



(51) International Patent Classification:

A61K 31/27 (2006.01) A61K 36/185 (2006.01)
A61K 31/55 (2006.01) A61K 36/324 (2006.01)
A61K 31/473 (2006.01)

(21) International Application Number:

PCT/US2018/066867

(22) International Filing Date:

20 December 2018 (20.12.2018)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

62/608,435 20 December 2017 (20.12.2017) US

(71) Applicant: **ATTILLAPS HOLDINGS** [US/US]; 1600
Fillmore St., #241, Denver, CO 80206 (US).

(72) Inventor; and

(71) Applicant: **SPALLITTA, Frank, Anthony** [US/US];
1600 Fillmore St., #241, Denver, CO 80206 (US).

(74) Agent: **CHAPMAN, Gary, B.**; Leydig, Voit & Mayer,
Ltd., 4940 Pearl East Circle, Suite 200, Boulder, CO 80301
(US).

(81) Designated States (*unless otherwise indicated, for every
kind of national protection available*): AE, AG, AL, AM,
AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ,
CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO,
DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN,
HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP,
KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME,
MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ,
OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA,
SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN,
TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(54) Title: TREATMENT OF OPHTHALMOLOGICAL CONDITIONS WITH ACETYLCHOLINESTERASE INHIBITORS

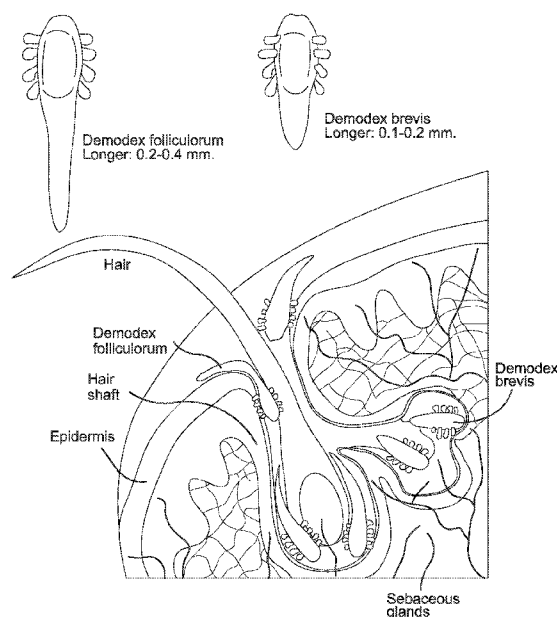


FIG. 2

(57) **Abstract:** Embodiments of the invention involve treating ophthalmology conditions by the topical or oral use of acetylcholinesterase inhibitors. By effectively reducing or eliminating the population of demodex mites in affected areas and areas where demodex mites may exist, this treatment achieves a more complete remission of clinical signs and symptoms of the ophthalmological afflictions than any previously described method. Embodiments of the invention are useful for treating ocular afflictions caused by demodex-induced inflammatory eye conditions, including meibomian gland dysfunction, conjunctivitis, keratoconjunctivitis, hyperemia, blepharitis and dry eye disease.



(84) Designated States (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

— *with international search report (Art. 21(3))*

TREATMENT OF OPHTHALMOLOGICAL CONDITIONS WITH ACETYLCHOLINESTERASE INHIBITORS

[0001] This application claims the benefit of priority to U.S. Provisional Patent Application No. 62/608,435 filed on December 20, 2017. This application is a
5 continuation-in-part application of U.S. Patent Application Serial Nos. 15/319,897 filed December 19, 2016 (86-14 US) and 14/444,748 (41-14 US) filed July 28, 2014. 15/319,897 is a national stage application of PCT/US15/36448 filed June 18, 2015 which claims the benefit of U.S. Provisional App. No. 62/014,520 filed June 19, 2014. 14/444,748 claims the benefit of U.S. Provisional App. Nos. 61/859,572 filed July 29,
10 2013, 61/861,072 filed August 1, 2013 and 61/953,290 filed March 14, 2014. Each of these applications are hereby incorporated by reference in their entireties to the extent they are not inconsistent with the disclosure herein.

BACKGROUND

[0002] Provided herein are methods for treatment of various ophthalmological
15 afflictions in humans employing topically applied or orally dosed acetylcholinesterase inhibitors and/or carbamates, such as ethyl carbamates to inactivate certain organisms associated with the ophthalmological affliction. For example, by reducing or eliminating demodex organisms from affected hair follicles, skin, eyes, eyelids, eyelashes, or meibomian gland areas, the methods reduce clinical signs of the
20 ophthalmological afflictions which are primarily due to allergic and inflammatory responses of the body to the demodex organisms and bacteria that are associated or carried by the demodex organism.

[0003] Eyelid involvement may be manifested by mild conjunctival irritation or inflammation of the meibomian (oil) glands on the eyelid margin. Chronic eyelid
25 irritation can result in loss of eyelashes. Meibomian gland dysfunction caused by the irritation and inflammation can lead to chronic dry eye disease and/or blepharitis.

[0004] Although hypothesized as a root cause of dry eye associated with Meibomian gland dysfunction, demodex folliculorum has yet to reach consensus and no commercially viable pharmacological solutions are available for treating demodex
30 brevis and demodex folliculorum in the eyelid. Reaction to the presence or metabolic activity of demodex mites in eyelash follicles has been discussed as a cause of

blepharitis but previous studies where topical miticides other than acetylcholinesterase inhibitors have been used have shown inconsistent and marginal results. Demodex seems to play a role in meibomian gland dysfunction which can eventually cascade into chronic dry eye disease.

5 **[0005]** The etiology of dry eye disease is becoming increasingly well understood. Dry eye disease is progressive in nature, and fundamentally results from insufficient tear coverage on the surface of the eye. This poor tear coverage prevents healthy gas exchange and nutrient transport for the ocular surface, promotes cellular desiccation and creates a poor refractive surface for vision. Poor tear coverage
10 typically results from: excessive evaporation of aqueous tear resulting from dysfunction of the meibomian glands. The meibomian gland at the rim of the eyelids inside the tarsal plate are responsible for the supply of meibum, an oily substance that prevents evaporation of the eye's tear film. If the flow of meibum is blocked by demodex (or indirectly by an inflammatory response to demodex), cylindrical dandruff
15 caused by demodex, or inflammation resulting from demodex or bacteria proliferated by demodex, a lack of meibum causes a cascading effect that leads to chronic dry eye disease. Low tear volume causes a hyperosmolar environment that induces an inflamed state of the ocular surface. This inflammatory response induces apoptosis of the surface cells which in turn prevents proper distribution of the tear film on the
20 ocular surface so that any given tear volume is rendered less effective. This initiates a vicious positive feedback loop cycle where more inflammation can ensue causing more surface cell damage, etc. Additionally, the neural control loop, which controls reflex tear activation, is disrupted because the sensory neurons in the surface of the eye are damaged. As a result, fewer tears are secreted and a second vicious cycle
25 develops that results in further progression of the disease because fewer tears cause nerve cell loss, which results in even fewer tears, etc.

[0006] There is a wide spectrum of treatments for dry eye disease, however, none provides substantial efficacy for treatment of the condition. Treatment options include: artificial tear substitutes, ointments, gels, warm compresses, environmental
30 modification, topical cyclosporine, omega-3 fatty acid supplements, punctal plugs and moisture chamber goggles. Patients with severe disease may further be treated with punctal cautery, systemic cholinergic agonists, systemic anti-inflammatory

agents, mucolytic agents, autologous serum tears, PROSE scleral contact lenses and tarsorrhaphy. Despite these treatment options, dry eye disease continues to be considered one of the most poorly treated diseases in ophthalmology. Accordingly, it would be desirable to have a more effective treatment for dry eye.

5 **[0007]** Conventionally, dry eye is treated with the frequent use of artificial tears or other lubricating eye drops. Artificial tears usually are the first step in dry eye treatment. RESTASIS® eye drops are the most commonly prescribed pharmaceutical intervention for dry eye (RESTASIS® (Cyclosporine Ophthalmic Emulsion) 0.05%) and contains cyclosporine which is an immunosuppressant.

10 Cyclosporine can increase tear production that has been reduced by inflammation in the eye(s). RESTASIS® eye drops are used to treat chronic dry eye that may be caused by inflammation. Ocular rosacea is characterized by ocular manifestations such as dry eye, tearing and burning, swollen eyelids, recurrent styes and potential vision loss from corneal damage. There is a need in the art for safe and effective
15 treatment of ophthalmological conditions, specifically demodex-induced inflammatory eye conditions such as Meibomian gland dysfunction, conjunctivitis, keratoconjunctivitis, hyperemia, dry eye, and blepharitis. Accordingly, provided herein are various treatments that specifically target this demodex-induced mechanism.

20 SUMMARY OF THE INVENTION

[0008] Provided herein are treatment methods for demodex-induced inflammatory eye conditions, including that alleviate, abrogate, or otherwise reduce or stop any one or more clinical symptoms associated with demodex-induced inflammatory eye conditions by administering or applying specific acetylcholinesterase inhibitors.

25 **[0009]** Embodiments of the invention described herein involve treating ophthalmological afflictions by the topical, intraocular or oral use of one or more than one specified acetylcholinesterase inhibitors. By effectively reducing or eliminating the population of Demodex mites in affected areas and areas where demodex mites may exist, this treatment achieves a more complete remission of clinical signs and
30 symptoms of the ophthalmological afflictions than any previously described method. Embodiments of the invention are useful for treating ophthalmological afflictions related to demodex-induced inflammatory eye conditions, including meibomian gland

dysfunction, conjunctivitis, keratoconjunctivitis, hyperemia (excess blood supply to the eye), blepharitis and dry eye disease.

