

(12) STANDARD PATENT
(19) AUSTRALIAN PATENT OFFICE

(11) Application No. **AU 2006340367 B2**

- (54) Title
Composition for treating bacterial, viral, fungal diseases, inflammation and pain
- (51) International Patent Classification(s)
A01N 25/00 (2006.01)
- (21) Application No: **2006340367** (22) Date of Filing: **2006.03.20**
- (87) WIPO No: **WO07/108807**
- (43) Publication Date: **2007.09.27**
(44) Accepted Journal Date: **2012.12.20**
- (71) Applicant(s)
Olatec Industries LLC
- (72) Inventor(s)
Torrence, Christopher
- (74) Agent / Attorney
Shelston IP, L 21 60 Margaret St, Sydney, NSW, 2000
- (56) Related Art
US 5084482 A
US 5053429 A
UNNIKRISHNAN M. et al. AGENTS AND ACTIONS, vol. 31(1-2), 1990, pages
110-112

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
27 September 2007 (27.09.2007)

PCT

(10) International Publication Number
WO 2007/108807 A1

(51) International Patent Classification:
A01N 25/00 (2006.01)

(21) International Application Number:

PCT/US2006/010665

(22) International Filing Date: 20 March 2006 (20.03.2006)

(25) Filing Language:

English

(26) Publication Language:

English

(71) Applicant (for all designated States except US): **OLATEC INDUSTRIES LLC** [US/US]; 128 Doral Greens Drive West, Rye Brook, New York 10573 (US).

(72) Inventor; and

(75) Inventor/Applicant (for US only): **TORRENCE, Christopher** [US/US]; 95 Leroy Avenue, Darien, Connecticut 06820 (US).

(74) Agent: **KUNG, Viola, T.**; Howrey LLP, 2941 Fairview Park Drive, Box 7, Falls Church, Virginia 22042 (US).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,

CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

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

Declaration under Rule 4.17:

— as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))

Published:

— with international search report

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



WO 2007/108807 A1

(54) Title: COMPOSITION FOR TREATING BACTERIAL, VIRAL, FUNGAL DISEASES, INFLAMMATION AND PAIN

(57) Abstract: The present invention is directed to compositions and methods of treating bacterial, viral, fungal diseases; inflammation or inflammatory-related disorders; pain; and skin conditions. The composition comprises an organic solvent extract, which is prepared by the method comprising the steps of: (a) mixing methionine with water, (b) adding an aqueous hypochlorite solution to the methionine solution and mixing, (c) adding a water-immiscible organic solvent to (b) and mixing, and (d) separating the organic solvent phase from the water phase to obtain the organic solvent extract.

COMPOSITION FOR TREATING BACTERIAL, VIRAL, FUNGAL DISEASES, INFLAMMATION AND PAIN

FIELD OF THE INVENTION

5 The present invention relates to compositions and methods of treating bacterial, viral, fungal diseases; inflammation or inflammatory-related disorders; pain; and skin conditions. The composition is prepared by the organic solvent extraction of a reaction product of methionine and hypochlorite.

BACKGROUND OF THE INVENTION

10 The human body is susceptible to many different types of infections from a variety of sources. Viral infection, usually in the form of the common cold, affects virtually everyone each year. While the coughing and sneezing associated with colds may be merely annoying, other common viral infections can be far more serious.
15 Influenza, for example, remains a leading cause of hospitalization and death among Americans, accounting for an average of 36,000 fatalities and 114,000 hospitalizations per year.

 Bacterial infections can be just as dangerous. Staphylococcus infections, which are caused by bacteria from the staphylococcus family, account for many serious post-
20 surgical complications. Staphylococcus infection is also the leading culprit in cases of food poisoning, and can be responsible for such life-threatening conditions as Toxic Shock Syndrome (TSS), pneumonia, bone infections (osteomyelitis), mastitis in nursing mothers, endocarditis (infection of the inside of the heart), and bacteremia (blood
25 infection). People who are otherwise healthy typically do not become severely ill from staphylococcus infections, but individuals with weakened immune systems, including the elderly, newborns, and persons with chronic illnesses, such as diabetes, cancer, lung disease, kidney disease, or HIV/AIDS, are at special risk.

