained release form to provide the rate controlled release of any one or more of the components or active ingredients to optimize the therapeutic effects. Suitable dosage forms for sustained release include layered tablets containing layers of varying disintegration rates or controlled release polymeric matrices impregnated with the active components and shaped in tablet form or capsules containing such impregnated or encapsulated polymeric matrices.

Aerosol preparations suitable for inhalation may include solutions and solids in powder form, which may be in combination with a pharmaceutically acceptable carrier such as inert compressed gas, e.g. nitrogen.

For preparing suppositories, a low melting wax such as a mixture of fatty acid glycerides such as cocoa butter is first melted, and the active ingredient is dispersed homogeneously therein by stirring or similar mixing. The molten homogeneous mixture is then poured into convenient sized molds, allowed to cool and thereby solidify.

Also included are solid form preparations which are intended to be converted, shortly before use, to liquid form preparations for either oral or parenteral administration. Such liquid forms include solutions, suspensions and emulsions.

The peptide of the present invention may also be deliverable transdermally. The transdermal compositions may take the form of creams, lotions, aerosols and/or emulsions and can be included in a transdermal patch of the matrix or reservoir type as are conventional in the art for this purpose.

The transdermal formulation of the peptide of the invention is understood to increase the bioavailability of said peptide into the circulating blood. One problem in the administration of peptides is the loss of bioactivity due to the formation of insolubles in aqueous environments or due to degradation. Therefore stabilization of peptides for maintaining their fluidity and maintaining their biological activity upon administration to the patients in need thereof needs to be achieved.

Prior efforts to provide active agents for medication include incorporating the medication in a polymeric matrix whereby the active ingredient is released into the systemic circulation. Known sustained-release delivery means of active agents are disclosed, for example, in U.S. Pat. No. 4,325,988, U.S. Pat. No. 4,188,373, U.S. Pat. No. 4,100,271, U.S. Pat. No. 447,471, U.S. Pat. No. 4,474,752, U.S. Pat. No. 4,474,753, or U.S. Pat. No. 4,478,822 relating to polymeric pharmaceutical vehicles for delivery of pharmaceutically active chemical materials to mucous membranes. The pharmaceutical carriers are aqueous solutions of certain polyoxy-
ethylene-polyoxypropylene condensates. These polymeric pharmaceutical vehicles are described as providing for increased drug absorption by the mucous membrane and prolonged drug action by a factor of two or more. The subseries are block copolymers of polyoxypropylene and polyoxyethylene used for stabilization of drugs such as insulin.

Aqueous solutions of telomer-polyethylene-polyoxypropylene block copolymers (poloxomers) are useful as stabilizers for the peptide. Aside from serving as a stabilizer for the peptide, poloxomers provide excellent vehicles for the delivery of the drug peptide, and they are physiologically acceptable. Poloxamers, also known by the trade name Pluronics (e.g., Pluronic F127, Pluronic P85, Pluronic F68) have surfactant properties that make them useful in industrial applications. Among other things, they can be used to increase the water solubility of hydrophobic oily substances or otherwise increase the miscibility of two substances with different hydrophobicities. For this reason, these polymers are commonly used in industrial applications, cosmetics, and pharmaceuticals. They have also been used as model systems for drug delivery applications. In situ gelation of pharmaceutical compositions based on poloxamer that are biologically triggered are known in the art (e.g., U.S. Pat. No. 5,256,396), describing compositions containing poloxamer 407 and water at specified concentrations.

The term capsule refers to a special container or enclosure made of methyl cellulose, polyvinyl alcohols, or denatured gelatins or starch for holding or containing compositions comprising the active ingredients. Hard shell capsules are typically made of blends of relatively high gel strength bone and pork skin gelatins. The capsule itself may contain small amounts of dyes, opaquing agents, plasticizers and preservatives.

Tablet means compressed or molded solid dosage form containing the active ingredients with suitable diluents. The tablet can be prepared by compression of mixtures or granulations obtained by wet granulation, dry granulation or by compaction well known to a person skilled in the art.

Oral gels refers to the active ingredients dispersed or solubilized in a hydrophilic semi-solid matrix.

Powders for constitution refer to powder blends containing the active ingredients and suitable diluents which can be suspended in water or juices. One example for such an oral administration form for newborns, toddlers and/or infants is a human breast milk substitute which is produced from milk powder and milk whey powder, optionally and partially substituted with lactose.

Human breast milk is a complex fluid, rich in nutrients and in non-nutritional bioactive components. It contains all of the nutrients needed by the newborn baby. These include the metabolic components (fat, protein, and carbohydrates), water, and the raw materials for tissue growth and development, such as fatty acids, amino acids, minerals, vitamins, and trace elements.

More than 98% of the fat in is in the form of triglycerides. Oleic acid and palmitic acid are the most abundant fatty acids in breastmilk triglycerides, with comparatively high proportions of the essential fatty acids, and linolenic acid, followed by long-chain polyunsaturated fatty acids, such as arachidonic acid and docosahexaenoic acid. These long-chain fatty acids are constituents of brain and neural tissue and are needed in early life for mental and visual development. The lipid component of breast milk is the transport vehicle for fat-soluble micronutrients such as prostaglandins and vitamins A, D, E, and K.

Proteins account for approximately 75% of the nitrogen-containing compounds in breast milk. Non-protein nitrogen substances include urea, nucleotides, peptides, free amino acids, and DNA. The proteins of breast milk can be divided into two categories: micellar caseins and aqueous whey proteins, present in the ratio of about 40:60. Casein forms micelles of relatively small volume and produces a soft, flocculent curd in the infant’s stomach. The major whey proteins are lactalbumin, lactoferrin, secretory IgA, and serum albumin, with a large number of other proteins and peptides present in smaller amounts.

The principal carbohydrate is lactose, a disaccharide produced in the mammary epithelial cell from glucose by a reaction involving lactalbumin. In addition to the nutritional components, breast milk contains a wealth of bioactive components that have beneficial non-nutritional functions. These include a wide range of specific and non-specific antimicrobial factors; cytokines and anti-inflammatory substances; and hormones, growth modulators, and digestive enzymes (Table 1), many of which have multiple activities. These components may be of particular importance for young infants because of the immaturity of the host defense and digestive systems early in life.

<table>
<thead>
<tr>
<th>TABLE 4</th>
<th>Examples of the non-nutritional components of breast milk</th>
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</thead>
<tbody>
<tr>
<td>Antimicrobial factors</td>
<td>secretory IgA, IgM, IgG</td>
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<tr>
<td></td>
<td>lactoferrin</td>
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<tr>
<td></td>
<td>lysozyme</td>
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<tr>
<td></td>
<td>complement C3</td>
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<td></td>
<td>leucocytes</td>
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<td></td>
<td>bifidus factor</td>
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<tr>
<td></td>
<td>lipids and fatty acids</td>
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<td></td>
<td>antiviral mucins, GAGs</td>
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<td></td>
<td>oligosaccharides</td>
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<td></td>
<td>Cytokines and anti-inflammatory factors</td>
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<td></td>
<td>tumor necrosis factor</td>
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<tr>
<td></td>
<td>interferon</td>
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<td></td>
<td>prostaglandins</td>
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<td></td>
<td>antichymotrypsin</td>
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<tr>
<td></td>
<td>antifibrin</td>
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<tr>
<td></td>
<td>platelet-activating factor</td>
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<tr>
<td>Hormones</td>
<td>feedback inhibitor of lactation (FIL)</td>
</tr>
<tr>
<td></td>
<td>insulin</td>
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<tr>
<td></td>
<td>prolactin</td>
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<td></td>
<td>thyroid hormones</td>
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<td></td>
<td>corticosteroids</td>
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<td></td>
<td>ACTH</td>
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<td></td>
<td>oxytocin</td>
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<td></td>
<td>calcitonin</td>
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<td></td>
<td>parathyroid hormone</td>
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<tr>
<td></td>
<td>erythropoietin</td>
</tr>
<tr>
<td>Growth factors</td>
<td>feedback inhibitor of lactation (FIL)</td>
</tr>
<tr>
<td></td>
<td>nerve (NGF)</td>
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<tr>
<td></td>
<td>insulin-like (IGF)</td>
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<tr>
<td></td>
<td>transforming (TGF)</td>
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<td></td>
<td>taurine</td>
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<td></td>
<td>polyamines</td>
</tr>
</tbody>
</table>
Besides breast milk, infant formula is the only other infant milk which the medical community considers nutritionally acceptable for infants under the age of one year. Cow's milk is not recommended because of its high protein and electrolyte (salt) content which may harm infant's immature kidneys. The nutrient content of infant formula should comprise: Protein, Fat, Linoleic acid, Vitamins: A, C, D, E, K, thiamin (B1), riboflavin (B2), B6, B12, Niacin, Folic acid, Pantothenic acid, Calcium, Metals: magnesium, iron, zinc, manganese, copper; Phosphorus, iodine, Sodium chloride, Potassium chloride. In addition, formulas not made with cow's milk must include biotin, choline, and inositol. Hypoallergenic formulas reduce the likelihood of certain medical complications in babies with specific health problems. Baby formula can be synthesized from raw amino acids. This kind of formula is sometimes referred to as elemental infant formula or as medical food because of its specialized nature. Powder blends containing the active ingredients and suitable diluents which can be suspended in water or juices can be produced by spray drying.