[0010] An exemplary method embodiment comprises a step of orally-administering or topically-applying to an individual having the ophthalmological affliction, including demodex-induced inflammatory eye conditions, an acetylcholinesterase inhibitor in a dosage sufficient to inactivate demodex brevis and/or demodex folliculorum mites from hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual, resulting in amelioration or cessation of the manifestations of allergic and/or inflammatory responses to the mites that cause symptoms and signs of the ophthalmological affliction in the individual. In an embodiment, the acetylcholinesterase inhibitor is not an organophosphate, including acetylcholinesterase inhibitors and organophosphates described in WO 2015/017328 and WO 2015/195928 including, but not limited to, dichlorvos, metrifonate, malathion, isopropyl fluorophosphate and echothiophate. In an exemplary embodiment, the acetylcholinesterase inhibitor is topically applied. In an embodiment, for example, the topically-applied acetylcholinesterase inhibitor is formulated in a saline solution, carrier lotion, cream, soap, wash, shampoo, gel, impregnated wipe, swab or via a spoolie brush. Preferably, the administration is around the eye and regions adjacent thereto, but not directly into the eye.

[0011] Optionally, a concentration of the acetylcholinesterase inhibitor in the topically-applied lotion, cream, soap, wash, shampoo, saline solution or gel is about 0.001 to 1 percent by weight or about 0.01 to 1 percent by weight. In an exemplary embodiment, a concentration of the acetylcholinesterase inhibitor in the topically-applied saline solution, lotion, cream, soap, wash, shampoo or gel is a lowest concentration effective for killing the demodex mites. In one embodiment, a dosage of acetylcholinesterase inhibitor in the topically-applied lotion, cream, soap, wash, shampoo, saline solution or gel is less than about 150 mg/kg of body mass or between about 0.01 mg per kg of body mass and 50 mg/kg of body mass. In an exemplary embodiment, a dosage of acetylcholinesterase inhibitor in the topically-applied saline solution, lotion, cream, soap, wash, shampoo or gel is a lowest dose effective for killing the demodex mites. Optionally, the topically-applied acetylcholinesterase inhibitor is encapsulated inside microliposomes before being

formulated into the carrier saline solution, lotion, cream, soap, wash, shampoo or gel.

[0012] In general, methods of the invention include those where the topically-applied acetylcholinesterase inhibitor is applied to said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands areas affected by the ophthalmological affliction. In certain embodiments, however, the topically-applied acetylcholinesterase inhibitor is further applied to areas not affected by the ophthalmological affliction. For example, in one embodiment, the topically-applied acetylcholinesterase inhibitor is applied to the hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands areas of the body where demodex brevis and/or demodex folliculorum mites exist. In an exemplary embodiment, the topically-applied acetylcholinesterase inhibitor is applied to all areas.

[0013] Optionally, methods of the invention further comprise a step of applying the acetylcholinesterase inhibitor to the individual's clothing, linens or both clothing and linens. Such application is useful, for example, for preventing the individual's clothing or linens from being a source of demodex mites to reintroduce onto the individual's skin. Similarly, methods of the invention optionally further comprise a step of orally- administering or topically-applying the acetylcholinesterase inhibitor to others having contact with the individual in a dosage sufficient to fill and eliminate demodex brevis and/or demodex folliculorum mites from hair follicles and/or skin of the others. For example, in embodiments, the others comprise household members, children, spouses, partners, family members or domestic pets.

[0014] In an exemplary embodiment of the methods of the invention, the topically-applied acetylcholinesterase inhibitor is applied to the hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual. In one embodiment, for example, the topically-applied acetylcholinesterase inhibitor penetrates an outer layer of the skin of the individual, thereby exposing the demodex brevis and/or demodex folliculorum mites present below the outer layer of the skin in the eyelid to the acetylcholinesterase inhibitor. In one embodiment, for example, the topically-applied acetylcholinesterase inhibitor penetrates to a subdermal region of the eyelid of the individual, thereby exposing the demodex brevis and/or demodex folliculorum mites present in and around the meibomian glands to the acetylcholinesterase

inhibitor. Certain formulations of the topical acetylcholinesterase inhibitor useful with the methods of the invention optionally comprise one or more compositions that increase a permeability of the skin, such as dimethyl sulfoxide (DMSO).

[0015] In an exemplary embodiment, the topically-applied acetylcholinesterase inhibitor is applied to affected skin areas at least once and not more than twice daily for a period of about two to twelve weeks. In one embodiment, the topically-applied acetylcholinesterase inhibitor is applied to the affected areas and/or to non-affected areas during a first application period, thereby killing and eliminating adult demodex brevis and/or demodex folliculorum mites from the hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands. In one embodiment, the topically-applied acetylcholinesterase inhibitor is further applied to the affected areas and/or to non-affected areas during a second application period, thereby killing and eliminating from the hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual demodex brevis and/or demodex folliculorum mites that have matured from a larval form and/or an egg form present on and/or in the skin during the first application period. In one embodiment, the topically-applied acetylcholinesterase inhibitor is further applied to the affected areas and/or to non-affected areas during a third application period, thereby killing and eliminating from the hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual demodex brevis and/or demodex folliculorum mites that have matured from a larval form and/or an egg form present on and/or in the skin and/or the hair follicles during the first application period and/or the second application period.

[0016] Optionally, the first application period and the second application period are separated by at least five but no more than ten days. Optionally, the first application period and the second application period are separated by at least seven days. In an exemplary embodiment, the first application period and the second application period are separated by a time sufficient to allow the larva form to mature into an adult form and/or to allow the egg form to mature into the adult form.

[0017] Optionally, the second application period and the third application period are separated by at least five but no more than ten days. Optionally, the second application period and the third application period are separated by at least seven days. In an exemplary embodiment, the second application period and the third

application period are separated by a time sufficient to allow the larva form to mature into an adult form and/or to allow the egg form to mature into the adult form.

[0018] In exemplary embodiments, the acetylcholinesterase inhibitor is orally-administered or topically-applied in a continued intermittent regime sufficient for
5 prophylactic control of demodex mite population in the hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual.

[0019] In another embodiment, the acetylcholinesterase inhibitor is orally-administered. In a specific embodiment, for example, the orally-administered acetylcholinesterase inhibitor is administered as an oral dose of the
10 acetylcholinesterase inhibitor of about 1 50 mg per kg of body mass or less or between about 0.01 mg per kg of body mass and 50 mg per kg of body mass. In an exemplary embodiment, the orally-administered acetylcholinesterase inhibitor is administered as an oral dose of the acetylcholinesterase inhibitor of a lowest dose effective for killing the demodex mites. In certain embodiments, the orally-
15 administered acetylcholinesterase inhibitor is formulated as a prodrug or pharmaceutically acceptable salt.

[0020] Optionally, the orally-administered acetylcholinesterase inhibitor is administered as a daily dose of 10 mg per kg of body mass. Optionally, the orally-administered acetylcholinesterase inhibitor is administered as a daily dose of 7.5 mg
20 per kg of body mass. Optionally, the orally-administered acetylcholinesterase inhibitor is administered as a three times per day dose of 5 mg per kg of body mass. Optionally, the orally-administered acetylcholinesterase inhibitor is repeated about two to four times with spacing of three to seven days between them. In embodiments where the administration is confined to a local region around the eye, lower doses
25 may be used.

[0021] In various embodiments, the elimination of the demodex brevis and/or demodex folliculorum mites from hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual results in a reduction in population of one or more bacteria in the eyes, eyelids, eyelashes of the individual. For example, in some
30 embodiments, the allergic and/or inflammatory responses to the mites result from a presence of one or more bacteria associated with the mites in the hair follicles, skin,

eyes, eyelids, eyelashes, or meibomian glands of the individual. In specific embodiments, the one or more bacteria comprise one or more bacteria from the genus staphylococcus or from the genus bacillus. For example, in one embodiment, the one or more bacteria comprise bacillus oleronius bacteria. In one embodiment, for example, the one or more bacteria comprise staphylococcus epidermidis bacteria. Optionally, the one or more bacteria are present in a digestive system of the demodex brevis and/or demodex folliculorum mites.

[0022] Another exemplary method for treating an ophthalmological affliction comprises a step of topically-applying to an individual having the ophthalmological affliction an active ingredient in a dosage sufficient to fill and eliminate demodex brevis and/or demodex folliculorum mites from hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual, resulting in cessation of the manifestations of allergic and/or inflammatory responses to the mites that cause symptoms and signs of the ophthalmological affliction in the individual, wherein the topically-applied active ingredient is applied to hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands areas affected by the ophthalmological affliction and to skin areas not affected by the ophthalmological affliction. In a specific embodiment, the topically-applied active ingredient is applied to all facial skin of the individual, thereby filling and eliminating the demodex brevis and/or demodex folliculorum mites from all facial skin of the individual. Again, methods of the invention are useful, for example, for treating ocular conditions including meibomian gland dysfunction, blepharitis and dry eye disease. In an exemplary embodiment, the ophthalmological condition is caused by, exacerbated by or otherwise comorbid with an infestation of the skin and/or hair follicles by demodex mites.

[0023] In an aspect, the acetylcholinesterase inhibitor is a reversible inhibitor. Compounds that are reversible competitive or noncompetitive inhibitors of cholinesterase include those having therapeutic uses, including: Carbamates, Physostigmin, Neostigmine, Pyridostigmine, Ambenonium, Demecarium, Rivastigmine, Phenanthrene derivatives, Galantamine, Caffeine - noncompetitive (also an Adenosine receptor antagonist)[13][14], Piperidines, Donepezil, Tacrine, also known as tetrahydroaminoacridine (THA'), Edrophonium, Huperzine A[15][16], Ladostigil, Ungeremine[17], and Lactucopicrin.