 Individuals with weakened immune systems are at risk from fungal infections. Fungal infections cause conditions in millions of people in the form of sinus infections,
30 athlete's foot, and yeast infections.

 The general term "pain" used herein represents all categories of pain, such as traumatic pain resulting from injury, post surgical pain, inflammatory pain; pain associated with disease

diabetic neuropathy, causalgia, brachial plexus avulsion, occipital neuralgia, fibromyalgia, gout, and other forms of neuralgic, neuropathic and idiopathic pain syndromes; pain of varying severity, i.e. mild, moderate and severe pain; acute and chronic pain; and specific organ pain, such as ocular and corneal pain, bone pain, heart pain, skin/burn pain, visceral (kidney, gall bladder, etc.), joint, dental and muscle pain.

Connective tissues are subjected to a constant barrage of stress and injury. Acute or chronic impacts and the natural progression of various degenerative diseases all produce painful inflammation in joint regions, such as the neck, back, arms, hips, ankles and feet. These afflictions are common and often debilitating.

Current therapies of pain include the use of opioid narcotic analgesics such as morphine and fentanyl, nonsteroidal anti-inflammatory drugs (NSAIDS) such as aspirin, ibuprofen and cyclooxygenase inhibitors, or ion channel blockers such as lidocaine and novacaine. These therapies all have limitations, for example, they cause tolerance, dependence, constipation, respiratory depression and sedation (opioids). NSAIDS have gastrointestinal side effects and increase bleeding time, and are not effective in treating severe pain.

Inflammation is a localized reaction of live tissue due to an injury, which may be caused by various endogenous and exogenous factors. The exogenous factors include physical, chemical, and biological factors. The endogenous factors include inflammatory mediators, antigens, and antibodies. Endogenous factors often develop under the influence of an exogenous damage. An inflammatory reaction is often followed by an altered structure and penetrability of the cellular membrane. At the tissue and organ level, inflammation is indicated by pain, swelling, reddening, increased temperature, and loss of function in some cases.

Inflammation is influenced by various exogenous and endogenous agents. Endogenous factors, namely, mediators, antigens, and autogens define the nature and type of an inflammatory reaction, especially its course in the zone of injury. In the case where a tissue damage is limited to the creation of mediators, an acute form of inflammation develops. If immunologic reactions are also involved in the process, through the interaction of antigens, antibodies, and autoantigens, a long-term inflammatory process will develop. Various exogenous agents, for example, infection, injury, radiation, also provide the course of inflammatory process on a molecular level by damaging cellular membranes which initiate biochemical reactions.

Nonsteroidal anti-inflammatory drugs (NSAIDS), such as aspirin, can block certain links

of an inflammatory process, but these drugs cannot stabilize damaged cellular membranes, which makes their influence on an inflammatory process limited and insufficient.

There is a need for a composition and a method for treating bacterial, viral, fungal diseases; inflammation or inflammatory-related disorders; pain; and skin conditions. The
5 composition should be economic and easy to manufacture, and the method should be effective and have no significant side effects.

SUMMARY OF THE INVENTION

The present invention is directed to an organic solvent extract prepared by the reaction of an aqueous methionine solution with an aqueous hypochlorite solution, and the extraction
10 of the reactive product with a water-immiscible organic solvent such as mineral oil. The reactive product is stable in the water-immiscible organic solvent for at least several months at room temperature.

The present invention is also directed to a composition for use in a method for treating bacterial, viral, fungal diseases; inflammation or inflammatory-related disorders; pain;
15 or skin conditions. The method comprises the step of administering to a subject in need thereof a composition comprising the organic solvent extract as prepared according to the present invention. The composition can be applied by any of the accepted modes of systemic administration including topical, oral, and parenteral (such as intravenous, intramuscular, subcutaneous or rectal). Topical administration and oral administration are
20 preferred.