Spray drying has been found to be the most suitable process for removing the last part of the water, since spray drying can convert milk concentrate into a powder while still keeping the valuable properties of the milk. The principle of all spray dryers is to transform the concentrate into many small droplets which are then exposed to a fast current of hot air. Because of the very large surface area of the droplets, the water evaporates almost instantaneously and the droplets are transformed into powder particles.

Powdered milk is a powder made from dried milk solids. Powdered milk has a far longer shelf life than liquid milk and does not need to be refrigerated due to its low moisture content.

Instant milk powder is produced by partially rehydrating the dried milk powder particles causing them to become sticky and agglomerate. The water is then removed by drying resulting in an increased amount of air incorporated between the powder particles.

Milk powder manufacture is a process carried out on a large scale. It involves the gentle removal of water, while retaining all the desirable natural properties of the milk like colour, flavour, solubility, nutritional value.

Milk powder process includes spray drying, fluid bed processing, extraction, evaporation and freeze drying. Other processes are freeze concentration, filtration, and homogenisation.

The artificial mother milk formulations or mother milk substitutes of the present invention are preferably prepared by adding to a mother milk formulation including commercially available mother milk formulations especially in power form the peptide of the present invention. The peptide is preferably added in an amount of 3-100 µg peptide or per 100 ml (commercially available) mother milk formulation, more preferably in an amount of 5-70 µg/100 ml and most preferably in an amount of 10-40 µg/100 ml mother milk formulation.

Suitable diluents are substances that usually make up the major portion of the composition or dosage form. Suitable diluents include sugars such as lactose, sucrose, mannitol and sorbitol, starches derived from wheat, corn rice and potato, and celluloses such as microcrystalline cellulose. The amount of diluents in the composition can range from about 5 to about 95% by weight of the total composition, preferably from about 25 to about 75%, more preferably from about 30 to about 60% by weight, and most preferably from about 40 to 50% by weight.

The term disintegrants refers to materials added to the composition to help it break apart (disintegrate) and release the medicaments. Suitable disintegrants include stearates, cold water soluble modified stearates such as sodium carboxymethyl starch, natural and synthetic gums such as locust bean, karaya, guar, tragacanth and agar, cellulose derivatives such as methylcellulose and sodium carboxymethylcellulose, microcrystalline celluloses and cross-linked microcrystalline celluloses such as sodium croscarmellose, alginates such as alginic acid and sodium alginate, clays such as bentonites, and effervescent mixtures. The amount of disintegrant in the composition can range from about 1 to about 40% by weight of the composition, preferably 2 to about 30% by weight of the composition, more preferably from about 3 to 20% by weight of the composition, and most preferably from about 5 to about 10% by weight.

Binders characterize substances that bind or "glue" powders together and make them cohesive by forming granules, thus serving as the "adhesive" in the formulation. Binders add cohesive strength already available in the diluents or bulking agent. Suitable binders include sugars such as sucrose, starches derived from wheat, corn rice and potato; natural gums such as acacia, gelatin and tragacanth; derivatives of seaweed such as alginic acid, sodium alginate and ammonium calcium alginate; cellulose materials such as methylcellulose and sodium carboxymethylcellulose and hydroxypropyl-methylcellulose; polyvinylpyrrolidone; and inorganics such as magnesium aluminum silicate. The amount of binder in the composition can range from about 1 to 30% by weight of the composition, preferably from about 2 to about 20% by weight of the composition, more preferably from about 3 to about 10% by weight, even more preferably from about 3 to about 6% by weight.

Lubricant refers to a substance added to the dosage form to enable the tablet, granules, etc. after it has been compressed, to release from the mold or die by reducing friction or wear. Suitable lubricants include metallic stearates such as magnesium stearate, calcium stearate or potassium stearate; stearic acid; high melting point waxes; and water soluble lubricants such as sodium chloride, sodium benzoate, sodium acetate, sodium oleate, polyethylene glycols and d'L-leucine. Lubricants are usually added at the very last step before compression, since they must be present on the surfaces of the granules and in between them and the parts of the
tablet press. The amount of lubricant in the composition can range from about 0.05 to about 15% by weight of the composition, preferably 0.2 to about 5% by weight of the composition, more preferably from about 0.3 to about 3%, and most preferably from about 0.3 to about 1.5% by weight of the composition.

[0240] Gildents are materials that prevent caking and improve the flow characteristics of granulations, so that flow is smooth and uniform. Suitable gildents include silicon dioxide and talc. The amount of gildent in the composition can range from about 0.01 to 10% by weight of the composition, preferably 0.1% to about 7% by weight of the composition, more preferably from about 0.2 to 5% by weight, and most preferably from about 0.5 to about 2% by weight.

[0241] Coloring agents are excipients that provide coloration to the composition or the dosage form. Such excipients can include food grade dyes and food grade dyes adsorbed onto a suitable adsorbent such as clay or aluminum oxide. The amount of the coloring agent can vary from about 0.01 to 10% by weight of the composition, preferably from about 0.05 to 6% by weight, more preferably from about 0.1 to about 4% by weight of the composition, and most preferably from about 0.1 to about 1%.

[0242] The peptide of the invention can be used to form multiparticles, discrete particles, well known dosage forms, whose totality represents the intended therapeutically useful dose of a drug. When taken orally, multiparticles generally disperse freely in the gastrointestinal tract, and maximize absorption. A specific example is described in U.S. Pat. No. 6,068,859, disclosing multiparticles that provide controlled release of azithromycin. Another advantage of the multiparticles is the improved stability of the drug. The poloxamer component of the multiparticulate is very inert, thus minimizing degradation of the drug.

[0243] However, formulation problems result from the melt-congeal process often used to form multiparticles. The multiparticles are preferably formed into round beads or spheres. Some carriers, when melted and then solidified, do not form round beads but may solidify into rods, strings, or other non-spherical shapes. The result is very irregularly shaped multiparticles that are difficult to process into dosage forms. This problem is solved by e.g. WO 2007104173 where the particles consist of a poloxamer, a resin, and/or a tocopherol, creating together with the medicament (e.g. insulin) micelles. Micelle formation is essential for the absorption of many nutrients within the human body. Bile salts formed in the liver and secreted by the gall bladder allow micelles of fatty acids to form. This allows the absorption of complicated lipids and lipid soluble vitamins within the micelle by the small intestine. Micelles are approximately spherical in shape. Preferably, peptide of the invention are formulated with a poloxamer and a resin to form micelles suitable for oral administration to patients in need of the medicament.

[0244] Liquid form preparations include solutions, suspensions and emulsions. As an example may be mentioned water or water-propylene glycol solutions for parenteral injections or addition of sweeteners and opacifiers for oral solutions, suspensions and emulsions. Liquid form preparations may also include solutions for intranasal administration.