[0024] In an aspect, the acetylcholinesterase inhibitor is a quasi-reversible inhibitor. Compounds which function as quasi-irreversible inhibitors of cholinesterase tend to have use as pesticides. These include organophosphates and carbamates.

Examples of organophosphates include: Echothiophate, Diisopropyl

- 5 fluorophosphates, Cadusafos, Chlorpyrifos, Dichlorvos, Dimethoate, Metrifonate (irreversible), Malathion and Parathion. Examples of carbamates include: Aldicarb; Bendiocarb; Bufencarb; Carbaryl; Car bendazim; Carbetamide; Carbofuran ; Carbosulfan ; Chlorbufam; Chlorpropham; Ethiofencarb; Formetanate; Methiocarb; Methomyl ; Oxamyl ; Phenmedipham; Pinmicarb; Pirimicarb; Propamocarb;
- 10 Propham; Propoxur; Huperzine A; Galantamine; Onchidal ; Coumarins.

[0025] In another embodiment, the acetylcholinesterase inhibitor corresponds to a compound currently used in medicine, including those having an established safety profile in humans. Examples include: Aricept; Aricept ODT; Cognex; donepezil; Exelon; galantamine; Namzaric; Razadyne; rivastigmine; tacrine; phospholine;

15 neostigmine; parathion ; malathion; dyflos; physostigmine; endrophonium; pyridostigmine; ecothiophate.

[0026] To further reduce the possibility of unwanted side effects, provided herein are methods that confine the amount and location of the acetylcholinesterase inhibitor to regions that provide the most benefit. In the case of the ophthalmological

20 treatments provided herein, the application may be a topical administration to an eye region of the individual. Optionally, the eye region corresponds to an area of between at least 1 cm and 10 cm around an outermost perimeter defined by the eye, eyelid, eyebrow and eyelashes. In this manner, the inhibitor is confined to a region around the eye, thereby effectively treating the Meibomian glands, and mite-

25 containing regions adjacent thereto that would otherwise tend to result in a relatively quick mite recovery/migration, while avoiding direct application to the eye.

[0027] Also provided herein is a method of treating an individual having a demodex-induced ophthalmological affliction of Meibomian gland dysfunction, the method comprising the steps of: applying to the individual in need thereof an

30 acetylcholinesterase inhibitor in a dosage sufficient to inactivate demodex brevis mites and/or demodex folliculorum mites from hair follicles, eyelids, eyelashes and Meibomian glands in and around an eye region; wherein a sufficient amount of

demodex brevis mites and/or demodex folliculorum mites are inactivated to ameliorate or cease manifestations of allergic and/or inflammatory responses to the mites that cause symptoms or signs of Meibomian gland dysfunction. Although some variation in the precise amount of inactivation is tolerated by the instant methods, inactivation may refer to at least 50%, at least 70%, at least 90%, or at least 95% of the relevant mite population that is killed, unable to mature, and/or unable to effectively reproduce.

[0028] The allergic and/or inflammatory responses to demodex may be manifested as Meibomian gland dysfunction, conjunctivitis, keratoconjunctivitis, hyperemia dry eye or blepharitis, wherein administration of an acetylcholinesterase inhibitor to inactivate demodex brevis and/or demodex folliculorum mites in an eye region reduces or eliminates allergic and/or inflammatory responses to the mites that cause symptoms and signs of the ophthalmological affliction in the individual.

[0029] The application may be a topical administration to an eye region of the individual. The "eye region" may be quantifiably defined as corresponding to an area of between at least 1 cm and 10 cm around an outermost perimeter defined by the eye, eyelid, eyebrow and eyelashes, and any subranges thereof.

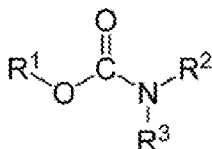
STATEMENTS REGARDING CHEMICAL COMPOUNDS AND NOMENCLATURE

[0030] In an embodiment, a composition or compound used with the methods of the invention is isolated or purified. In an embodiment, an isolated or purified compound is at least partially isolated or purified as would be understood in the art. In an embodiment, the composition or compound of the invention has a chemical purity of 95%, optionally for some applications 99%, optionally for some applications 99.9%, optionally for some applications 99.99%, and optionally for some applications 99.999% pure.

[0031] Many of the compounds used in the methods of the invention contain one or more ionizable groups, ionizable groups include groups from which a proton can be removed (e.g., -COOH) or added (e.g., amines) and groups which can be quaternized (e.g., amines). All possible ionic forms of such molecules and salts thereof are intended to be included individually in the disclosure herein. With regard to salts of the compounds herein, one of ordinary skill in the art can select from

among a wide variety of available counterions that are appropriate for preparation of salts of this invention for a given application. In specific applications, the selection of a given anion or cation for preparation of a salt can result in increased or decreased solubility of that salt.

- 5 **[0032]** The term "carbamate" generally refers to an organic compound derived from carbamic acid (NH₂COOH), such as NR₂R₃COOR₁:



- [0033]** In an aspect, each of the groups R1-R3 are independently selected to correspond to any of the R groups of the chemicals listed herein. In an aspect, any of
- 10 R1-R3 are hydrogen.

- [0034]** Examples of carbamates for use with the methods described herein include, but are not limited to, neostigmine, rivastigmine, meprobamate, carisoprodol, felbamate, tybamate. Preferred carabamates are those that have been demonstrated to have miticidal or insecticidal capabilities and that can be provided to
- 15 a mite on or in hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands at a level sufficient to inactivate or kill the mite without permanently adversely affecting the host patient. The carbamate may be a naturally occurring compound, such as a purified and isolated naturally occurring compound. Alternatively, the carbamate may be a synthetically produced carbamate, as known in the art. Any of the compounds
- 20 provided herein may be provided in the form of a derivative, prodrug, or a pharmaceutically acceptable salt thereof.

- [0035]** In an aspect, the carbamate is selected from the group consisting of: aldicarb, bendiocarb, bufencarb, carbaryl, carbendazim, carbetamide, carbofuran, carbosulfan, chlorbufam, chlorpropham, ethiofencarb, formetanate, methiocarb,
- 25 methomyl, oxamyl, phenmedipham, pinmicarb, pirimicarb, propamocarb, propham, propoxur, butocarboxim, carbanolate, promacyl, thiocarboxime, thiofanox, benomyl, and metolcarb or a derivative, prodrug or pharmaceutically acceptable salt thereof.

[0036] In an aspect, the carbamate is an ethyl carbamate of the form = ethyl. R2 and R3 are optionally independently selected as hydrogen.

[0037] The compounds used in the methods of this invention can contain one or more chiral centers. Accordingly, this invention is intended to include racemic mixtures, diastereomers, enantiomers, tautomers and mixtures enriched in one or more stereoisomer. The scope of the invention as described and claimed encompasses the racemic forms of the compounds as well as the individual enantiomers and non-racemic mixtures thereof.

[0038] Pharmaceutically acceptable salts comprise pharmaceutically-acceptable anions and/or cations. As used herein, the term "pharmaceutically acceptable salt" can refer to acid addition salts or base addition salts of the compounds in the present disclosure. A pharmaceutically acceptable salt is any salt which retains at least a portion of the activity of the parent compound and does not impart significant deleterious or undesirable effect on a subject to whom it is administered and in the context in which it is administered. Pharmaceutically acceptable salts include metal complexes and salts of both inorganic and organic acids. Pharmaceutically acceptable salts include metal salts such as aluminum, calcium, iron, magnesium, manganese and complex salts. Pharmaceutically acceptable salts include, but are not limited to, acid salts such as acetic, aspartic, alkylsulfonic, arylsulfonic, axetil, benzenesulfonic, benzoic, bicarbonic, bisulfuric, bitartaric, butyric, calcium edetate, camsylic, carbonic, chlorobenzoic, -32- cilexetil, citric, edetic, edisylic, estolic, esyl, esylic, formic, fumaric, gluceptic, gluconic, glutamic, glycolic, glycolylarsanilic, hexamic, hexylresorcinoic, hydrabamic, hydrobromic, hydrochloric, hydroiodic, hydroxynaphthoic, isethionic, lactic, lactobionic, maleic, malic, malonic, mandelic, methanesulfonic, methylnitric, methylsulfuric, mucic, muconic, napsylic, nitric, oxalic, p-nitromethanesulfonic, pamoic, pantothenic, phosphoric, monohydrogen phosphoric, dihydrogen phosphoric, phthalic, polygalactouronic, propionic, salicylic, stearic, succinic, sulfamic, sulfanilic, sulfonic, sulfuric, tannic, tartaric, teoclic, toluenesulfonic, and the like. Pharmaceutically acceptable salts may be derived from amino acids, including but not limited to cysteine. Other pharmaceutically acceptable salts may be found, for example, in Stahl et al., Handbook of Pharmaceutical Salts: Properties, Selection, and Use, Wiley-VCH; Verlag Helvetica Chimica Acta, Zurich,

2002. (ISBN 3-906390-26-8). Pharmaceutically- acceptable cations include among others, alkali metal cations (e.g., Li^+ , Na^+ , K^+), alkaline earth metal cations (e.g., Ca^{2+} , Mg^{2+}), non-toxic heavy metal cations and ammonium (NH_4^+) and substituted ammonium (N(R')_4^+ , where R' is hydrogen, alkyl, or substituted alkyl, i.e., including, methyl, ethyl, or hydroxyethyl, specifically, trimethyl ammonium, triethyl ammonium, and triethanol ammonium cations). Pharmaceutically- acceptable anions include among other halides (e.g., Cl^- , Br^-), sulfate, acetates (e.g., acetate, trifluoroacetate), ascorbates, aspartates, benzoates, citrates, and lactate.

