

NON-SURGICAL METHOD OF TREATMENT FOR CATARACT

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority to U.S. Provisional Patent Application No. 61/672,569, filed July 17, 2012, which is hereby incorporated by reference in its entirety.

STATEMENT OF GOVERNMENT INTEREST

[0002] This invention was made with government support under RR024986 awarded by the National Institutes of Health. The government has certain rights in the invention.

FIELD OF THE INVENTION

[0003] The disclosure generally relates to inhibitors of α -crystallin aggregation, uses thereof, and methods of screening for therapeutically effective modulators of protein aggregation.

BACKGROUND OF THE INVENTION

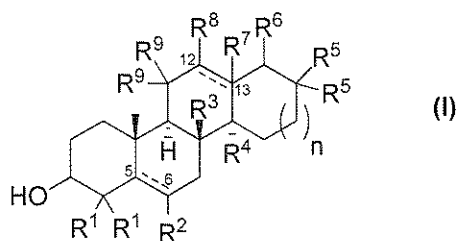
[0004] Cataract, or a clouding of the eye lens, is a condition affecting over half of all adults over the age of 80, with approximately 25 million patients suffering from the condition in the United States. Moreover, cataracts are thought to be the leading cause of blindness worldwide. α A-crystallin (cryAA) and α B-crystallin (cryAB) comprise thirty percent of the protein content of the eye lens, where they are responsible for maintaining lens transparency (Haslbeck et al., *Nat Struct Mol Biol* 12, 842 (2005)). cryAA and cryAB belong to a family of small heat shock proteins (sHSPs) that contain a conserved crystallin domain (Bloemendal et al., *Prog Biophys Mol Biol* 86, 407 (2004); Haslbeck, *supra*). Once synthesized, these lens sHSPs are never degraded, so any damage accumulates through life and eventually leads to aging-associated cataract (Haslbeck, *supra*; Perng et al., *J Biol Chem* 274, 33235 (1999); Meehan et al., *J Biol Chem* 279, 3413 (2004); Meehan et al., *J Mol Biol* 372, 470 (2007)). Similarly, destabilizing mutations in cryAB, such as R120G, result in hereditary forms of cataract with early onset (Vicart et al., *Nat Genet* 20, 92 (1998)). In hereditary cataract, cryAB is prone to aggregation and forms amyloid-like fibrils *in vitro* (Andley et al., *PLoS One* 6, e17671 (2011)).

[0005] Currently, treatment for cataracts includes surgery to excise the clouded lens and insert an artificial replacement. Surgery can be costly and is not appropriate for all patients.

Therapies which target the underlying mechanism of protein aggregation would benefit these patients. Thus, there remains a need in the art for therapeutics and methods of screening for therapeutics that block aggregation-prone proteins, such as cryAB, from forming pathological aggregates (e.g., aggregates associated with cataracts).

SUMMARY

[0005a] In a first aspect of the present disclosure, there is provided a use of a composition comprising a therapeutically effective amount of a compound of formula I for the manufacture of a medicament for treating or preventing cataract in an individual in need thereof:



wherein:

each R^1 is H or each R^1 is Me;

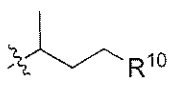
R^2 is H or OH;

the dashed line between carbons 5 and 6 indicates an optional double bond;

R^3 is H or Me;

R^4 is H or Me;

n is 0 or 1;

R^6 is  and each R^5 is independently H or Me;

the dashed line between carbons 12 and 13 is an optional double bond,

R^7 is H or Me, with the proviso that R^7 is not present when the double bond between carbons 12 and 13 is present;

R^8 is H or OH;

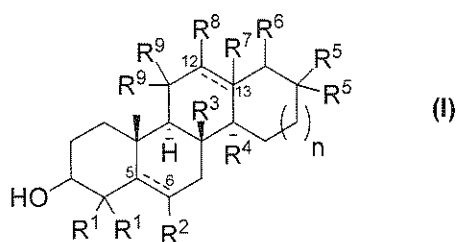
both R^9 groups together form an oxo (=O) or each R^9 is hydrogen; and

R^{10} is branched C_1 - C_6 alkyl, substituted with a hydroxyl group;

or pharmaceutically acceptable salt thereof.