[0245] Other preferred pharmaceutical compositions are buffered solutions. The term buffer, buffer system, buffer solution and buffered solution, when used with reference to hydrogen ion concentration or pH, refers to the ability of a system, particularly an aqueous solution, to resist a change of pH on adding acid or alkali, or on dilution with a solvent. Preferred buffer systems can be selected from the group consisting of formate (pKa=3.75), lactate (pKa=3.86), benzoic acid (pKa=4.2), oxalate (pKa=4.29), fumarate (pKa=4.38), aniline (pKa=4.63), acetate buffer (pKa=4.76), citrate buffer (pKa2=4.76, pKa3=6.4), glutamate buffer (pKa=4.3), phosphate buffer (pKa=7.20), succinate (pKa=4.93, pKa2=5.62), pyridine (pKa=5.23), phthalate (pKa=5.41), histidine (pKa=6.04), MES (2-(N-morpholino)ethanesulfonic acid; pKa=6.15), maleic acid (pKa=6.26), cacodylate (dimethylsulfate, pKa=6.27), carboxylic acid (pKa=6.35), ADA (N-2-acetamidoiminodiacetic acid (pKa=6.62), PIPES (4-piperazinebisis-ethanesulfonic acid; BIS-TRIS-propane, 1,3-bis[tris(hydroxymethyl)methylamino]-propane), pKa=6.80), ethylenediamine (pKa=6.85), ACES 2-[2-amino-2-oxoethyl] amino-ethanesulfonic acid; pKa=6.9), imidazole (pKa=6.95), MOPS (3-(N-morphin)-propanesulfonic acid; pKa=7.20), diethylmalonic acid (pKa=7.2), TES (2-[tris(hydroxymethyl)methyl]amino ethanesulfonic acid; pKa=7.50) and HEPES (N-2-hydroxyethylpiperazin-N'-2-ethanesulfonic acid; pKa=7.55) buffers or other buffers having a pKa between 3.8 to 7.7.

[0246] Preferred is the group of carboxylic acid buffers such as acetate and carboxylic diacid buffers such as fumarate, taurine and phthalate and carboxylic tricarboxylic buffers such as citrate. Another group of preferred buffers is represented by inorganic buffers such as sulfate, borate, carbonate, oxalate, calcium hydroxide and phosphate buffers. Another group of preferred buffers are nitrogen containing buffers such as imidazole, diethylenediamine, and piperezine.

[0247] Also preferred are sulfonic acid buffers such as TES, HEPES, ACES, PIPES, [[2-hydroxy-1,1-bis(hydroxymethyl) amino]-1-propanesulfonic acid (TAPS), 4-[2-hydroxyethyl]piperazin-1-propanesulfonic acid (EPAPS), 4-Morpholinepropanesulfonic acid (MOPS) and N,N-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid (BES).

[0248] Another group of preferred buffers are glycine buffers such as glycine, glycyl-glycine, glycyl-glycyl-glycine, N,N,N,N,N,N-trimethyllysine, 3-methylhistidine, 5-hydroxylysine, O-phosphoserine, y-carboxyglutamate, E-N-acetyllysine, o-N-methylarginine, citrulline, ornithine and derivatives thereof.

<table>
<thead>
<tr>
<th>pH range</th>
<th>pKa 25°C C.</th>
<th>Buffer</th>
</tr>
</thead>
<tbody>
<tr>
<td>2.7-4.2</td>
<td>3.40</td>
<td>malate (PK1)</td>
</tr>
<tr>
<td>3.0-4.5</td>
<td>3.75</td>
<td>formate</td>
</tr>
<tr>
<td>3.0-6.2</td>
<td>4.76</td>
<td>citrate (PK2)</td>
</tr>
<tr>
<td>3.2-5.2</td>
<td>4.21</td>
<td>succinate (PK1)</td>
</tr>
<tr>
<td>3.6-5.6</td>
<td>4.76</td>
<td>acetate</td>
</tr>
<tr>
<td>3.8-5.6</td>
<td>4.87</td>
<td>propionate</td>
</tr>
<tr>
<td>4.0-6.0</td>
<td>5.13</td>
<td>malate (PK2)</td>
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<td>4.9-5.9</td>
<td>5.23</td>
<td>pyridine</td>
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<td>5.0-6.0</td>
<td>5.33</td>
<td>piperazine (PK1)</td>
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<td>5.0-7.4</td>
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<td>5.5-6.5</td>
<td>5.64</td>
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TABLE 5

Also preferred are the following buffers:
<table>
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<th>effective pH range</th>
<th>pKa 25°C.</th>
<th>buffer</th>
</tr>
</thead>
<tbody>
<tr>
<td>5.5-6.7</td>
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<td>6.40</td>
<td>citrate (pK3)</td>
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<td>5.5-7.2</td>
<td>6.24</td>
<td>maleate (pK2)</td>
</tr>
<tr>
<td>5.5-7.4</td>
<td>1.70, 6.04, 9.09</td>
<td>histidine</td>
</tr>
<tr>
<td>5.8-7.2</td>
<td>6.46</td>
<td>bis-tris</td>
</tr>
<tr>
<td>5.8-8.0</td>
<td>7.20</td>
<td>phosphate (pK2)</td>
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<tr>
<td>6.0-12.0</td>
<td>9.50</td>
<td>ethanolamine</td>
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<tr>
<td>6.0-7.2</td>
<td>6.59</td>
<td>ADA</td>
</tr>
<tr>
<td>6.0-8.0</td>
<td>6.35</td>
<td>carbonate (pK1)</td>
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<td>6.1-7.5</td>
<td>6.78</td>
<td>ACES</td>
</tr>
<tr>
<td>6.1-7.5</td>
<td>6.76</td>
<td>PIPES</td>
</tr>
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<td>6.2-7.6</td>
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<td>MOPS</td>
</tr>
<tr>
<td>6.2-7.8</td>
<td>6.95</td>
<td>imidazole</td>
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<tr>
<td>6.3-9.5</td>
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<td>BIS-TRIS propane</td>
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<td>6.4-7.8</td>
<td>7.09</td>
<td>BES</td>
</tr>
<tr>
<td>6.5-7.9</td>
<td>7.14</td>
<td>MOPS</td>
</tr>
<tr>
<td>6.6-8.2</td>
<td>7.48</td>
<td>HEPES</td>
</tr>
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<td>6.8-8.2</td>
<td>7.40</td>
<td>TES</td>
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<tr>
<td>6.9-8.5</td>
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<td>7.52</td>
<td>DIPS</td>
</tr>
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<td>TAPSO</td>
</tr>
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<td>7.0-8.3</td>
<td>7.76</td>
<td>triethanolamine (TEA)</td>
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<td>0.91, 2.10, 6.70, 9.32</td>
<td>pyrophosphate</td>
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<td>7.1-8.5</td>
<td>2.85</td>
<td>HUPPSO</td>
</tr>
<tr>
<td>7.2-8.5</td>
<td>7.78</td>
<td>POPSO</td>
</tr>
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</table>

**TABLE 5—continued**

Also preferred are the following buffers:

<table>
<thead>
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<th>effective pH range</th>
<th>pKa 25°C.</th>
<th>buffer</th>
</tr>
</thead>
<tbody>
<tr>
<td>6.0-8.0</td>
<td>6.35</td>
<td>carbonate (pK1)</td>
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<tr>
<td>6.1-7.5</td>
<td>6.78</td>
<td>ACES</td>
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<tr>
<td>6.1-7.5</td>
<td>6.76</td>
<td>PIPES</td>
</tr>
<tr>
<td>6.2-7.6</td>
<td>6.87</td>
<td>MOPS</td>
</tr>
<tr>
<td>6.2-7.8</td>
<td>6.95</td>
<td>imidazole</td>
</tr>
<tr>
<td>6.3-9.5</td>
<td>6.80, 9.00</td>
<td>BIS-TRIS propane</td>
</tr>
<tr>
<td>6.4-7.8</td>
<td>7.09</td>
<td>BES</td>
</tr>
<tr>
<td>6.5-7.9</td>
<td>7.14</td>
<td>MOPS</td>
</tr>
<tr>
<td>6.6-8.2</td>
<td>7.48</td>
<td>HEPES</td>
</tr>
<tr>
<td>6.8-8.2</td>
<td>7.40</td>
<td>TES</td>
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<tr>
<td>6.9-8.5</td>
<td>7.60</td>
<td>MOBS</td>
</tr>
<tr>
<td>7.0-8.2</td>
<td>7.52</td>
<td>DIPS</td>
</tr>
<tr>
<td>7.0-8.2</td>
<td>7.61</td>
<td>TAPSO</td>
</tr>
<tr>
<td>7.0-8.3</td>
<td>7.76</td>
<td>triethanolamine (TEA)</td>
</tr>
<tr>
<td>7.0-9.0</td>
<td>0.91, 2.10, 6.70, 9.32</td>
<td>pyrophosphate</td>
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<tr>
<td>7.1-8.5</td>
<td>2.85</td>
<td>HUPPSO</td>
</tr>
<tr>
<td>7.2-8.5</td>
<td>7.78</td>
<td>POPSO</td>
</tr>
</tbody>
</table>

**[0250]** Preferred are the buffers having an effective pH range of from 2.7 to 8.5, and more preferably of from 3.8 to 7.7. The effective pH range for each buffer can be defined as pKa−1 to pKa+1, where Ka is the ionization constant for the weak acid in the buffer and pKa=−log K.