[0039] Also provided herein are methods of treating an individual having a demodex-induced inflammatory eye condition. The method may comprise the steps of: applying to the individual in need thereof an acetylcholinesterase inhibitor in a dosage sufficient to inactivate demodex brevis mites and/or demodex folliculorum mites from an eye region of the individual, wherein the eye region includes one or more of hair follicles, eyelids, eyelashes and Meibomian glands in and around an eye region, including such that a sufficient amount of demodex brevis mites and/or demodex folliculorum mites are inactivated to ameliorate or cease manifestations of allergic and/or inflammatory responses to the demodex mites that cause the inflammatory eye condition. This sufficient amount, depending on the sensitivity of the individual, may correspond to an at least 50%, 70%, 90% or 95% elimination, including in and around the eye region and/or a whole body decrease.

[0040] The inflammatory eye condition may correspond to one or more of: Meibomian gland dysfunction, dry eye, blepharitis, conjunctivitis, keratoconjunctivitis, or hyperemia.

[0041] For any of the applications described herein, the application may be a topical administration to the eye region of the individual, preferably avoiding application directly to the eye surface.

[0042] The eye region may correspond to an area of between at least 1 cm and 10 cm around an outermost perimeter defined by the eye, eyelid, eyebrow and eyelashes.

[0043] Without wishing to be bound by any particular theory, there may be discussion herein of beliefs or understandings of underlying principles relating to the

devices and methods disclosed herein. It is recognized that regardless of the ultimate correctness of any mechanistic explanation or hypothesis, an embodiment of the invention can nonetheless be operative and useful.

BRIEF DESCRIPTION OF THE DRAWINGS

5 [0044] FIG. 1 summarizes a topical administration method to an eye region of an individual in need of treatment.

[0045] FIG. 2 is an illustration of *Demodex folliculorum* and *Demodex brevis* mites, including relative sizes and locations in the skin, hair follicles and glands.

10 [0046] FIG. 3 illustrates *Demodex* mites in the eye region, including eye lash and meibomian glands.

[0047] FIG. 4 is a table summary of *Demodex* survival time for various acetylcholinesterase inhibitors.

DETAILED DESCRIPTION OF THE INVENTION

15 [0048] In general, the terms and phrases used herein have their art-recognized meaning, which can be found by reference to standard texts, journal references and contexts known to those skilled in the art. The following definitions are provided to clarify their specific use in the context of the invention.

[0049] "Inactivate" is used broadly herein to refer to the functional ability to decrease the impact of *demodex brevis* and/or *demodex folliculorum* mites. For example, the inactivation may be by death of the mite. Alternatively, the inactivation may refer to the inability of the mite to reproduce, so that the mite die off occurs as the mites age and die without reproduction. So long as the treatment leads to an adverse effect on the *demodex brevis* and/or *demodex folliculorum* mites that corresponds to improved clinical outcome, such as symptom improvement, the treatment is considered herein to inactivate *demodex brevis* and/or *demodex folliculorum* mites.

25

[0050] "Efficiently transported" refers to the ability of the treatment agent to act against mites that are located beneath the skin surface, such as into an epidermal or subdermal region so that the mites are timely inactivated.

[0051] “Substantially all” refers to, unless defined in the contrary, at least 90%, at least 95% or at least 99% of the relevant population, so in the context of demodex mites, it refers to inactivation (e.g., killed or eliminated or otherwise unable to propagate) and/or application to hair (number) or skin (surface area).

5 **[0052]** As used herein, “Demodex” refers to *D. folliculorum* and *D. brevis* mites, including *Demodex* mites in humans that may contribute to a demodex-induced inflammatory eye condition in humans.

[0053] *D. folliculorum* and *d. brevis* mites may play a role in ophthalmological conditions. An increased demodex population has been observed in patients with
10 ophthalmological afflictions. For most people, demodex mites live harmlessly in the hair follicles, skin, eyelids, eyelashes, or meibomian glands as a result of either down-regulating host immunity or simply dodging host immune defenses. There is continual debate within the ophthalmology community as to whether or not they are the causative agents of such ophthalmological diseases as meibomian gland
15 dysfunction, dry eye disease and blepharitis (inflammation of the eyelids).

[0054] Human beings are the one and only host of these ubiquitous mites [1]. In fact, these two mites are considered to be the most common ectoparasite of humans [6]. Women tend to have a higher rate of demodex infections [5]. The rate of infestation also seems to be correlated with age, with 84% of people at age 60
20 harboring mites and increasing to 100% in those 70 years and older [7]. Whether those that are immunocompromised are more susceptible to higher infestation rates is unknown, though some studies indicate that AIDs and leukemia patients may be more prone to greater than average numbers [5].

[0055] The mites are most commonly found in the scalp, face and upper chest
25 area, with *D. folliculorum* exhibiting a predilection for the hair follicles and *D. brevis* for the sebaceous ducts and meibomian glands at the rim of the eyelids (the sebaceous ducts transfer the waxy sebum that lubricates the skin and hair from the sebum glands; the meibonmian glands are a special type of such gland) [4][5]. *D. folliculorum* are a communal bunch, tending to congregate in the follicle area of the
30 hair or eyelashes with their posterior ends protruding from the follicular pores. *D. brevis*, on the other hand, tend to be more solitary and will occupy the sebaceous

glands singly [6]. Both species are tiny, less than 0.4 mm, with elongated, clear bodies and four pairs of stout legs. *D. brevis* is usually a tad shorter, ~ 0.1 mm, than *D. folliculorum*. They both have ridged scales along their cephalothorax and sharp, piercing teeth [6].

5 **[0056]** Short-lived creatures, a mite's life cycle from egg to larva to adult lasts from 14-18 days. Adults emerge from the follicles and ducts to reproduce at the surface of the skin where females will then deposit eggs in the sebaceous glands. Larva will mature via two nymphal stages in the glands until entering the follicles and ducts as adults to begin the cycle anew [6]. It is hypothesized that both species of
10 mites feed upon sebum as a primary food source but may also munch on follicular and glandular epithelia. They are thought to be obligate ectoparasites, incapable of living outside their human host.

[0057] Some studies have discovered a greater than average mite density, greater than five mites per cm², do seem to play a role in skin diseases for patients
15 [6]. Researchers have suggested that blockage of the hair follicles and sebaceous ducts by mites may result in epithelial hyperplasia, elicit a phagocytic, granulomatous reaction or bring about an inflammatory response due to their waste products [5]. The fact that treatment with certain antibiotics can reduce the severity of meibomian gland dysfunction, blepharitis, dry eye disease strongly suggests a microbial
20 component to mite-related diseases.

[0058] In 2007, researchers isolated from *D. folliculorum* a bacterium *Bacillus oleronium* that provoked inflammatory responses in 73% of rosacea patients but only 29% of controls [21]. These results suggest that patients with rosacea including
25 ocular rosacea were sensitized to the bacteria and may be immunologically sensitive to the mites, bacteria or both [21].

[0059] Two antigenic proteins found on the bacterium's cell surface in particular appeared to be responsible for the inflammatory response by stimulating peripheral blood mononuclear cell proliferation; one 83 kDa protein showed similarity with heat-shock proteins while the other 62 kDa protein shared amino acid sequence
30 homology with a protease enzyme found to be involved signal transduction as well as carbohydrate metabolism [21]. Stronger proof of the pathogenic role of *B.*

oleroniusin ocular rosacea may also be found in the sensitivity of the bacterium to many antibiotics proven to be effective in the treatment of rosacea, specifically tetracycline, doxycycline and minocycline [21].

[0060] In an exemplary embodiment, an acetylcholinesterase inhibitor is administered topically to a patient with an active ophthalmological condition in which the underlying cause is a demodex mite. Because the target organisms, demodex brevis and demodex folliculorum, are ectoparasites in the mite family, an effective treatment must be capable of eradicating the entire lifecycle of such a microscopic insect, including egg, larval, and adult stages. For this reason, this embodiment treats such patients with several doses. Such spacing allows time for demodex eggs to hatch into immature mites that are killed before they can mature into egg-producing adults. After the acetylcholinesterase inhibitor carries out its miticidal activity on demodex brevis and demodex folliculorum organisms, inflammatory responses to them begin to diminish but remnants of the dead mites still elicit some flushing and lesion formation until the cleanup processes of the body remove them, a process requiring six to twelve weeks. During this initial phase of acetylcholinesterase inhibitor administration, conventional ophthalmological medications such as artificial tear substitutes, ointments, gels, warm compresses, environmental modification, topical cyclosporine, omega-3 fatty acid supplements, punctal plugs and moisture chamber goggles can be utilized in combination treatment with acetylcholinesterase inhibitor compounds. Patients with severe disease may further be treated with punctal cautery, systemic cholinergic agonists, systemic anti-inflammatory agents, mucolytic agents, autologous serum tears, PROSE scleral contact lenses and tarsorrhaphy can optionally be employed to suppress early flareups and to give early clinical response. No such medications are needed to treat manifestations of the ophthalmological condition after six to twelve weeks have elapsed. After prolonged intervals of freedom from symptoms, should classic signs begin to reappear, treatment can be repeated. The acetylcholinesterase inhibitor is formulated into a cosmetically-acceptable topical saline solution, lotion, cream, shampoo, or gel and applied especially to hair follicles, skin, eyes, eyelids, eyelashes, surrounding the eyes and any area possibly inhabited by demodex brevis and demodex folliculorum. Because of the well-known barrier effect the skin presents to the penetration of topical medications, such a route of

treatment with acetylcholinesterase inhibitor is anticipated to require once or twice daily applications for as long as twelve to six weeks to achieve sufficient follicle penetration and effective miticidal activity. A topical formulation that could achieve this effect would contain about 5% or less of the acetylcholinesterase inhibitor. The lesser the percentage of the acetylcholinesterase inhibitor that can be used while still receiving the miticidal effect and successfully treating the ocular condition is ideal for limiting any possible side effects of the chemical. Further, full facial treatment is optionally useful for preventing reintroduction of the mites onto facial skin and glands.