[0005b] In a second aspect of the present disclosure, there is provided a use of 5a-cholestan-3b-ol-6-one for the manufacture of a medicament for treating or preventing cataract.

[0006] The present disclosure provides a method of treating or preventing cataract, the method comprising administering to an individual in need thereof an effective amount of a composition comprising a compound of formula I:



wherein:

both R^1 are H or both R^1 are Me;

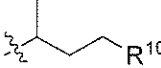
R^2 is H or OH;

dashed line between carbons 5 and 6 indicates an optional double bond;

R^3 is H or Me;

R^4 is H or Me;

n is 0 or 1;

(a) R^6 is  and each R^5 is independently H or Me or (b) R^6 and one R^5 taken together form an optionally substituted 6-membered ring and the other R^5 is Me;

the dashed line between carbons 12 and 13 is an optional double bond, with the proviso that R^7 is not present when the double bond between carbons 12 and 13 is present, and R^7 is H or Me when the double bond between carbons 12 and 13 is not present;

R^8 is H or OH;

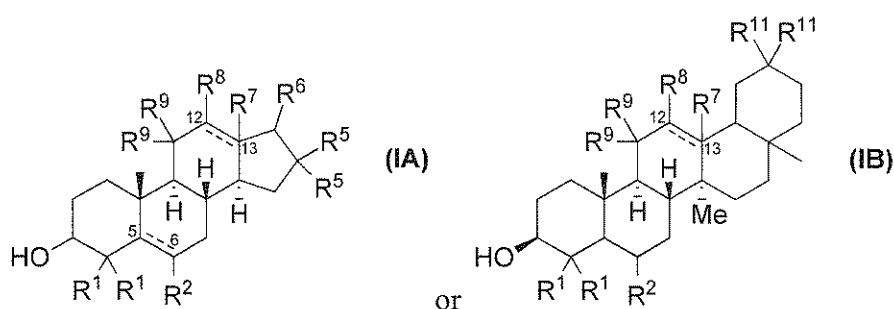
both R^9 together form an oxo (=O) or both R^9 are hydrogen; and

R^{10} is CO_2H or linear or branched $\text{C}_1\text{-C}_6$ alkyl;

or a prodrug or pharmaceutically acceptable salt thereof.

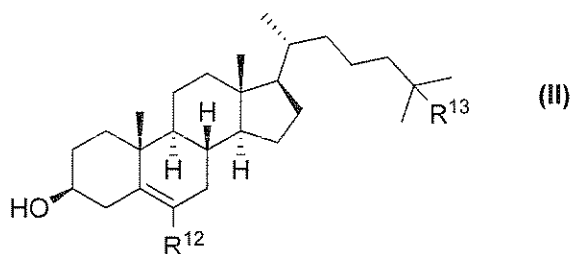
[0007] The present disclosure also provides an ophthalmic pharmaceutical composition comprising a pharmaceutically acceptable ophthalmic carrier and a compound of formula I.

[0008] In various embodiments of the method and/or composition, the compound of formula I has a structure of formula IA or formula IB:



wherein each R^{11} is independently alkyl, CO_2H , or CO_2alkyl .

[0009] In a more specific embodiment of the method and/or composition, the compound has a structure of formula II:



wherein R^{12} is H or OH and R^{13} is H or OH. In one embodiment of the method and/or composition, the compound is 5-cholesten-3 β ,25-diol.

[0010] In various embodiments of the method, the composition is administered topically, subconjunctivally, retrobulbarly, periocularly, subretinally, suprachoroidally, or intraocularly. In various embodiments of the method, the cataract is an age-related cataract or a diabetic cataract. In some embodiments of the method, the individual has a hereditary form of cataract with early onset. In more specific embodiments of the method, the individual has a R120G mutation and/or a D109H mutation in cryAB.

[0011] In some embodiments of the composition, the pharmaceutically acceptable ophthalmic carrier is a cyclodextrin. In a specific embodiment, the cyclodextrin is (2-hydroxypropyl)- β -cyclodextrin.

[0012] Additionally, the present disclosure includes a high-throughput method of screening compounds for modulation of protein thermal stability, the method comprising: (a) contacting a protein with each of a plurality of test compounds; and (b) measuring the melting transition (T_m) of the protein in the presence of each of the plurality of test compounds, wherein a compound that decreases or increases the apparent T_m by at least 2 standard deviations is a pharmacological protein chaperone.