**[0251]** Most preferred are buffers suitable for pharmaceutical use e.g. buffers suitable for administration to a patient such as acetate, carbonate, citrate, fumarate, glutamate, lactate, phosphate, phthalate, and succinate buffers. Particularly preferred are citrate buffer, carbonate buffer, pyrophosphate buffer, for use in the present invention. The term “carboxylic acid buffers” as used herein shall refer to carboxylic mono acid buffers and carboxylic diacid buffers as well as carboxylic triacid buffers.

**[0252]** Of course also combinations of buffers, especially of the buffers mentioned herein are useful for the present invention.

**[0253]** Some suitable pharmaceutical buffers are a citrate buffer (preferably at a final formulation concentration of from about 20 to 200 mM, more preferably at a final concentration of from about 30 to 120 mM) or an acetate buffer (preferably at a final concentration of from about 20 to 200 mM) or a phosphate buffer (preferably at a final formulation concentration of about 20 to 200 mM).

**[0254]** Techniques for the formulation and administration of the peptide of the present invention may be found in “Remington’s Pharmaceutical Sciences” Mack Publishing Co., Easton Pa. A suitable composition comprising the peptide mentioned herein may be a solution of the peptide in a suitable liquid pharmaceutical carrier or any other formulation such as tablets, pills, film tablets, coated tablets, dragees, capsules, powders and deposits, gels, syrups, slurries, suspensions, emulsions, and the like.

**[0255]** A particularly preferred pharmaceutical composition is a lyophilised (freeze-dried) preparation (lyophilisate) suitable for administration by inhalation or for intravenous administration. To prepare the preferred lyophilised preparation the peptide of the invention are solubilised in a 4 to 5% (w/v) mannitol solution and the solution is then lyophilised. The mannitol solution can also be prepared in a suitable buffer solution as described above.

**[0256]** Further examples of suitable cryo-/lyoprotectants (otherwise referred to as bulking agents or stabilizers) include thiol-free albumin, immunoglobulins, polyalkyleneoxides (e.g. PEG, polypropylene glycols), trehalose, glucose, sucrose, sorbitol, dextran, maltose, raffinose, stachyose and other saccharides (cf. for instance WO 97/29782), while mannitol is used preferably. These can be used in conventional amounts in conventional lyophilization techniques. Methods of lyophilisation are well known in the art of preparing pharmaceutical formulations.

**[0257]** For administration by inhalation the particle diameter of the lyophilised preparation is preferably between 2 and 5 μm, more preferably between 3 to 4 μm. The lyophilised preparation is particularly suitable for administration using an inhalator, for example the OPTINEB® or VENTA-NEB® inhalator (NEBU-TEC, Elseneif, Germany). The lyophilised product can be rehydrated in sterile distilled water or any other suitable liquid for inhalation administration.

**[0258]** Alternately for intravenous administration the lyophilised product can be rehydrated in sterile distilled water or any other suitable liquid for intravenous administration.

**[0259]** After rehydration for administration in sterile distilled water or another suitable liquid the lyophilised preparation should have the approximate physiological osmolality of the target tissue for the rehydrated peptide preparation i.e. blood for intravenous administration or lung tissue for inhalation administration. Thus it is preferred that the rehydrated formulation is substantially isotonic.

**[0260]** The preferred dosage concentration for either intravenous, oral, or inhalation administration is between 100 to 2000 μmole/ml, and more preferably is between 200 to 800 μmole/ml. These are also the preferred ranges of the peptide in the mother milk substitute or artificial mother milk formulation or the pharmaceutical compositions disclosed herein.

Dietary Supplement

**[0261]** Still another aspect of the present invention relates to the use of disclosed peptides as a dietary supplement. That dietary supplement is preferably for oral administration and especially but not limited to administration to newborns, toddlers, and/or infants. A dietary supplement is intended to supplement the diet. The “dietary ingredients” in these products may in addition include: vitamins, minerals, herbs or other botanicals, amino acids, and substances such as enzymes, organ tissues, glandulars, and metabolites. Dietary supplements may be manufactured in forms such as tablets, capsules, softgels, gelcaps, liquids, or powders.

Method of Treatment

**[0262]** Another aspect of the present invention relates to a method of prophylaxis and/or treatment of cancer, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, an infectious disease, a lung disease, a heart and vascular disease or a metabolic disease or any other disease disclosed herein comprising administering to a patient in need thereof a pharmaceutical composition.
comprising the peptide according to the present invention in a therapeutically effective amount effective to treat the aforementioned disease.

Accordingly, the terms “prophylaxis” or “treatment” includes the administration of the peptide of the present invention to prevent, inhibit, or arrest the symptoms of an infectious disease, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, or a heart and vascular disease. In some instances, treatment with the peptide of the present invention will be done in combination with other protective compounds to prevent, inhibit, or arrest the symptoms of an infectious disease, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, or a heart and vascular disease.

The term “active agent” or “therapeutic agent” as used herein refers to an agent that can prevent, inhibit, or arrest the symptoms and/or progression of an infectious, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, or a heart and vascular disease or any other disease disclosed herein.

The term “therapeutic effect” as used herein refers to the effective provision of protection effects to prevent, inhibit, or arrest the symptoms and/or progression of an infectious, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, or a heart and vascular disease.

The term “a therapeutically effective amount” as used herein means a sufficient amount of the peptide of the invention to produce a therapeutic effect, as defined above, in a subject or patient in need of treatment.

The terms “subject” or “patient” are used herein mean any mammal, including but not limited to human beings, including a human patient or subject to which the compositions of the invention can be administered. The term mammals include human patients and non-human primates, as well as experimental animals such as rabbits, rats, and mice, and other animals.

The peptide of the present invention can be used for the prophylaxis and/or treatment of cancer, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, an infectious disease, a lung disease, a heart and vascular disease or a metabolic disease or any other disease mentioned herein in combination administration with another therapeutic compound. As used herein the term “combination administration” of a compound, therapeutic agent or known drug with the peptide of the present invention means administration of the drug and the peptide at such time that both the known drug and the peptide will have a therapeutic effect. In some cases this therapeutic effect will be synergistic. Such concomitant administration can involve concurrent (i.e. at the same time), prior, or subsequent administration of the drug with respect to the administration of the peptide of the present invention. A person of ordinary skill in the art would have no difficulty determining the appropriate timing, sequence and dosages of administration for particular drugs and peptide of the present invention.

Definition of Peptide Activity

A peptide is deemed to have therapeutic activity if it demonstrated any one of the following activities listed in a) to g).

a) The peptide could inhibit the activity of an over active biological pathway.
b) The peptide could inhibit the production of an over produced biological molecule.
c) The peptide could inhibit the activity of an over produced biological molecule.
d) The peptide could increase the activity of an under active biological pathway.
e) The peptide could increase the production of an under produced biological molecule.
f) The peptide could mimic the activity of an under produced biological molecule.
g) The peptide could prevent, inhibit, or arrest the symptoms and/or progression of cancer, an infectious disease, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, or a heart and vascular disease or any other disease disclosed herein.

As used herein “inhibition” is defined as a reduction of the activity or production of a biological pathway or molecule activity of between 10 to 100%. More preferably the reduction of the activity or production of a biological pathway or molecule activity is between 25 to 100%. Even more preferably the reduction of the activity or production of a biological pathway or molecule activity is between 50 to 100%.

As used herein “increase” is defined as an increase of the activity or production of a biological pathway or molecule of between 10 to 100%. More preferably the increase of the activity or production of a biological pathway or molecule activity is between 25 to 100%. Even more preferably the increase of the activity or production of a biological pathway or molecule activity is between 50 to 100%.

As used herein “mimic” is defined as an increase in the activity of a biological pathway dependent on the under produced biological molecule of between 10 to 100%. More preferably the increase of the activity of the biological pathway is between 25 to 100%. Even more preferably the increase of the activity the biological pathway is between 50 to 100%.

Pepptides

The following of the invention peptide was for tested for the activity as a therapeutic agent for the prophylaxis and/or treatment of cancer, an infectious disease, an autoimmune disease, a fibrotic disease, an inflammatory disease, a neurodegenerative disease, or a heart and vascular disease:

Peptide having the amino acid sequence:

H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-L-threoninol (Octreotide).