DETAILED DESCRIPTION OF THE INVENTION

The present invention describes an organic solvent extract that is useful in treating a variety of diseases or disorders. The organic solvent extract is prepared by the method comprising the steps of: (a) mixing methionine with water, (b) adding an aqueous
25 hypochlorite solution to the methionine solution and mixing at a temperature between 0°C to ambient temperature, (c) adding a water-immiscible organic solvent to (b) and mixing, and (d) separating the organic solvent phase from the water phase to obtain the organic solvent extract.

The organic solvent extract is prepared by first preparing a reactive product, and
30 then stabilizing the reactive product by extracting it into a water-immiscible organic solvent.

The reactive product is prepared by first mixing methionine with water until a suspension is formed. Methionine can be L-methionine, D-methionine, or a mixture thereof. The mixing is _____

preferably carried out under an inert gas, e.g. argon. The mixing time is typically a few seconds to a few minutes. The mixing can be done by any means of mixing, for example, by a blender.

An aqueous solution of hypochlorite such as sodium hypochlorite is added to the methionine solution and thoroughly mixed. Then the reaction mixture is rested until the mixture is cleared and the excess methionine forms foam on the top. The foam is optionally removed. The
5 reaction is carried out at a temperature between 0°C to ambient temperature, preferably at a low temperature such as 0-5°C. The reaction is preferably carried out under an inert gas such as argon.

The reactive product is not stable in water and is extracted with a water-immiscible organic solvent. The water-immiscible organic solvent useful in this invention is preferably a semi-polar or
10 non-polar solvent having a polarity of about 0.1-7.5, such as mineral oil, hexane, heptane, methylene chloride, vegetable oil, or ethyl acetate. A non-polar solvent is more preferred. The extraction is carried out immediately after the methionine/hypochlorite reaction, preferably, within 5 minutes, preferably within 2 minutes, and more preferably within 1 minute.

The water-organic solvent mixture is allowed to settle. The organic phase is separated from
15 the water phase by any means known to a skilled person such as pouring or pipetting, and the organic solvent extract is obtained. Any non-soluble residues in the organic solvent extract are optionally removed by filtration, decanting, centrifugation, or any means known to a skilled person. The reactive product is stable in the organic solvent at room temperature (22-28 °C) for at least a month, preferably, 3 months, more preferably 6 months or a year.

20 In a typical reaction, 700-800 ml of distilled water, 20-100 g of methionine, and 300-500 ml of 3-12 % hypochlorite are used. In a typical extraction, about 200-300 ml or more water-immiscible organic solvent is used. The amounts of the above reagents can be scaled up or scaled down.

The organic solvent extract of the present invention is useful for treating a variety of
25 diseases or disorders. The organic solvent extract can be used as is, or it can be administered in the form of a pharmaceutical composition that additionally contains a pharmaceutically acceptable carrier. In one embodiment, the organic solvent extract can be incorporated into any acceptable carrier, including creams, gels, lotions or other types of suspensions that can stabilize the active ingredient and deliver it to the affected area by topical applications. In another
30 embodiment, the pharmaceutical composition can be in the dosage forms such as tablets, capsules, granules, fine granules, powders, syrups, suppositories, injections, or the like. The

above pharmaceutical composition can be prepared by conventional methods.

In one embodiment, the present invention provides a composition for treating a bacterial disease such as staphylococcus infection.

In another embodiment, the present invention provides a composition for treating a
5 viral disease such as influenza infection.

In another embodiment, the present invention provides a composition for treating a fungal disease such as athlete's foot, yeast infection, and sinus infection caused by fungus infection.

In another embodiment, the present invention provides a composition for treating a
10 skin condition such as skin damages by burns or sun, skin blotches, or wart.

In another embodiment, the present invention provides a composition for treating inflammation or inflammatory-related disorders. The present invention alleviates symptoms associated with inflammation or inflammatory-related disorders, such as pain, swelling, reddening, increased temperature, or loss of function in some cases.