[0013] In various embodiments of the method, the protein is an amyloid-forming protein or a protein underlying a loss-of-function disease. In some embodiments, the amyloid-forming protein is selected from the group consisting of Hsp27, α A-crystallin, α B-crystallin, β B2-crystallin, β B1-crystallin, γ D-crystallin, Hsp22, Hsp20, tau, Alpha-synuclein, IAPP, beta-amyloid, PrP, Huntingtin, Calcitonin, Atrial natriuretic factor, Apolipoprotein AI, Serum amyloid A, Medin, Prolactin, Transthyretin, Lysozyme, Beta 2 microglobulin, Gelsolin, Keratoepithelin, Cystatin, Immunoglobulin light chain AL, and S-IBM. In other embodiments, the protein underlying a loss-of-function disease is selected from the group consisting of mutant β -glucosidase, cystic fibrosis transmembrane receptor, hexosaminidase A, hexosaminidase B, β -galactosidase, and alpha-glucosidase.

[0014] In some embodiments of the method, the T_m is determined using a high-throughput differential scanning fluorimetry device.

[0015] In one embodiment of the method, the measuring step comprises: (b1) heating the protein in the presence of each of a plurality of test compounds from 50°C to 80°C, (b2) cooling the protein to 25°C, (b3) maintaining the protein at 25°C for 10 seconds, and (b4) measuring the fluorescence of the protein.

[0016] In various embodiments, the method further comprises repeating steps (b1)-(b4) between 2 and 30 times, wherein each repeat of step (b1) is performed at an incrementally higher temperature. In specific embodiments, the amyloid-forming protein is heated from 65°C to 80°C in 1°C increments. In other embodiments, (b1) further comprises, after heating, equilibrating the

amyloid-forming protein and test compound between 60 and 180 seconds. In various embodiments of the method, the equilibrating step is 130 seconds.

[0017] Additionally, the present disclosure includes a high-throughput screening system, comprising: (a) an amyloid-forming protein; (b) a device capable of measuring the melting transition (T_m) of the amyloid-forming protein; and (c) a plurality of test compounds.

[0018] In various embodiments of the screening system, the protein is selected from the group consisting of Hsp27, α A-crystallin, α B-crystallin, β B2-crystallin, β B1-crystallin, γ D-crystallin, Hsp22, Hsp20, tau, Alpha-synuclein, IAPP, beta-amyloid, PrP, Huntingtin, Calcitonin, Atrial natriuretic factor, Apolipoprotein AI, Serum amyloid A, Medin, Prolactin, Transthyretin, Lysozyme, Beta 2 microglobulin, Gelsolin, Keratoepithelin, Cystatin, Immunoglobulin light chain AL, and S-IBM.

[0019] In some embodiments of the screening system, the device is a high-throughput differential scanning fluorimetry device.

[0020] The use of the compound of any one of structural formulas I, IA, IB, or II in any of the methods disclosed herein or for preparation of medicaments for administration according to any of the methods disclosed herein is specifically contemplated. In this regard, the present disclosure provides a compound of any one of structural formulas I, IA, IB, or II for use in a method of treating or preventing cataract, wherein the method comprises administering to an individual in need thereof an effective amount the compound.

[0021] Other features and advantages of the present disclosure will become apparent from the following detailed description. It should be understood, however, that the detailed description and the specific examples, while indicating specific embodiments of the disclosure, are given by way of illustration only, because various changes and modifications within the spirit and scope of the disclosure will become apparent to those skilled in the art from this detailed description. The entire document is intended to be related as a unified disclosure, and it should be understood that all combinations of features described herein are contemplated, even if the combination of features are not found together in the same sentence, or paragraph, or section of this document. In addition to the foregoing, the present disclosure includes, as an additional aspect, all embodiments of the present disclosure narrower in scope in any way than the variations specifically mentioned above. For example, if aspects

of the present disclosure are described as “comprising” a feature, embodiments also are contemplated “consisting of” or “consisting essentially of” the feature.