Furthermore the present invention relates to the use of the above-mentioned peptide as pharmacologically active agents in medicine, i.e. as medicament. Advantage of the peptide of the invention is that the peptide is less toxic in comparison to the commonly used drugs for the certain indications mentioned herein and that the peptide have less side effects, can be used for a long term treatment of certain diseases and can be easily administered. Moreover the peptide are selective for certain targets and under physiological conditions no toxic or noxious degradation products are formed.

As used herein, the term “peptide(s) or “peptide(s) of the invention” shall also refer to salts, deprotected form, acetylated form of the peptide, deacetylated form of the peptide, enantiomers, diastereomers, racemates, prodrugs and hydrates of the above-mentioned peptide. Diastereomers of the peptide are obtained when the stereochemical or chiral
center of one or more amino acids is changed. The enantiomer
has the opposite stereochemistry at all chiral centers.

[0278] The term “prodrug” refers to any precursor com-
 pound which is able to generate or to release the above-
 mentioned peptide under physiological conditions. Such pro-
 drugs, i.e. such precursor molecules are for instance larger
 peptides which are selectively cleaved in order to form the
 peptide of the invention. Further prodrugs are protected
 amino acids having especially protecting groups at the car-
 boxylic acid and/or amino group.

[0279] Suitable protecting groups for amino groups are the
 benzoyloxy carbonyl, t-butoxy carbonyl (BOC), formyl, and
 acetyl or acyl group. Suitable protecting groups for the
 carboxylic acid group are esters such as benzyl esters or t-buty-
 l esters.

[0280] The present invention also includes the above pep-
 tidies having amino acid substitutions, deletions, additions,
 the substitutions and additions including the standard D and L
 amino acids and modified amino acids such as for example
 amidated and acetylated amino acids, wherein the therapeutic
 activity of the base peptide sequence as shown above is main-
 tained.

[0281] In the listed peptide sequences “Ac” indicates an
 acetylated residue and “NH,” indicates an amidated residue.
 “cyclo” indicates a cyclic peptide, and “D” indicates a D
 optical isomer. Decasetylated amino or NH-group refers to the
 free amino (—HH₂) group.

[0282] The following abbreviations are used for the com-
 mon amino acids referred to herein.

<table>
<thead>
<tr>
<th>TABLE 6</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abbreviation</td>
</tr>
<tr>
<td>Ala</td>
</tr>
<tr>
<td>Arg</td>
</tr>
<tr>
<td>Asn</td>
</tr>
<tr>
<td>Asp</td>
</tr>
<tr>
<td>Cys</td>
</tr>
<tr>
<td>Gln</td>
</tr>
<tr>
<td>Glu</td>
</tr>
<tr>
<td>Gly</td>
</tr>
<tr>
<td>His</td>
</tr>
<tr>
<td>Ile</td>
</tr>
<tr>
<td>Leu</td>
</tr>
<tr>
<td>Lys</td>
</tr>
<tr>
<td>Met</td>
</tr>
<tr>
<td>Phe</td>
</tr>
<tr>
<td>Pro</td>
</tr>
<tr>
<td>Pyl</td>
</tr>
<tr>
<td>Ser</td>
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<td>Sec</td>
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<tr>
<td>Thr</td>
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<tr>
<td>Trp</td>
</tr>
<tr>
<td>Tyr</td>
</tr>
<tr>
<td>Val</td>
</tr>
<tr>
<td>Asx</td>
</tr>
<tr>
<td>Glx</td>
</tr>
<tr>
<td>Xaa</td>
</tr>
<tr>
<td>Xle</td>
</tr>
</tbody>
</table>

[0283] Some modified amino acids are indicated as fol-
 lows:
“D-2-NaI” is 2-naphthyl-D-alanine,
“SertBu” is t-butyl serine,
“Azagly” is aza glycine,
“Me” is methyl,
Met(O) is methionine sulfoxide,
“Pyr” and “pGlu” are pyroglutamic acid,
“Tyr(SO₃H)” is sulphated tyrosine,
“Tyr(Me)” is methyltyrosine,
“NHEt” is ethylamide.

EXAMPLES

[0284] The peptides as listed above were tested for activity
using the assays described in Examples 1 to 17. The tested
peptides are all commercially available.

Example 1

HIV-1 Experiments

[0285] CEM-SS cells were passaged in T-75 flasks prior to
use in the antiviral assay. On the day preceding the assay, the
cells were split 1:2 to assure they were in an exponential
growth phase at the time of infection. Total cell viability
quantification was performed using a hemacytometer and
trypan blue exclusion. Cell viability was greater than 95% for
the cells to be utilized in the assay. The cells were resus-
pended at 5×10⁶ cells/ml in tissue culture medium and added
to the peptide-containing microtiter plates in a volume of 50
microliters.

[0286] The virus used was the lymphocytotropic strain
HIV-1_RR. Virus was obtained from NIH AIDS Research
and Reference Reagent Program and was grown in CEM-SS cells
for the production of stock virus pools. For each assay, a pre-
titered aliquot of virus was removed from the freezer
(−80°C) and allowed to thaw slowly to room temperature in
a biological safety cabinet. The virus was resuspended and
diluted into tissue culture medium such that the amount of
virus added to each well in a volume of 50 microliters was the
amount determined to give between 85% to 95% cell killing
after 6 days post-infection. TCID₅₀ calculations by endpoint
titration in CEM-SS cells indicated that the multiplicity of
infection was approximately 0.01. AZT (nucleoside reverse
transcriptase inhibitor; NRTI) and indinavir (protease
inhibitor; PI) were used as positive control antiviral compo-
unds.

Plate Format

[0287] Each plate contained cell control wells (cells only),
virus control wells (cells plus virus), drug cytotoxicity wells
(cells plus peptide only), peptide colorimetric control wells
(peptide only) as well as experimental wells (peptide—10
micromers per ml—plus cells plus virus). Samples were
evaluated for antiviral efficacy with triplicate measurements
and with duplicate measurements to determine cellular cyto-
toxicity, if detectable.

[0288] At assay termination, the plates were stained with
the soluble tetrazolium-based dye MTS (CellTiter 96
Reagent, Promega) to determine cell viability and quantify
peptide toxicity. MTS is metabolized by the mitochondrial
enzymes of metabolically active cells to yield a soluble for-
mazan product, allowing the rapid quantitative analysis of
cell viability and peptide cytotoxicity. This reagent is stable,
single solution that does not require preparation before use.
At assay termination, 20-25 microliters of MTS reagent
was added per well and the microtiter plates were then
incubated for 5 hours at 37°C, and 5% CO₂ to assess cell viability.
Adhesive plate sealers were used in place of lids, the sealed
plates were inverted several times to mix the soluble forma-
zan product and the plate was read spectrophotometrically at
490/560 nm with a Molecular Devices Vmax plate reader.

[0289] The overall assay performance was valid based
upon judgement of the positive control compounds AZT and
indinavir exhibiting the expected levels of antiviral activity. Macroscopic observation of the cells in each well of the microtiter plate confirmed the cytotoxicity results obtained following staining of the cells with the MTS metabolic dye.

Results from HIV Experiments:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% CPE reduction in HIV-1 infected CEM-SS cells</th>
<th>% Cell viability</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>0.0</td>
<td>100</td>
</tr>
<tr>
<td>AZT (positive control)</td>
<td>99</td>
<td>97</td>
</tr>
<tr>
<td>Indinavir (positive control)</td>
<td>100</td>
<td>95</td>
</tr>
</tbody>
</table>

**Example 2**

**HBV Experimental Assay System**

[H0290] HepG2-2.2.15 is a stable cell line containing the hepatitis B virus (HBV) ayw strain genome (ATCC Cat. No. CRL-11997). Antiviral compounds blocking any late step of viral replication such as transactivation, translation, pregenome encapsidation, reverse transcription, particle assembly and release can be identified and characterized using this cell line. In this assay, an active compound will reduce the production of secreted HBV from cells, measured by utilizing real time quantitative PCR (TaqMan) assay to directly and accurately measure HBV DNA copies. The analysis of this data allows to calculate:

[H0291] Antiviral activity

[H0292] Compound Cytotoxicity

[H0293] HepG2-2.2.15 cells were plated in 96-well microtiter plates. After 16-24 hours the confluent monolayer of HepG2-2.2.15 cells was washed and the medium was replaced with complete medium containing test peptide—10 micrograms per ml—in duplicate. Lamivudine (3TC) was used as the positive control, while media alone was added to the cells as a negative control (virus control). Three days later the culture medium was replaced with fresh medium containing the peptide. Six days following the initial administration of the peptide, the cell culture supernatants was collected, treated with pronase and DNase and then used in a real-time quantitative TaqMan PCR assay. The PCR-amplified HBV DNA was detected in real-time by monitoring increases in fluorescence signals that result from the exonuclease degradation of a quenched fluorescence probe molecule that hybridizes to the amplified HBV DNA. For each PCR amplification, a standard curve was simultaneously generated using dilutions of purified HBV DNA. Antibacterial activity was calculated from the reduction in HBV DNA levels (% virus control). A novel dye uptake assay was then employed to measure cell viability, which is used to calculate toxicity (% cell control).