10 **[0061]** Referring to **FIG. 1**, a method of treating an individual **10** having a demodex-induced ophthalmological affliction of Meibomian gland dysfunction, the method comprising the steps of: applying to the individual in need thereof an acetylcholinesterase inhibitor (indicated by arrow **20**) in a dosage sufficient to inactivate demodex brevis mites and/or demodex folliculorum mites from hair follicles
15 **100**, eyelids **110**, eyebrows **120**, eyelashes **130** and Meibomian glands **140** in and around an eye region. In this manner, a sufficient amount of demodex brevis mites and/or demodex folliculorum mites are inactivated to ameliorate or cease manifestations of allergic and/or inflammatory responses to the mites that cause symptoms or signs of Meibomian gland dysfunction.

20 **[0062]** The allergic and/or inflammatory responses may be manifested by dry eye or blepharitis, and the symptoms related thereto. The application may be by a topical administration of acetylcholinesterase inhibitor **20** to an eye region **205** of the individual **10**. Optionally, the eye region may be quantifiably defined, such as corresponding to an area of the eye region **205**, including a distance **210**, such as
25 between at least 1 cm and 10 cm around an outermost perimeter **200** defined by the eye, eyelid, eyebrow and eyelashes. For clarity, the eye region **205** is illustrated as rectangular in shape, but the invention provided herein is compatible with any shape and region size, although it is preferable to control mite population in the area around the glands **140** so as to avoid relatively fast natural mite migration back to the glands
30 **140** (or region adjacent thereto) with corresponding manifestation of unwanted clinical system. Accordingly, the methods provided herein, upon completion, may be

characterized as ameliorating or cessation of symptoms and signs of the ophthalmological affliction of at least 30 days – 6 months, and any range therein.

[0063] **FIGs 2-3** illustrate the *demodex brevis* and *folliculorum* mites in the skin area generally and the eye region, respectively, demonstrating the need to ensure treatment applications are able to achieve sub-dermal penetration.

Medical use of Carbamates:

[0064] Urethane (ethyl carbamate) was once produced commercially in the United States as an antineoplastic agent and for other medicinal purposes. It was found to be toxic and largely ineffective. It is occasionally used as a veterinary medicine.

[0065] In addition, some carbamates are used in human pharmacotherapy, for example, the cholinesterase inhibitors neostigmine and rivastigmine, whose chemical structure is based on the natural alkaloid physostigmine. Other examples are meprobamate and its derivatives like carisoprodol, felbamate, and tybamate, a class of anxiolytic and muscle relaxant drugs widely used in the 60s before the rise of benzodiazepines, and still used nowadays in some cases.

[0066] The cholinesterase inhibitors neostigmine and rivastigmine may be efficacious if they have similar miticidal capabilities compared to many other carbamate compounds.

[0067] Drug class and mechanism: Rivastigmine is an oral medication used to treat patients with Alzheimer's disease. Rivastigmine is in a class of drugs called cholinesterase inhibitors that also includes tacrine (Cognex), donepezil (Aricept), and galantamine (Razadyne - formerly known as Reminyl). Cholinesterase inhibitors inhibit (block) the action of acetylcholinesterase, the enzyme responsible for the destruction of acetylcholine. Acetylcholine is one of several neurotransmitters in the brain, chemicals that nerve cells use to communicate with one another. Reduced levels of acetylcholine in the brain are believed to be responsible for some of the symptoms of Alzheimer's disease.

[0068] In vitro efficacy and mite lifetimes: Relative efficacy of various formulations are evaluated with a demodex survival time experiments. **FIG. 4** tabulates demodex

survival time, expressed in terms of LT50 (time at which 50% of demodex mites are killed) and/or average minutes to death for various inhibitors.

We claim:

1. A method of treating an ophthalmological affliction, the method comprising the step of orally-administering or topically-applying to an individual having the ophthalmological affliction an acetylcholinesterase inhibitor in a dosage sufficient to inactivate demodex brevis and/or demodex folliculorum mites from hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual, resulting in amelioration or cessation of the manifestations of allergic and/or inflammatory responses to the mites that cause symptoms and signs of the ophthalmological affliction in the individual.
2. The method of claim 1, wherein said ophthalmological affliction is a demodex-induced inflammatory eye condition.
3. The method of claim 2, wherein said demodex-induced inflammatory eye condition is one or more of: Meibomian gland dysfunction, conjunctivitis, keratoconjunctivitis, hyperemia, dry eye, and blepharitis.
4. The method of claim 1, wherein said acetylcholinesterase inhibitor is a miticide or insecticide.
5. The method of claim 4, wherein said acetylcholinesterase inhibitor is selected from the group consisting of: couramin, boswellia sacra, lactucopicrin, celastus paniculatus, dichlorvos, malathion, metrifonate, physostigmine, galantamine, demacarium, donepezil, tacrine, diisopropyl fluorophosphate, echothiophate and huperzine A.
6. The method of claim 1, wherein said acetylcholinesterase inhibitor is efficiently transported into the epidermis or a subdermal region upon contact with said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual.
7. The method of claim 1, wherein said step of orally-administering or topically-applying said acetylcholinesterase inhibitor kills at least a portion of said

demodex brevis and/or demodex folliculorum mites or renders at least a portion of said demodex brevis and/or demodex folliculorum mites unable to reproduce.

8. The method of claim 1, wherein said step of orally-administering or topically-
5 applying said acetylcholinesterase inhibitor kills or eliminates said demodex brevis and/or demodex folliculorum mites.
9. The method of any of claims 1-8, wherein the ophthalmological affliction is one or more of meibomian gland dysfunction, blepharitis, dry eye disease.
10. The method of any of claims 1-8, wherein the ophthalmological affliction affects the eye or eyelids, or both.
11. The method of claim 1, wherein said acetylcholinesterase inhibitor is a
15 carbamate.
12. The method of claim 11, wherein said carbamate is a miticide or insecticide.
13. The method of claim 11, wherein said carbamate is a naturally occurring
20 compound.
14. The method of claim 11, wherein said carbamate is an ethyl carbamate or a prodrug or pharmaceutically acceptable salt thereof.
- 25 15. The method of claim 11 wherein said carbamate is selected from the group consisting of: aldicarb, bendiocarb, bufencarb, carbaryl, carbendazim, carbetamide, carbofuran, carbosulfan, chlorbufam, chloroprotham, ethiofencarb, formetanate, methiocarb, methomyl, oxamyl, phenmedipham, pinmicarb, pirimicarb, propamocarb, protham, propoxur, butocarboxim,
30 carbanolate, promacyl, thiocarboxime, thiofanox, benomyl, and metolcarb or a derivative, prodrug or pharmaceutically acceptable salt thereof.
16. The method of claim 1, wherein said acetylcholinesterase inhibitor is a naturally occurring acetylcholinesterase inhibitor.

17. The method of claim 16, wherein said naturally occurring acetylcholinesterase inhibitor is a miticide or insecticide.
- 5 18. The method of claim 16, wherein said naturally occurring acetylcholinesterase inhibitor is witch hazel or a substance derived therefrom.
19. The method of claim 16, wherein said naturally occurring acetylcholinesterase inhibitor is Boswellia sacra resin or a substance derived therefrom.
- 10 20. The method of claim 16, wherein said naturally occurring acetylcholinesterase inhibitor is a coumarin or a derivative, prodrug or pharmaceutically acceptable salt thereof.
- 15 21. The method of claim 16, wherein said naturally occurring acetylcholinesterase inhibitor is selected from the group consisting of: huperzine A, galantamine, , coumarins, celastus paniculatus, and boswellia or a derivative, prodrug or pharmaceutically acceptable salt thereof.
- 20 22. The method of any of claims 1-21, wherein the topically-applied acetylcholinesterase inhibitor is formulated in a saline solution, carrier lotion, cream, soap, wash, shampoo or gel.
- 25 23. The method of claim 22, wherein a concentration of the acetylcholinesterase inhibitor in the topically-applied lotion, cream, soap, wash, shampoo or gel is about 0.001 to 5 percent by weight or about 0.01 to 1 percent by weight.
24. The method of claim 22, wherein a concentration of the acetylcholinesterase inhibitor in the topically-applied lotion, cream, soap, wash, shampoo or gel is a
30 lowest concentration effective for killing the demodex mites.
25. The method of claim 22, wherein a dosage of the acetylcholinesterase inhibitor in the topically-applied lotion, cream, soap, wash, shampoo or gel is less than

about 150 mg/kg of body mass or between about 0.01 mg per kg of body mass and 50 mg/kg of body mass.

26. The method of claim 22, wherein a dosage of the acetylcholinesterase inhibitor
5 in the topically-applied lotion, cream, soap, wash, shampoo or gel is a lowest dose effective for killing the demodex mites.