DETAILED DESCRIPTION OF THE INVENTION

[0022] The invention provides inhibitors of α -crystallin aggregation and methods of using α -crystallin aggregation inhibitors to, e.g., treat or prevent cataracts in a subject having or at risk of developing cataracts. Inhibitors of α -crystallin aggregation of the invention are, e.g., sterols represented by Formula I, Formula IA, Formula IB, and Formula II and may be formulated in ophthalmic pharmaceutical compositions comprising a pharmaceutically acceptable ophthalmic carrier. Because cataracts affect such a large portion of the population (over half of all adults over the age of 80) and the predominant treatment is surgical intervention, the discoveries described herein represent a significant advancement in the non-surgical methodologies available for treating cataracts. Moreover, many non-surgical treatments currently in use tend to inhibit further aggregation of α -crystallin. Remarkably, compounds of the invention are able to reverse aggregation of α -crystallin and inhibit further aggregation of α -crystallin.

[0023] The invention further provides high-throughput methods of screening compounds for modulation of protein thermal stability, the method comprising contacting a protein with each of a plurality of test compounds; and measuring the melting transition (T_m) of the protein in the presence of each of the plurality of test compounds, wherein a compound that decreases or increases the apparent T_m by at least 2 standard deviations is identified as a pharmacological protein chaperone.

Methods of treating or preventing cataract

[0024] In some embodiments, the invention provides a method of treating or preventing cataract, the method comprising administering to an individual in need thereof an effective amount of a composition comprising a compound of any one of structural formulae I, IA, IB, or II, or e.g., a compound in Table 1. In some embodiments, the cataract is an age-related cataract, a diabetic cataract, a cataract associated with surgery, a cataract resulting from exposure to radiation, a cataract resulting from a genetic illness, a cataract resulting from an infection, or a cataract resulting from medication. In some embodiments, the individual has a hereditary form of cataract with early onset. For example, hereditary forms of cataract include individuals with a R120G mutation and/or a D109H mutation in cryAB.

[0025] An individual “in need of” treatment according to the invention is an individual that is suffering from a cataract. For example, the individual may have an age-related cataract or a cataract resulting from having diabetes. Similarly, an individual “in need of” treatment according to the invention is an individual that is at risk for developing a cataract. Individuals at risk of developing a cataract include, but are not limited to, individuals with a family history of developing cataracts, individuals with a mutation linked to a cataract with early onset, such as individuals with a R120G mutation and/or a D109H mutation in cryAB, individuals exposed to radiation, diabetics, and the like. For example, in one aspect, the individual has been diagnosed with cataract in one eye, and the compound is administered to prevent or slow cataract formation in the contralateral eye.

[0026] “Treating” cataract does not require a 100% abolition of a cataract. Similarly, “prevention” does not require 100% inhibition of cataract formation. Any decrease in cloudiness or deceleration of cataract progression constitutes a beneficial biological effect in a subject. In this regard, the invention reduces the cataract, e.g., at least about 5%, at least about 10% or at least about 20% compared to levels observed in the absence of the inventive method (e.g., in a biologically-matched control subject or specimen that is not exposed to the compound of the inventive method). In some embodiments, the cataract is reduced by at least about 30%, at least about 40%, at least about 50%, or at least about 60%, at least about 70%, at least about 80%, at least about 90%, or more (about 100%).

[0027] In some embodiments, the “treating” cataracts according to inventive method inhibits cataract formation by, e.g., at least about 5%, at least about 10% or at least about 20% compared to levels observed in the absence of the inventive method (e.g., in a biologically-matched control subject or specimen that is not exposed to the compound of the inventive method). In some embodiments, cataract formation is inhibited by at least about 30%, at least about 40%, at least about 50%, or at least about 60%, at least about 70%, at least about 80%, at least about 90%, or more (about 100%) compared to cataract formation in the absence of the compound of the inventive method. Cataracts generally are detected using any of a number of optic tests including, but not limited to, visual acuity testing, ophthalmoscopy, slit-lamp examination, keratometry, tonometry, contrast testing, glare sensitivity, wavefront mapping.