Results from HBV Experiments:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% inhibition of HBV replication in HEP G2 cells</th>
<th>% Cell viability</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>13.4</td>
<td>97.8</td>
</tr>
<tr>
<td>3TC (positive control)</td>
<td>92.0</td>
<td>95.8</td>
</tr>
</tbody>
</table>

**Example 3**

**HCMV Experimental Assay System**

[H0294] MRC-5 cells (human embryonal lung fibroblasts) were obtained from the American Type Culture Collection (ATCC CCL-171; Rockville, Md.) and grown in Eagle’s Minimum Essential Medium with Earle’s BSS (EMBM) supplemented with 10% fetal bovine serum (FBS), 0.1 mM non-essential amino acids, 1.0 mM sodium pyruvate, 2.0 mM L-Glutamine, 100 units/ml Penicillin and 100 micrograms/ml Streptomycin. Cells were split twice a week 1:2.

[H0295] HCMV strain AD169 was obtained from ATCC (ATCC VR-538). Virus stocks were prepared by infecting 80% confluent MRC-5 cells at a minimal multiplicity of infection in MRC-5 growth medium containing 2% FBS. Monolayers were incubated at 37°C, 5% CO₂, until 90%–95% viral cytopathic effect (CPE) was observed (10-13 days). Culture medium was then collected from the cells, centrifuged at low speed to remove cellular debris, aliquoted in 1 ml volumes and stored at –80°C as stock virus.

[H0296] MRC-5 cells were seeded at 75,000 cells/well in 24 well plates using MRC-5 growth medium. The plates were incubated overnight at 37°C, 5% CO₂. The following day, medium was removed and 100 plaque forming units (pfu) of HCMV was added to the wells. Virus was allowed to adsorb onto the cells for 1 hour at 37°C, 5% CO₂. Peptide was diluted—10 micrograms per ml—in assay medium containing 0.5% Methylcellulose.

[H0297] After the incubation period, 1 ml of each peptide solution was added to the wells without aspirating the virus inoculums. The plates were incubated for 7-10 days to allow for plaque formation. Ganciclovir was used as positive control. Cultures were examined microscopically and toxicities were noted. The Media was aspirated from the wells and the cells were fixed and stained using 20% methanol containing Crystal Violet followed by enumeration of plaques by microscopic inspection.

[H0298] For cytotoxicity testing, MRC-5 cells were seeded at 2,500 cells/well in 96 well plates using growth medium. The plates were incubated overnight at 37°C, 5% CO₂. The following day, peptide was added and tested in duplicates. After a 6 days incubation period, cell viability was measured using CellTiter 96 Solution (Promega). Plates were incubated for additional 4 hours at 37°C. Adhesive plate sealers were used in place of lids, the sealed plates were inverted several times to mix the soluble formazan product and the plate was read spectrophotometrically at 490/560 nm with a Molecular Devices Vmax plate reader.

[H0299] The overall assay performance was valid based upon judgement of the positive control compound Ganciclovir exhibiting the expected levels of antiviral activity. Macroscopic observation of the cells in each well of the microtiter
plate confirmed the cytotoxicity results obtained following staining of the cells with the MTS metabolic dye. Results from HCMV Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% plaque reduction</th>
<th>% Cell viability</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>0.0</td>
<td>100</td>
</tr>
<tr>
<td>Ganciclovir</td>
<td>100</td>
<td>100</td>
</tr>
</tbody>
</table>

Example 4
Methicillin Resistant Staphylococcus Aureus (MRSA) Assay

[0300] The antibacterial assay was conducted using clear, U-bottom 96-well microtiter plates. Cation-adjusted Mueller-Hinton Broth (MHB) was used for testing MRSA. The peptide of the invention (0.1 ml of each—10 micrograms per ml)—was dispensed into wells in duplicate. Then the wells were inoculated with 5x10^3 CFU/ml MRSA in 0.1 ml volume. For control purposes, each plate included 4 wells containing media without bacterial inoculum and 4 wells containing medium with inoculum but without peptide. The plates were incubated for 12 h at 37°C, and read visually 18-24 hours post-incubation. Growth control of MRSA was examined first to determine adequacy of media preparations and growth conditions. Acceptable growth is defined as ≥ 2 mm wide button of cells at the bottom of each sample well, or obvious turbidity in the culture supernatant. Test wells were examined and scored as positive/negative for activity. A positive score for activity is based on complete inhibition of macroscopic growth of the test Pseudomonas aeruginosa.

Results from MRSA Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>0.0</td>
</tr>
</tbody>
</table>

Example 5
Pseudomonas aeruginosa Assay

[0301] The antibacterial assay was conducted using clear, U-bottom 96-well microtiter plates. Cation-adjusted Mueller-Hinton Broth (MHB) was used for testing Pseudomonas aeruginosa. The peptide of the invention (0.1 ml of each—10 micrograms per ml)—was dispensed into wells in duplicate. Then the wells were inoculated with 5x10^3 CFU/ml Pseudomonas aeruginosa in 0.1 ml volume. For control purposes, each plate included 4 wells containing media without bacterial inoculum and 4 wells containing medium with inoculum but without peptide. The plates were incubated for 12 h at 37°C, and read visually 18-24 hours post-incubation. Growth control of Pseudomonas aeruginosa was examined first to determine adequacy of media preparations and growth conditions. Acceptable growth is defined as ≥ 2 mm wide button of cells at the bottom of each sample well, or obvious turbidity in the culture supernatant. Test wells were examined and scored as positive/negative for activity. A positive score for activity is based on complete inhibition of macroscopic growth of the test Pseudomonas aeruginosa.

Results from Pseudomonas aeruginosa Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>100.0</td>
</tr>
</tbody>
</table>

Example 6
Streptococcus pneumoniae Assay

[0302] The antibacterial assay was conducted using clear, U-bottom 96-well microtiter plates. Cation-adjusted Mueller-Hinton Broth (MHB) was used for testing Streptococcus pneumoniae. The peptide of the invention (0.1 ml of each—10 micrograms per ml)—was dispensed into wells in duplicate. Then the wells were inoculated with 5x10^3 CFU/ml Streptococcus pneumoniae in 0.1 ml volume. For control purposes, each plate included 4 wells containing media without bacterial inoculum and 4 wells containing medium with inoculum but without peptide. The plates were incubated for 12 h at 37°C, and read visually 18-24 hours post-incubation. Growth control of Streptococcus pneumoniae was examined first to determine adequacy of media preparations and growth conditions. Acceptable growth is defined as ≥ 2 mm wide button of cells at the bottom of each sample well, or obvious turbidity in the culture supernatant. Test wells were examined and scored as positive/negative for activity. A positive score for activity is based on complete inhibition of macroscopic growth of the test Streptococcus pneumoniae.

Results from Streptococcus pneumoniae Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>100.0</td>
</tr>
</tbody>
</table>

Example 7
Mycobacterium tuberculosis Assay

[0303] The antibacterial assay was conducted using clear, U-bottom 96-well microtiter plates. Middlebrook 7H12 assay medium was used for testing drug-resistant Mycobacterium tuberculosis. The peptide of the invention (0.1 ml of each—10 micrograms per ml—10 micrograms per ml)—was dispensed into wells in duplicate. Then the wells were inoculated with 5x10^3 CFU/ml. Mycobacterium tuberculosis in 0.1 ml volume. For control purposes, each plate included 4 wells containing media without bacterial inoculum and 4 wells containing medium with inoculum but without peptide. The plates were incubated for seven days at 37°C, and read visually thereafter. Growth control of Mycobacterium tuberculosis was examined first to determine adequacy of media preparations and growth conditions. Acceptable growth is defined as ≥ 2 mm wide button of cells at the bottom of each sample well, or obvious turbidity in the culture supernatant. Test wells were examined and scored as positive/negative for activity. A positive score for activity is based on complete inhibition of macroscopic growth of the test Mycobacterium tuberculosis. The drug-resistant Mycobacterium tuberculosis
that was used in the assay is resistant against following medications: para-aminosalicylic acid (PAS), streptomycin and isoniazid (INH).