27. The method of claim 22, wherein the topically-applied acetylcholinesterase inhibitor is encapsulated inside microliposomes before being formulated into the
10 carrier lotion, cream, soap, wash, shampoo or gel.

28. The method of claim 22, wherein the topically-applied acetylcholinesterase inhibitor is applied to said hair follicles, skin, eyes, eyelids, eyelashes, or Meibomian Gland areas affected by the ophthalmological affliction.
15

29. The method of claim 28, wherein the topically-applied acetylcholinesterase inhibitor is further applied to said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian gland areas not affected by the ophthalmological affliction.

20 30. The method of claim 1, wherein the topically-applied acetylcholinesterase inhibitor is applied to said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian gland areas of a region on the face surrounding the eyes where demodex brevis and/or demodex folliculorum mites exist.

25 31. The method of claim 30, wherein the topically-applied acetylcholinesterase inhibitor is applied to substantially all said hair follicles, skin, eyes, eyelids, eyelashes, or Meibomian Gland areas in an eye region of said individual.

32. The method of any of claims 1-31, further comprising a step of applying the
30 acetylcholinesterase inhibitor to an individual's clothing, linens or both clothing and linens.

33. The method of any of claims 1-32, wherein the topically-applied acetylcholinesterase inhibitor is applied to an eye region corresponding to hair

follicles, skin, eyes, eyelids, eyelashes, and/or meibomian gland areas at least once and not more than twice daily for a period of two to six weeks.

34. The method of any of claims 1-33, wherein the individual has affected skin areas, and the topically-applied acetylcholinesterase inhibitor is applied to the affected skin areas and/or to non-affected skin areas during a first application period, thereby killing and eliminating adult demodex brevis and/or demodex folliculorum mites from the hair follicles in the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual.

35. The method of claim 34, wherein the topically-applied acetylcholinesterase inhibitor is further applied to the affected said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian Gland areas and/or to non-affected hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands areas during a second application period, thereby killing and eliminating from the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual demodex brevis and/or demodex folliculorum mites that have matured from a larval form and/or an egg form present on and/or in the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands during the first application period.

36. The method of claim 35, wherein the topically-applied acetylcholinesterase inhibitor is further applied to the affected skin areas and/or to non-affected skin areas during a third application period, thereby killing and eliminating from the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual demodex brevis and/or demodex folliculorum mites that have matured from a larval form and/or an egg form present on and/or in the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands during the first application period and/or the second application period.

37. The method of claim 35 or claim 36, wherein the first application period and the second application period are separated by at least five but no more than ten days.

38. The method of any of claims 35-37, wherein the first application period and the second application period are separated by at least seven days.

39. The method of any of claims 35-38, wherein the first application period and the second application period are separated by a time sufficient to allow the larva form to mature into an adult form and/or to allow the egg form to mature into the adult form.

40. The method of any of claims 36-39, wherein the second application period and the third application period are separated by at least five but no more than ten days.

41. The method of any of claims 36-39, wherein the second application period and the third application period are separated by at least seven days.

42. The method of any of claims 36-41, wherein the second application period and the third application period are separated by a time sufficient to allow the larva form to mature into an adult form and/or to allow the egg form to mature into the adult form.

43. The method of claim 1, wherein the acetylcholinesterase inhibitor is orally-administered or topically-applied in a continued intermittent regime sufficient for prophylactic control of demodex mite population in the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual.

44. The method of claim 43, wherein the acetylcholinesterase inhibitor is administered as an oral dose of the acetylcholinesterase inhibitor of about 150 mg per kg of body mass or less or between about 0.01 mg per kg of body mass and 50 mg per kg of body mass.

45. The method of claim 44, wherein the acetylcholinesterase inhibitor is administered as an oral dose of the acetylcholinesterase inhibitor of a lowest dose effective for killing the demodex mites.

46. The method of claim 45, wherein the orally-administered acetylcholinesterase inhibitor is administered as a daily dose of 10 mg per kg of body mass.
47. The method of claims 45, wherein the orally-administered acetylcholinesterase inhibitor is administered as a daily dose of 7.5 mg per kg of body mass.
48. The method of claim 45, wherein the orally-administered acetylcholinesterase inhibitor is administered as a three times per day dose of 5 mg per kg of body mass.
49. The method of claim 44, wherein the orally-administered acetylcholinesterase inhibitor is repeated about two to four times with spacing of three to seven days between them.
50. The method of claim 1, wherein the orally-administered acetylcholinesterase inhibitor is formulated as a prodrug or pharmaceutically acceptable salt.
51. The method of claim 1, wherein the inactivation of the demodex brevis and/or demodex folliculorum mites from the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual results in a reduction in population of one or more bacteria in the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual.
52. The method of claim 51, wherein the allergic and/or inflammatory responses to the mites result from a presence of one or more bacteria associated with the mites in the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual.
53. The method of claim 51 or claim 52, wherein the one or more bacteria comprise one or more bacteria from the genus staphylococcus or from the genus bacillus.
54. The method of claim 51, wherein the one or more bacteria comprise bacillus oleronius bacteria.

55. The method of any of claims 51-54, wherein the one or more bacteria comprise staphylococcus epidermidis bacteria.

56. The method of any of claims 51-55, wherein the one or more bacteria are
5 present in a digestive system of the demodex brevis and/or demodex folliculorum mites.

57. A method of treating an ophthalmological affliction, the method comprising a
step of topically-applying to an eye region of an individual having the
10 ophthalmological affliction that is a demodex-induced inflammatory eye condition an active ingredient comprising an acetylcholinesterase inhibitor in a dosage sufficient to inactivate demodex brevis and/or demodex folliculorum mites from said eye region, wherein said eye region comprises hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands of the individual, resulting
15 in amelioration or cessation of one or more manifestations of allergic and/or inflammatory responses to the mites that cause symptoms and signs of the ophthalmological affliction in the individual, wherein the topically-applied active ingredient is applied to the said hair follicles, skin, eyes, eyelids, eyelashes, or meibomian glands areas affected by the ophthalmological affliction and to
20 areas not affected by the ophthalmological affliction.

58. The method of claim 57, wherein said active ingredient is a carbamate.

59. The method of claim 58, wherein said carbamate is a miticide or insecticide.
25

60. The method of claim 58, wherein said carbamate is a naturally occurring compound.

61. The method of claim 58, wherein said carbamate is an ethyl carbamate or a
30 prodrug or pharmaceutically acceptable salt thereof.

62. The method of claim 58 wherein said carbamate is selected from the group consisting of: neostigmine and rivastigmine or a derivative, prodrug or pharmaceutically acceptable salt thereof.

63. The method of claim 58, wherein said carbamate is selected from the group consisting of: aldicarb, bendiocarb, bufencarb, carbaryl, carbendazim, carbetamide, carbofuran, carbosulfan, chlorbufam, chlorpropham, ethiofencarb, formetanate, methiocarb, methomyl, oxamyl, phenmedipham, pinmicarb, pirimicarb, propamocarb, propham, propoxur, butocarboxim, carbanolate, promacyl, thiocarboxime, thiofanox, benomyl, and metolcarb or a derivative, prodrug or pharmaceutically acceptable salt thereof.
64. The method of claim 57, wherein said active ingredient is a naturally occurring acetylcholinesterase inhibitor.
65. The method of claim 64, wherein said naturally occurring acetylcholinesterase inhibitor is a miticide or insecticide.
66. The method of claim 64, wherein said naturally occurring acetylcholinesterase inhibitor is witch hazel or a substance derived therefrom.
67. The method of claim 64, wherein said naturally occurring acetylcholinesterase inhibitor is Boswellia sacra resin or a substance derived therefrom.
68. The method of claim 64, wherein said naturally occurring acetylcholinesterase inhibitor is a coumarin or a derivative, prodrug or pharmaceutically acceptable salt thereof.
69. The method of claim 64, wherein said naturally occurring acetylcholinesterase inhibitor is selected from the group consisting of: huperzine A, galantamine, onchidal, coumarins, celastrus paniculatus, and boswellia or a derivative, prodrug or pharmaceutically acceptable salt thereof.
70. The method of claim 57, wherein the active ingredient comprises a miticide or insecticide.

71. The method of any of claims 57-70, wherein the ophthalmological affliction comprises one or more of Meibomian Gland Dysfunction, blepharitis, conjunctivitis, keratoconjunctivitis, hyperemia, or dry eye disease.
- 5 72. A method of treating an ophthalmological affliction that is a demodex-induced inflammatory eye condition, the method comprising a step of orally-administering or topically-applying to an individual having the ophthalmological affliction a carbamate in a dosage sufficient to inactivate demodex brevis and/or demodex folliculorum mites from hair follicles and/or said hair follicles, skin,
10 eyes, eyelids, eyelashes, or meibomian glands of the individual, resulting in amelioration or cessation of the manifestations of allergic and/or inflammatory responses to the mites that cause symptoms and signs of the ophthalmological affliction in the individual.
- 15 73. The method of claim 72, comprising topically applying the carbamate to an eye region of the individual.
74. The method of claim 72-73, wherein said carbamate is a miticide or insecticide.
- 20 75. The method of any of claims 72-74, wherein said carbamate is a naturally occurring compound.
76. The method of claim 72, wherein said carbamate is an ethyl carbamate or a prodrug or pharmaceutically acceptable salt thereof.
- 25 77. The method of claim 72, wherein said carbamate is selected from the group consisting of: neostigmine and rivastigmine or a derivative, prodrug or pharmaceutically acceptable salt thereof.
- 30 78. The method of claim 72, wherein said carbamate is selected from the group consisting of: aldicarb, bendiocarb, bufencarb, carbaryl, carbendazim, carbetamide, carbofuran, carbosulfan, chlorbufam, chlorpropham, ethiofencarb, formetanate, methiocarb, methomyl, oxamyl, phenmedipham, pinmicarb, pirimicarb, propamocarb, propham, propoxur, butocarboxim,

carbanolate, promacyl, thiocarboxime, thiofanox, benomyl, and metolcarb or a derivative, prodrug or pharmaceutically acceptable salt thereof.