[0028] An “effective amount” of a composition comprising a compound of any one of structural formulae I, IA, IB, or II, or a compound in Table 1 is an amount that inhibits or

reduces aggregation of an amyloid-forming protein such as cryAB in an individual. Inhibiting aggregation does not require a 100% inhibition of aggregation. Any inhibition of aggregation constitutes a beneficial biological effect in a subject. In this regard, the invention inhibits the aggregation of an amyloid-forming protein, e.g., at least about 5%, at least about 10% or at least about 20% compared to levels observed in the absence of the inventive method (e.g., in a biologically-matched control subject or specimen that is not exposed to the compound of the inventive method). In some embodiments, the formation of amyloid aggregates is inhibited by at least about 30%, at least about 40%, at least about 50%, or at least about 60%. In some embodiments, the inventive method inhibits amyloid formation by at least about 70%, at least about 80%, at least about 90%, or more (about 100%) compared to amyloid formation in the absence of the compound of the inventive method.

[0029] Similarly, an “effective amount” for reducing aggregation does not require a 100% abolition of aggregation. Any reduction of aggregation constitutes a beneficial biological effect in a subject. In this regard, the invention reduces the aggregation of an amyloid-forming protein, e.g., at least about 5%, at least about 10% or at least about 20% compared to levels observed in the absence of the inventive method (e.g., in a biologically-matched control subject or specimen that is not exposed to the compound of the inventive method). In some embodiments, amyloid aggregates are reduced by at least about 30%, at least about 40%, at least about 50%, or at least about 60%. In some embodiments, the inventive method reduces amyloid aggregates by at least about 70%, at least about 80%, at least about 90%, or more (about 100%) compared to amyloid aggregates in the absence of the compound of the inventive method.

Route of administration

[0030] As will be understood by those skilled in the art, the most appropriate method of administering a compound to a subject is dependent on a number of factors. In various embodiments, the compound according to the invention is administered locally to the eye, e.g., topically, subconjunctivally, retrobulbarly, periocularly, subretinally, suprachoroidally, or intraocularly. For example, in various embodiments, the composition is delivered locally to the eye via injection. Injectable solutions can be directly injected into the cornea, crystalline lens and vitreous or their adjacent tissues using a fine needle. The composition also can be administered as an intraocular perfusate.

[0031] Additional contemplated routes of administration include, but are not limited to, one or more of: oral (e.g., as a tablet, capsule, or as an ingestible solution), mucosal (e.g., as a nasal spray or aerosol for inhalation), nasal, parenteral (e.g., by an injectable form), gastrointestinal, intraspinal, intraperitoneal, intramuscular, intravenous, intrauterine, intradermal, intracranial, intratracheal, intravaginal, intracerebroventricular, intracerebral, subcutaneous, transdermal, rectal, buccal, epidural and sublingual.

[0032] In some embodiments, the mode for delivery of a composition of the invention to the eye is via a contact lens. The lens may be provided pre-treated with the desired compound. Alternatively, the lens is provided in a kit with components for preparing a coated lens, which are provided as lyophilized powders for reconstitution or as concentrated or ready-to-use solutions. The compositions can be provided as kits for single or multi-use.

[0033] In some embodiments, the mode for delivery of a composition of the invention to the eye is via an ophthalmic rod (Gwon et al., *Ophthalmology*. 1986 Sep;93(9 Suppl):82-5). In some embodiments, the mode for delivery of a composition of the invention to the eye is via an intraocular lens-hydrogel assembly (Garty et al., *Invest Ophthalmol Vis Sci*, 2011 Aug 3;52(9):6109-16).

Dose

[0034] The composition comprising the compound is provided in a therapeutically effective amount that achieves a desired biological effect at a medically-acceptable level of toxicity. The dosage of the compositions may vary depending on the route of administration and the severity of the disease. The dosage may also be adjusted depending on the body weight, age, sex, and/or degree of symptoms of each patient to be treated. The precise dose and route of administration will ultimately be at the discretion of the attendant physician or veterinarian. It will be appreciated that it may be necessary to make routine variations to the dosage depending on the age and weight of the patient as well as the severity of the condition to be treated. The frequency of administration depends on the formulation and the aforementioned parameters. For example, it may be desirable to apply eye drops at least once per day, including 2, 3, 4, or 5 times per day.