Results from *Mycobacterium tuberculosis* Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>100.0</td>
</tr>
</tbody>
</table>

**Example 8**

**Cell Cycle Assay**

[0304] Human A549 cells (carcinomic human alveolar basal epithelial cells) were utilized in the experiments employing the Propidium iodide cell cycle assay. The eukaryotic cell cycle is a series of events that take place in a cell leading to its replication. The regulation of the cell cycle involves steps crucial to the cell, including detecting and repairing genetic damage, and provision of various checks to prevent uncontrolled cell division. The molecular events that control the cell cycle are ordered and directional; that is, each process occurs in a sequential fashion. The cell cycle consists of four distinct phases: G₁ phase, S phase, G₂ phase (collectively known as interphase) and M phase. M phase is itself composed of two tightly coupled processes: mitosis, in which the cell’s chromosomes are divided between the two daughter cells, and cytokinesis, in which the cell’s cytoplasm divides forming distinct cells. Activation of each phase is dependent on the proper progression and completion of the previous one. Cells that have temporarily or reversibly stopped dividing are said to have entered a state of quiescence called G₀ phase. The relatively brief M phase consists of nuclear division and cytoplasmic division. The first phase within interphase, from the end of the previous M phase till the beginning of DNA synthesis is called G₁ (G indicating gap or growth). During this phase the biosynthetic activities of the cell resume at a high rate. This phase is marked by synthesis of various enzymes that are required in S phase, mainly those needed for DNA replication. The ensuing S phase starts when DNA synthesis commences; when it is complete, all of the activities of DNA replication and repair are completed. The cell then enters the G₂ phase, which lasts until the cell enters mitosis. Significant protein synthesis occurs during this phase, mainly involving the production of microtubules, which are required during the process of mitosis. Inhibition of protein synthesis during G₂ phase prevents the cell from undergoing mitosis.

[0305] Disregulation of the cell cycle components may lead to tumor formation.

[0306] Propidium iodide is an intercalating agent and a fluorescent molecule that can be used to stain DNA. Cells were incubated for 24 hours with test peptide—10 micrograms per ml—or left untreated. After that cells were trypsinized, suspended in medium +10% FCS, centrifuged (1000 rpm, 5 min) and the cell pellet resuspended in PBS (1 ml). The cells were pipetted into 2.5 ml absolute EtOH (final concentration approx. 70%) and incubated on ice for 15 min. Thereafter, cells were pelleted at 1500 rpm for 5 min and resuspended in Propidium iodide solution in PBS. After incubation for 40 min at 37°C, cells were analyzed in the FACS.

Results from Cell Cycle Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>G₀/G₁</th>
<th>S</th>
<th>G₂/M</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>59.1</td>
<td>31.8</td>
<td>9.2</td>
</tr>
</tbody>
</table>

**Example 9**

**T Cell Proliferation Assay**

[0307] Human Peripheral Blood Mononuclear Cells (PBMC) were obtained from normal human donors. The T cell proliferation was induced by stimulation of the cells with the T cell mitogen phytohemagglutinin (PHA), either in the absence (positive proliferation control), or in the presence of test peptide—10 micrograms per ml—to examine their effects on the T cell proliferating response. 10³/well PBMC were plated in 96-well microtiter plates and assayed in duplicate with the peptide. Cell cultures were incubated at 37°C for 3 days in a 5% CO₂ incubator and were thereafter pulsed with 1 microCi/well³H-thymidine for additional 12 hours of culture. At the end of incubation time, the plates were harvested and the cells counted by liquid scintillation for the incorporation of³H-thymidine as a measure of T cell proliferation.

Results from T cell Proliferation Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% of PHA induced control</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>100.0</td>
</tr>
</tbody>
</table>

**Example 10**

**B Cell Proliferation Assay**

[0308] Human Peripheral Blood Mononuclear Cells (PBMC) were obtained from normal human donors. The B cell proliferation was induced by stimulation of the cells with the B cell mitogen *Staphylococcus aureus* Cowans I (SAC) plus Interleukin-2, either in the absence (positive proliferation control), or in the presence of test peptide—10 micrograms per ml—to examine their effects on the B cell response. 10³/well PBMC were plated in 96-well microtiter plates and assayed in duplicate with the peptide. Cell cultures were incubated at 37°C for 3 days in a 5% CO₂ incubator and were thereafter pulsed with 1 microCi/well³H-thymidine for additional 12 hours of culture. At the end of incubation time, the plates were harvested and the cells counted by liquid scintillation for the incorporation of³H-thymidine as a measure of B cell proliferation.

Results from B Cell Proliferation Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% of SAC/IL2 induced control</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>115.3</td>
</tr>
</tbody>
</table>
Example 11
Phagocytosis Assay

[0309] RAW 264.7 (Mouse leukaemic monocyte macrophage cell line) cells were obtained from ATCC and grown in RPMI 1640 medium containing 10% FBS. Cells were incubated in 12x75 mm tubes at 37°C with test peptide—10 micrograms per ml—for 30 min prior to adding Fluorescein-labeled Escherichia coli bacteria as the agent to be ingested. After the cells were incubated for additional 60 min at 37°C, and allowed to ingest the Fluorescein-labeled Escherichia coli bacteria, cells were fixed with 1% paraformaldehyde. The samples were then analyzed by flow cytometry to determine the amount of phagocytosis as a function of brightness (the greater the phagocytic activity, the more fluorescence in the macrophage population). Data are reported as % positive and the mean fluorescence intensity (MFI) of positively stained cells.

Results from Phagocytosis Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% of control phagocytosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>105.9</td>
</tr>
</tbody>
</table>

Example 12
Apoptosis Induction Assay

[0310] Human A549 cells (carcinomic human alveolar basal epithelial cells) were utilized in the experiments employing the Annexin-5 apoptosis assay. Annexin-5 is a member of a highly conserved protein family that binds acidic phospholipids in a calcium-dependent manner. Annexin-5 possesses a high affinity for phosphatidylserine. Phosphatidylserine is translocated from the inner side of the plasma membrane to the outer layer when cells undergo death by apoptosis or cell necrosis and serves as a signal by which cell destined for death are recognized by phagocytes. Test peptide—10 micrograms per ml—were exposed for 24 hours to the A549 cells before they were analyzed for signs of apoptosis.

Results from Apoptosis Induction Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% of induction</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>2.6</td>
</tr>
</tbody>
</table>

Example 13
Apoptosis Prevention Assay

[0311] Human A549 cells (carcinomic human alveolar basal epithelial cells) were utilized in the experiments employing the Annexin-5 apoptosis assay. Annexin-5 is a member of a highly conserved protein family that binds acidic phospholipids in a calcium-dependent manner. Annexin-5 possesses a high affinity for phosphatidylserine. Phosphatidylserine is translocated from the inner side of the plasma membrane to the outer layer when cells undergo death by apoptosis or cell necrosis and serves as a signal by which cell destined for death are recognized by phagocytes. A549 cells were pretreated for 30 min with test peptide—10 micrograms per ml—followed by the exposure to C2 ceramide. Ceramide mediates cell apoptosis through the activation of the mitogen activating protein kinase (MAPK) and the stress activated kinase (JNK/SAPK). C2 ceramide is a synthetic, membrane soluble analog of ceramide.

Results from Apoptosis Prevention Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% prevention of ceramide induced apoptosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>0.0</td>
</tr>
</tbody>
</table>

Example 14
Th1/Th2 Cytokine Profiling Assay

[0312] The Balb/c mice (originated in 1923, it is a popular strain and is used in many different research disciplines. Also classified as an inbred from the production of 20 or more successive brother-sister matings, the Balb/c mouse is albino and small in size) were immunized on Days 1, 15, and 29 with Ovalbumin (Ovalbumin is the main protein found in egg white, commonly used to simulate an immunological reaction in test animals) in PBS (5 micrograms/injection). On day 50, spleens of the mice were harvested (3 weeks after last boost with Ovalbumin). Cells were cultured (2x10^7/well in triplicate) and incubated with culture medium or test peptide—10 micrograms per ml—for 30 min. Thereafter, additional Ovalbumin was added to the cells at 10 micrograms/ml for in vitro restimulation of the cells. 72 hours later, cell supernatants were harvested and assayed using the Becton Dickinson Mouse Th1/Th2 Cytokine CBA Kit. This kit can be used to measure Interleukin-2 (IL-2), Interleukin-4 (IL-4), Interleukin-5 (IL-5), Interferon-γ (IFN-γ), and Tumor Necrosis Factor-α (TNF-α) protein levels in a single sample. The kit performance has been optimized for analysis of physiologically relevant concentrations (µg/ml levels) of specific cytokine proteins in tissue culture supernatants and serum samples.