In yet another embodiment, the present invention provides a composition for
15 alleviate the symptoms of pain regardless of the cause of the pain. The general term "pain" treatable by the present method includes traumatic pain, neuropathic pain, organ pain, and pain associated with diseases. Traumatic pain includes pain resulting from injury, post-surgical pain and inflammatory pain. Neuropathic pain includes neuropathic and idiopathic
20 pain syndromes, and pain associated with neuropathy such as diabetic neuropathy, causalgia, brachial plexus avulsion, occipital neuralgia, fibromyalgia, gout, and other forms of neuralgia. Organ pain includes ocular, corneal, bone, heart, skin/burn, visceral (kidney, gall bladder, etc.), joint, dental and muscle pain. Pain associated with diseases includes pain associated with cancer, AIDS, arthritis, herpes and migraine. The present invention
25 reduces pain of varying severity, i.e. mild, moderate and severe pain; acute and chronic pain. For example, the present invention is effective in treating pain derived from inflammatory arthritis or degenerative arthritis, joint pain, muscle pain, tendon pain, and burn pain.

The methods for use of the compositions of the present invention comprise the step
30 of administering to a subject in need thereof a composition comprising the organic solvent extract as prepared according to the present invention. The composition of the present invention can be applied by any of the accepted modes of systemic administration including topical, oral, parenteral (such as intravenous, intramuscular, subcutaneous or rectal), and otherwise systemic routes of _____

administration. Dosing of the composition can vary based on the extent of the injury and each patient's individual response. Topical administration and oral administration are preferred.

When applied topically, the composition is preferably applied onto the affected area and rubbed into it. The composition can be applied 1-5 times daily, preferably 1-3 times daily, and preferably 3 times daily. The active compound passes through the skin and is delivered to the site of discomfort. Those of skill in the art will recognize that a wide variety of delivery mechanisms other than topical application, e.g., ingestible tablets, ingestible liquids, inhalers and aerosols are also suitable.

The following examples further illustrate the present invention. These examples are intended merely to be illustrative of the present invention and are not to be construed as being limiting.

EXAMPLES

Example 1. Preparation of Product

In a blender, 720-750 ml of distilled water at 0-4°C and 20 g DL methionine fine powder (Degussa Corp., Richfield Park, NJ) were combined and mixed for a few seconds under argon. After the methionine solid is uniformly suspended in the distilled water, 370-400 ml of Clorox bleach (6% sodium hypochlorite) at 0-4°C was added and mixed at a blend speed for 30 second under argon. The reaction was rested for 30 seconds to a minute until the solution was cleared and the foam formed on top. The foam was excess methionine and was optionally removed.

The solution was immediately (< 2 minutes) poured into an extraction blender and added with 240 ml of mineral oil (STO Oil Corp., San Marcos, TX) at room temperature, then blended at low speed for 20 seconds under argon.

The water-oil mixture was allowed to settle for 3-4 hours, and then the oil phase was removed and filtered through 20-micron filter paper. The filtrate was stored in an open container for 3 days to allow argon to dissipate, and the oil filtrate was ready to be used as a product.

Example 2. Hepatotoxicity Trials

An *in vivo* evaluation for hepatotoxicity of the product of Example 1 was performed on a rat population.

A control group of ten rats were untreated for control analysis of baseline serum enzymes and histology.

A second group of ten rats were gavaged twice daily with a standard saline solution to gauge the effects of repeating handling and gavaging on the liver.

5 A third group of twenty-five rats were gavaged twice daily with mineral oil alone.

A fourth group of twenty-five rats were gavaged twice daily with the product of Example 1.

After 28 days, the rats were anesthetized and their livers were tested according to standard methodology. It was concluded that consumption of the product of Example 1 did not
10 result in any signs of hepatotoxicity.

Example 3. Treatment of infections

The product of Example 1 was used to treat subjects who exhibited various infections.

The results are summarized in Table 1.

15

Table 1

Subject	Condition	Method of Application	Application	Results
Subject 1	Staphylococcal infection of leg	Topical	3 months	Symptoms relieved
Subject 2	Sinus condition	Topical	1 week	Symptoms relieved

Example 4. Treatment of joint and muscle pain

20 Subjects having joint pain, bug bite pain, or burn pain were treated with the product of Example 1. The results are summarized in Table 2.