79. The method of claim 1, wherein said acetylcholinesterase inhibitor is a
5 reversible competitive or noncompetitive inhibitor of acetylcholinesterase.

80. The method of claim 79, wherein said acetylcholinesterase inhibitor is selected
from the group consisting of: Carbamates, Physostigmin, Neostigmine,
Pyridostigmine, Ambenonium, Demecarium, Rivastigmine, Phenanthrene
10 derivatives, Galantamine, Caffeine, Piperidines, Donepezil, Tacrine or
tetrahydroaminoacridine (THA'), Edrophonium, Huperzine A, Ladostigil,
Ungeremine, and Lactucopicrin, or a derivative, prodrug or pharmaceutically
acceptable salt thereof.

15 81. The method of claim 1, wherein said acetylcholinesterase inhibitor is a quasi-
reversible inhibitor of acetylcholinesterase.

82. The method of claim 81, wherein said acetylcholinesterase inhibitor is selected
from the group consisting of: organophosphates and carbamates, or a
20 derivative, prodrug or pharmaceutically acceptable salt thereof.

83. The method of claim 82, wherein said acetylcholinesterase inhibitor is an
organophosphate selected from the group consisting of: Echothiophate,
Diisopropyl fluorophosphates, Cadusafos, Chlorpyrifos, Dichlorvos, Dimethoate,
25 Metrifonate, Malathion and Parathion, or a derivative, prodrug or
pharmaceutically acceptable salt thereof.

84. The method of claim 82, wherein said acetylcholinesterase inhibitor is a
carbamate selected from the group consisting of: Aldicarb; Bendiocarb;
30 Bufencarb; Carbaryl ; Carbendazim; Carbetamide; Carbofuran; Carbosulfan;
Chlorbufam; Chloroprotham; Ethiofencarb; Formetanate; Methiocarb;
Methomyl; Oxamyl; Phenmedipham; Pinmicarb; Pirimicarb; Propamocarb;
Protham; Propoxur; Huperzine A; Galantamine; Onchidal ; and Coumarins, or a
derivative, prodrug or pharmaceutically acceptable salt thereof.

85. The method of claim 82, wherein said acetylcholinesterase inhibitor is metrifonate or dichlorvos.
- 5 86. The method of claim 1, wherein said acetylcholinesterase inhibitor is a known compound used in medicine and having an established safety profile in humans.
- 10 87. The method of claim 86, wherein said acetylcholinesterase inhibitor is selected from the group consisting of: Aricept; Aricept ODT; Cognex; donepezil; Exelon; galantamine; Namzaric; Razadyne; rivastigmine; tacrine; phospholine neostigmine; parathion; malathion ; dyflos; physostigmine; endrophonium; pyridostigmine; ecothiapate, or a derivative, prodrug or pharmaceutically acceptable salt thereof.
- 15 88. The method of any of the above claims, further comprising the step of applying the acetylcholinesterase inhibitor with one or more of a spray bottle, impregnated swab/qtip, impregnated wipes, eye drops or spoolie brush.
- 20 89. The method of any of the above claims, wherein the topically-applied acetylcholinesterase inhibitor penetrates to a subdermal region of the eyelid of the individual, thereby exposing the demodex brevis and/or demodex folliculorum mites present in and around the meibomian glands to the acetylcholinesterase inhibitor.
- 25 90. A method of treating an individual having a demodex-induced inflammatory eye condition, the method comprising the steps of:
- 30 applying to the individual in need thereof an acetylcholinesterase inhibitor in a dosage sufficient to inactivate demodex brevis mites and/or demodex folliculorum mites from an eye region of the individual, wherein the eye region includes one or more of hair follicles, eyelids, eyelashes and Meibomian glands in and around an eye region;
- wherein a sufficient amount of demodex brevis mites and/or demodex folliculorum mites are inactivated to ameliorate or cease manifestations of

allergic and/or inflammatory responses to the demodex mites that cause the inflammatory eye condition.

- 5 91. The method of claim 91, wherein the inflammatory eye condition corresponds to one or more of: Meibomian gland dysfunction, dry eye, blepharitis, conjunctivitis, keratoconjunctivitis, or hyperemia,.
92. The method of any of claims 90-91, wherein the application is a topical administration to the eye region of the individual.
- 10 93. The method of claim 92, wherein the eye region corresponds to an area of between at least 1 cm and 10 cm around an outermost perimeter defined by the eye, eyelid, eyebrow and eyelashes.