Results from Th1/Th2 Cytokine Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>TNF-α spleen cells; Th1 response (% of control)</th>
<th>IFN-γ spleen cells; Th1 response (% of control)</th>
<th>IL-2 spleen cells; Th1 response (% of control)</th>
<th>IL-4 spleen cells; Th2 response (% of control)</th>
<th>IL-5 spleen cells; Th2 response (% of control)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>113.0</td>
<td>56.0</td>
<td>159.5</td>
<td>114.4</td>
<td>96.1</td>
</tr>
</tbody>
</table>
Example 15

TNF Alpha Production Assay

[0313] Human Peripheral Blood Mononuclear Cells (PBMC) were obtained from normal human donors. The macrophages were prepared by adherence of PBMC to the plastic wells of the plates. After 8 days in culture in the presence of recombinant human macrophage-colony stimulating factor at 2 ng/ml, differentiated macrophages were preincubated with test peptide—10 micrograms per ml—for 30 min, followed by in-well stimulation by the addition of lipopolysaccharide at a final concentration of 200 ng/ml. Not stimulated macrophages served as negative background control.

[0314] After overnight incubation, supernatants from the control and LPS-stimulated cultures were harvested and assayed for TNF alpha production employing a TNF alpha specific ELISA. Results from TNF Alpha Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% of LPS induction</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>105.8</td>
</tr>
</tbody>
</table>

Example 16

Endothelial Cell Migration Assay

[0315] Endothelial cell migration is a prerequisite for the process of neo-vascularization or angiogenesis which is crucial for on-site recruitment of blood vessel formation. Primary Human endothelial cells (HUV EC) were seeded in insert chambers with 3 micrometer pore size of multi-transwell plate for 6 hours at 37°C in Endothelial Cell Basal Medium (EBM) supplemented with 0.1% bovine serum albumin. Thereafter, designated concentration of test peptide—10 micrograms per ml—was added in duplicate wells. The endothelia were allowed to migrate for 22 hours at 37°C, then, migrated cells were fixed and stained with Hoechst 33342 dye. Images of 3 fields per insert were taken and the number of migrated cells per field were quantified using the ImageProPlus software. Data were analyzed for the average number of the migrated cells and standard deviation of six data points for each treatment condition. Active test peptide against HUV EC migration was determined based on 50% inhibition of migrated cells as compared with the control. Statistic p values were computed using the Student’s t-test. Results from Endothelial Cell Migration Assay:

<table>
<thead>
<tr>
<th>Compound</th>
<th>% inhibition of migration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Octreotide</td>
<td>-24</td>
</tr>
</tbody>
</table>

Example 17

Endothelial Tube Formation Assay

[0316] The endothelial tube formation assay is based on the ability of endothelial cells to form three-dimensional capillary-like tubular structures when cultured on a gel of basement membrane extract. The endothelial tube formation assay represents a powerful model for studying inhibition and induction of angiogenesis. Pre-labeled HUV EC with Calcein AM were seeded in a 96-well culture plate coated with extracellular matrix (Chemicon international Cat. ECM625) and treated with test peptide—10 micrograms per ml—in full growth medium. Positive control was vehicle only. The endothelial cells were allowed to form tubes for 20 hours and were then examined under an inverted fluorescence microscope. Duplicate wells for each treatment were photographed and quantitatively analyzed for an average tubule length using image analysis software ImageProPlus. Raw data were expressed as average tubule lengths in pixels ± standard deviation. Statistic p values were computed using the Student’s t-test.

[0317] Results from Endothelial Tube Formation Assay:

Example 18

Mother Milk Formulation

[0318] Methods to prepare mother milk or artificial mother milk formulations or mother milk substitutes are described in WO03043429, U.S. Pat. No. 5,962,062, WO0030461, EP0527283, EP0832565

[0319] One example of an artificial mother milk or mother milk substitute formulation is provided in the following while also the other formulations disclosed in the above mentioned references can be used and are included herewith by reference.

[0320] The milk substitute contains, by weight, approximately 15% skimmed milk solids, approximately 75% demineralized water, approximately 9% soya oil, approximately 0.02% of carrageenates, 0.2% lecithin, and approximately 0.2% of disodium hydrogenphosphate.

[0321] In a first step, the solubilizing aqueous medium is produced, comprises, by weight, approximately 75% of water, approximately 0.02% of carrageenate and approximately 0.2% of disodium hydrogenphosphate.

[0322] The skimmed milk powder is then added to the solution for 10 min at 60°C, and dissolved in the liquid.

[0323] Then soya oil and lecithin are added to the milk substitute composition at 60°C. The milk composition is allowed to stand 30 min at 55°C. After pasteurization, the peptide of the invention is added in liquid or powder form in such a quantity that the milk composition obtained comprises an amount of 5-50 micrograms, preferably 10-40 micrograms per 100 ml of milk composition. Optionally peptide 2 could be added in similar or smaller amounts to the obtained composition.

Example 19

Gel Formulation

[0324] 0.5 g of peptide
1.6 g of isopropanol
1.0 g of glycerol
1.6 g of polyoxyethylene-polyoxypropylene copolymer 12500 (Pluronic F127) 5.5 g of water are mixed for 10 minutes and then heated to 85°C under continuous stirring for 15 minutes. The solution is cooled to room temperature under stirring. During the cooling phase the solution begins to gel at a temperature of about 45°C to form a clear gel. The gel contains 5% of the peptide combination for medical use. Optionally peptide 2 could be added in an amount form 0.01 to 0.5 g.

Example 20 Lotion Formulation

**0325** 0.5 g of peptide 1.9 g of isopropanol 1.0 g of dimethylisosorbide 1.0 g of polyoxyethylene-polyoxypropylene copolymer 12500 (Pluronic F127) 5.6 g of water are stirred and heated at 50°C, until a clear solution has been formed. Then the composition is cooled to room temperature under stirring. The lotion contains 5% of peptide combination for medical use. Optionally peptide 2 could be added in an amount form 0.01 to 0.5 g.

16. The pharmaceutical composition of claim 13, wherein said composition is prepared as a lyophilized formulation or a buffered liquid formulation.

17. The pharmaceutical composition of claim 13, wherein said composition comprises at least one pharmaceutical acceptable carrier, cryoprotectant, lyoprotectant, excipient or diluent.

18. A method of treatment of cancer, autoimmune disease, fibrotic disease, inflammatory disease, neurodegenerative disease, infectious disease, lung disease, heart and vascular disease and metabolic disease, the method comprising, administering to a patient in need thereof, a therapeutically effective amount of a pharmaceutical composition comprising a peptide consisting of the sequence H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-L-threoninol (SEQ ID NO:1) or salts or hydrates thereof, wherein administration of the pharmaceutical composition treats said diseases.

19. The method of claim 18, wherein the cancer, autoimmune disease, fibrotic disease, inflammatory disease, neurodegenerative disease, infectious disease, lung disease, heart and vascular disease and metabolic disease is selected from Mycobacterium tuberculosis infection, Streptococcus pneumoniae infection, Streptococcus pneumoniae infection related diseases, hemolytic uremic syndrome, pneumonia, meningitis, cystic fibrosis complication, middle ear diseases and infections of the bloodstream.

20. The method of claim 18, wherein the peptide is administered by intravenous administration, oral administration, or administration by inhalation.

21. The method of claim 18, wherein the peptide is administered as a lyophilized formulation or as a buffered liquid formulation.

**SEQ LISTING**

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1  5
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1-12. (canceled)

13. A pharmaceutical composition comprising a peptide consisting of the sequence H-D-Phe-Cys-Phe-D-Trp-Lys-Thr-Cys-L-threoninol (SEQ ID NO:1).

14. The pharmaceutical composition of claim 13, wherein said composition is incorporated in a nutritional formulation.

15. The pharmaceutical composition of claim 14, wherein the nutritional formulation is an artificial mother milk formulation or mother milk substitute suitable for oral administration to newborns, toddlers and infants.