Table 2

Subject	Condition	Method of Application	Relief Time
Subject 1	Hip joint pain	Topical	Immediate
Subject 3	Arthritic pain	Topical	Immediate
Subject 4	Back/knee pain	Topical	Immediate
Subject 5	Knee pain	Topical	Immediate
Subject 6	Back/knee pain	Topical	Immediate
Subject 7	Hip pain	Topical	Immediate

Subject 8	Arthritic hand	Topical	Immediate
Subject 2	Hand burn pain	Topical	Immediate
Subject 1	Bug bite pain	Topical	Immediate

Table 2 shows that subjects with various pain, when applied with the product topically, resulted in immediate relief of the pain.

5 **Example 5. Treatment of Wounds or Injuries**

The product of Example 1 was applied to individuals having burns or skin conditions topically three times a day as indicated in Table 3. After application of the product, the symptoms of the subjects were relieved.

10

Table 3

Subject	Condition	Method of Application	Application Time
Subject 1	Burns	Topical	3 days
Subject 9	Sun face blotches	Topical	3 months
Subject 10	Purple sun burn	Topical	3 days
Subject 11	Sun damage to skin	Topical	2 months
Subject 12	Sun damage to skin	Topical	2 months
Subject 13	Bleeding facial skin	Topical	2 weeks
Subject 2	Wart	Topical	3 weeks
Subject 14	Diabetic sores of legs and feet	Topical	1 week

15 The invention, and the manner and process of making and using it, are now described in such full, clear, concise and exact terms as to enable any person skilled in the art to which it pertains, to make and use the same. It is to be understood that the foregoing describes preferred embodiments of the present invention and that modifications may be made therein without departing from the scope of the present invention as set forth in the claims. To particularly point out and distinctly claim the subject matter regarded as invention, the following claims conclude the specification.

CLAIMS:

1. An organic solvent extract prepared by the method comprising the steps of:
 - (a) mixing methionine with water,
 - (b) adding an aqueous hypochlorite solution to (a) and mixing at a temperature
5 between 0°C to ambient temperature,
 - (c) adding a water-immiscible organic solvent to (b) and mixing, and
 - (d) separating the organic solvent phase.

2. The organic solvent extract according to Claim 1, wherein said water-immiscible
10 organic solvent is hexane, heptane, methylene chloride, mineral oil, vegetable oil, or ethyl acetate.

3. The organic solvent extract according to Claim 2, wherein said water-immiscible
organic solvent is mineral oil or hexane.
15

4. The organic solvent extract according to any one of Claims 1-3, wherein the steps (a)
and (b) are carried out under inert gas.

5. The organic solvent extract according to any one of Claims 1-4, wherein the
20 methionine is D-methionine, L-methionine, or a mixture thereof.

6. The organic solvent extract according to any one of Claims 1-5, wherein the water-
immiscible organic solvent is ethyl acetate.

- 25 7. Composition comprising the organic solvent extract of any one of Claims 1-6 for use
in the treatment of bacterial infection.

8. Composition comprising the organic solvent extract of any one of Claims 1-6 for use
in the treatment of viral infection.
30

9. Composition comprising the organic solvent extract of any one of Claims 1-6 for use
in the treatment of fungal infection.

10. Composition comprising the organic solvent extract of any one of Claims 1-6 for use in a method of treating pain.

11. Composition comprising the organic solvent extract for use according to Claim 10,
5 wherein said pain is joint pain, muscle pain, arthritic pain, or burn pain.

12. Composition comprising the organic solvent extract of any one of Claims 1-6 for use in a method of treating skin conditions in a subject by topically administering the composition to the afflicted area of skin in a subject.

10

13. Composition comprising the organic solvent extract of any one of Claims 1-6 for use in a method of treating inflammation or inflammatory-related disorders in a subject by administering the composition to a subject suffering from inflammation or inflammatory-related disorders.

15