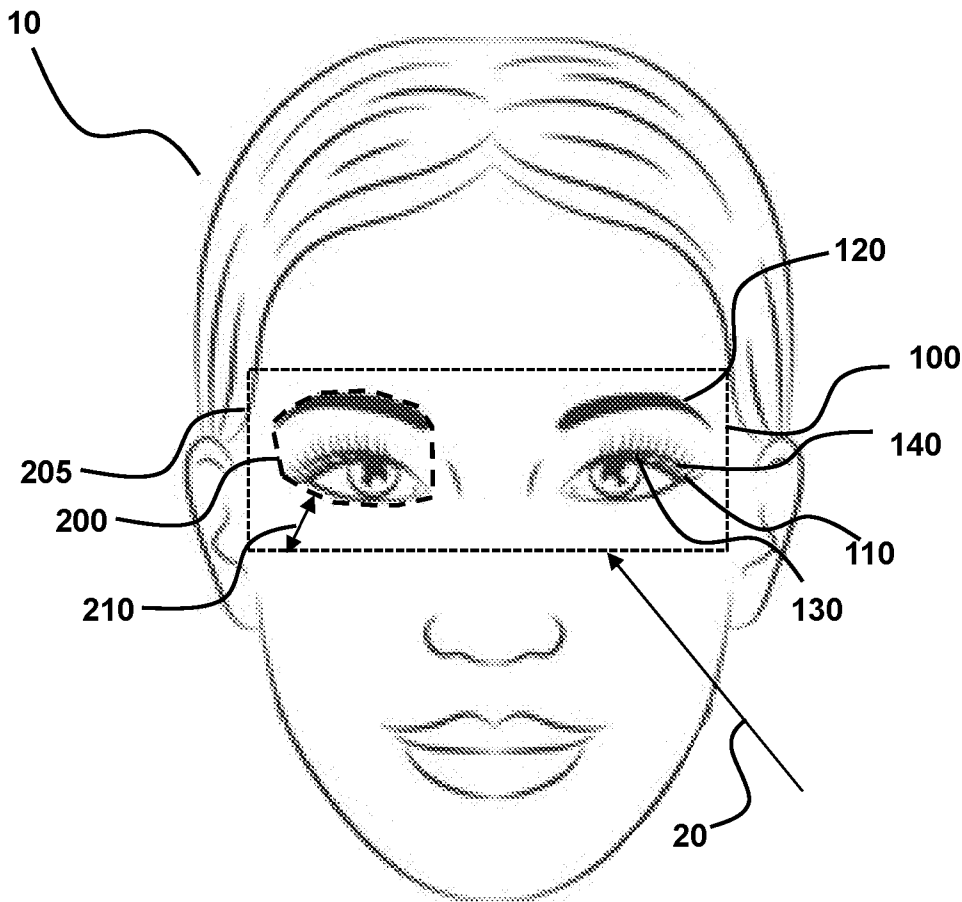


FIG. 1

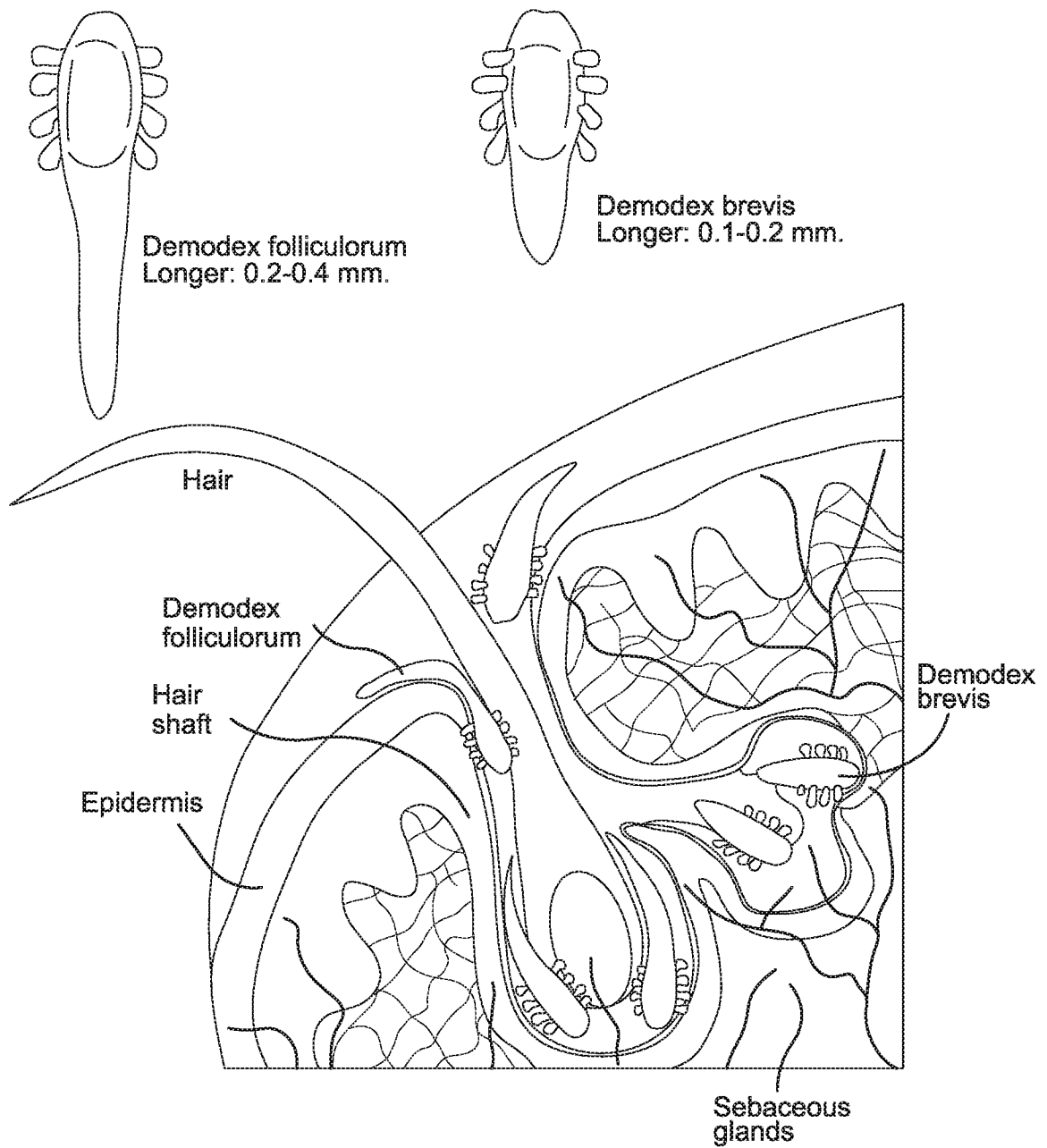


FIG. 2

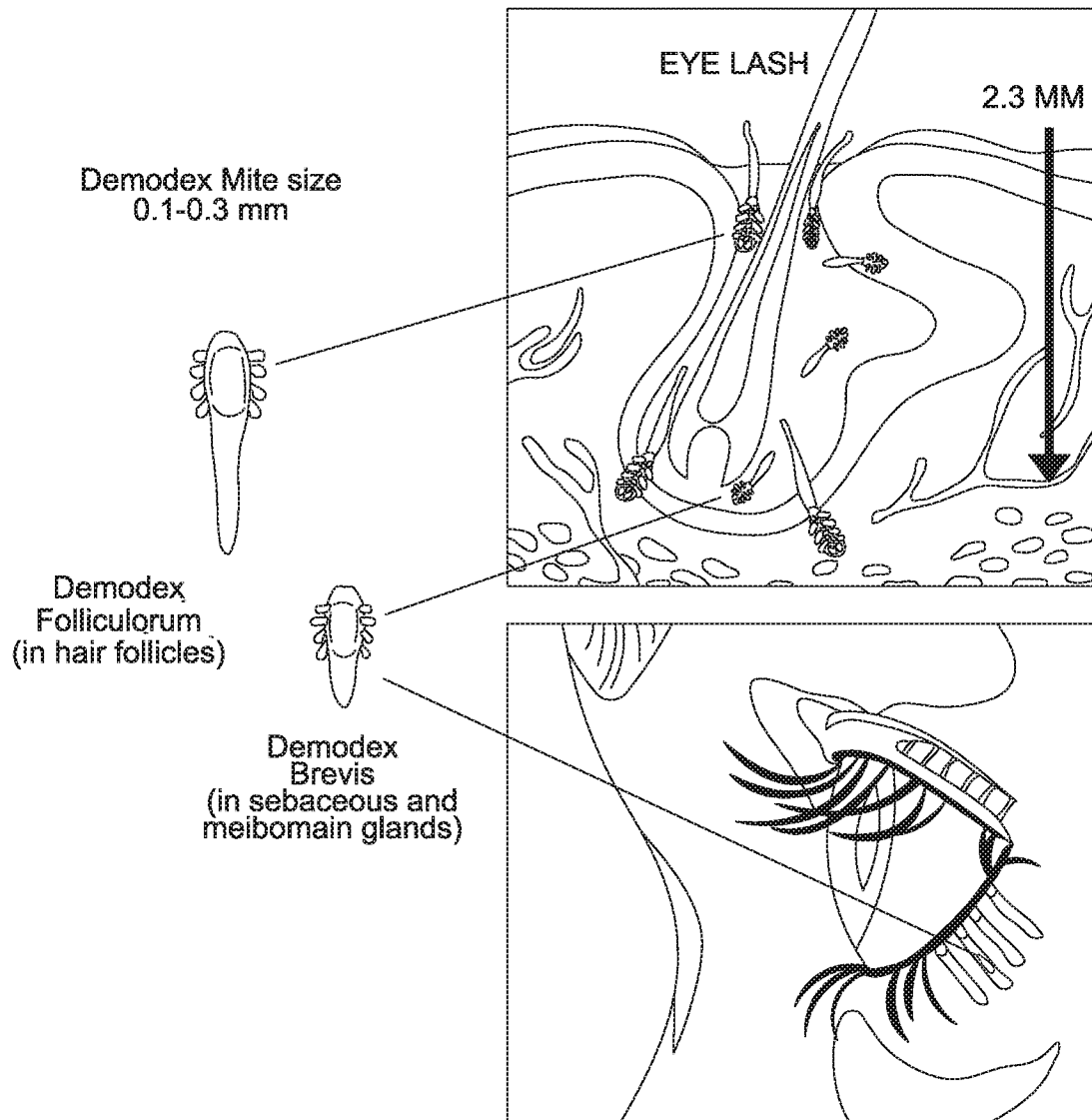


FIG. 3

Compounds	LT50 in Minutes	Average minutes to death	Total mites	LT50
FCO	181	198.02	51	26/51
Tonka Bean (couramin)	181	198.02	51	26/51
Frankincense (boswellia sacra)	24	26.71	51	26/51
Mineral Oil	50	50.78	51	26/51
Wild Lettuce (lactucopicrin)	494	497.51	121	61/121
Malkangni Seed Oil (celastrus paniculatus)	119	64.29	57	29/57
Dichlorvos 1% water	5	9.16	50	25/50
Malathion 1% water	113	110.34	56	28/56
Malathion 1% oil	388	379.06	50	25/50
Metrifonate 1% water	437	432.38	63	32/63
Metrifonate 1% oil	354		50	25/50
Physostigmine 1% water	248	257.90	51	26/51
Galantamine 1% water	181	216.02	60	30/60
Demacarium 1% water	703		65	33/65
Donepezil 1% water	539		58	29/58
Tacrine 1% water	603		50	25/50
Diisopropyl fluorophosphate	426		58	29/58
Echothiophate 1% water	561		63	32/63
Rivastigmine 1% water				
Neostigmine 1% water	No activity		19	
Warfarin 1% water	No activity		51	
Pyridostigmine 1% water	No activity		54	
Edrophonium 1% water	No activity		52	
Ambenonium 1% water	No activity		57	
Distilled Water	No activity		59	
Chloroquine 10% water	2688			56/112
Doxycycline 5% water	1896		56	28/56
Huperzine A (natural) 1% water	685		62	31/62
Huperzine A (synthetic) 1% water	450		57	29/57

FIG. 4

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US18/66867

A. CLASSIFICATION OF SUBJECT MATTER

IPC - A61K 31/27, 31/55, 31/473, 36/185, 36/324 (2019.01)

CPC - A61K 0/0014, 9/0053, 31/27, 31/55, 31/473, 31/662, 31/683, 36/185, 36/324

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

See Search History document

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

See Search History document

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

See Search History document

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2017/0135978 A1 (ATTILLAPS HOLDINGS) 18 May 2017; claims 2-5 and 88; paragraphs [0023], [0026]-[0028], [0030], [0036]-[0039], [0041]-[0043], [0048]-[0049]; [0056], [0064]-[0066], [0076], [0094]-[0097], [0099]	1-8, 9/1-8, 10/1-8, 11-21, 30-31, 43-52, 53/51-52, 54, 57-70, 71/57-70, 72-73, 74/72-73, 76-87, 90-91, 92/90-91
----- Y		----- 93/92/90-91
Y	US 8,128,968 B2 (GAO YY, et al.) 6 March 2012; column 5, lines 4-15 and 24-30; column 9, lines 2-13	93/92/90-91
A	GAO Y-Y, et al. "In Vitro and In vivo Killing of Ocular Demodex by Tea Tree Oil" British Journal of Ophthalmology 2005, vol. 58, pages 1468-1473	1
A	KOO H, et al. "Ocular Surface Discomfort and Demodex: Effect of Tea Tree Oil Eyelid Scrub in Demodex Blepharitis" J. Korean Med Sci 2012, vol. 27, pages 1574-1579	1

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

09 February 2019 (09.02.2019)

Date of mailing of the international search report

08 MAR 2019

Name and mailing address of the ISA/

Mail Stop PCT, Attn: ISA/US, Commissioner for Patents
P.O. Box 1450, Alexandria, Virginia 22313-1450
Facsimile No. 571-273-8300

Authorized officer

Shane Thomas

PCT Helpdesk: 571-272-4300
PCT OSP: 571-272-7774

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US18/66867

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☒ Claims Nos.: 22-29, 32-42, 55-56, 75, and 88-89
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.