[0035] Exemplary doses of the compounds for administration to a human (of approximately 70 kg body weight) via an ocular route are 0.1 mg to 1 g, e.g., 1 mg to 500 mg of the compound per unit dose. In various embodiments, the dose is about 1 µg/kg body

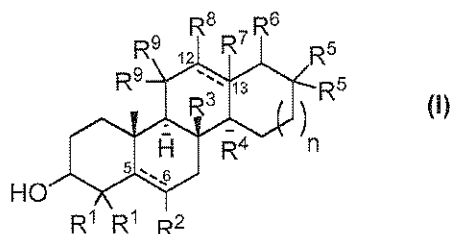
weight to 15 mg/kg body weight. For example, the dose can be 1 $\mu\text{g/kg}$, 5 $\mu\text{g/kg}$, 10 $\mu\text{g/kg}$, 20 $\mu\text{g/kg}$, 30 $\mu\text{g/kg}$, 40 $\mu\text{g/kg}$, 50 $\mu\text{g/kg}$, 60 $\mu\text{g/kg}$, 70 $\mu\text{g/kg}$, 80 $\mu\text{g/kg}$, 90 $\mu\text{g/kg}$, 100 $\mu\text{g/kg}$, 120 $\mu\text{g/kg}$, 140 $\mu\text{g/kg}$, 160 $\mu\text{g/kg}$, 180 $\mu\text{g/kg}$, 200 $\mu\text{g/kg}$, 300 $\mu\text{g/kg}$, 400 $\mu\text{g/kg}$, 500 $\mu\text{g/kg}$, 600 $\mu\text{g/kg}$, 700 $\mu\text{g/kg}$, 800 $\mu\text{g/kg}$, 900 $\mu\text{g/kg}$, 1 mg/kg, 2 mg/kg, 3 mg/kg, 4 mg/kg, 5 mg/kg, 6 mg/kg, 7 mg/kg, 8 mg/kg, 9 mg/kg, 10 mg/kg, 11 mg/kg, 12 mg/kg, 13 mg/kg, 14 mg/kg, or 15 mg/kg. Exemplary doses of the compounds for administration to a human (of approximately 70 kg body weight) via systemic route are 0.1 mg to 5 g, e.g., 1 mg to 2.5g of the compound per unit dose.

[0036] Preferred concentrations of the compound of formula I, IA, IB, or II, or a compound listed in Table 1, range from about 1 $\mu\text{g/ml}$ to 500 $\mu\text{g/ml}$, for example, about 1 $\mu\text{g/ml}$, about 2 $\mu\text{g/ml}$, about 3 $\mu\text{g/ml}$, about 4 $\mu\text{g/ml}$, about 5 $\mu\text{g/ml}$, about 10 $\mu\text{g/ml}$, about 20 $\mu\text{g/ml}$, about 30 $\mu\text{g/ml}$, about 40 $\mu\text{g/ml}$, about 50 $\mu\text{g/ml}$, about 60 $\mu\text{g/ml}$, about 70 $\mu\text{g/ml}$, about 80 $\mu\text{g/ml}$, about 90 $\mu\text{g/ml}$, about 100 $\mu\text{g/ml}$, about 120 $\mu\text{g/ml}$, about 140 $\mu\text{g/ml}$, about 160 $\mu\text{g/ml}$, about 180 $\mu\text{g/ml}$, about 200 $\mu\text{g/ml}$, about 250 $\mu\text{g/ml}$, about 300 $\mu\text{g/ml}$, about 350 $\mu\text{g/ml}$, about 400 $\mu\text{g/ml}$, about 450 $\mu\text{g/ml}$, or about 500 $\mu\text{g/ml}$. The inhibitor may be provided in combination with other pharmaceutically active agents.

[0037] The composition according to the invention is provided in a container, either as a concentrate that is diluted prior to use in an appropriate diluent or at the ready-to-use concentration. Preferably, single dosages are provided in sterile vials.

Compounds effective in treating or preventing cataract

[0038] In various embodiments, the compound of the inventive method or composition is a compound of formula I:



wherein:

both R^1 are H or both R^1 are Me;

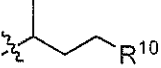
R^2 is H or OH;

dashed line between carbons 5 and 6 indicates an optional double bond;

R^3 is H or Me;

R^4 is H or Me;

n is 0 or 1;

(a) R^6 is  and each R^5 is independently H or Me or (b) R^6 and one R^5 taken together form an optionally substituted 6-membered ring and the other R^5 is Me;

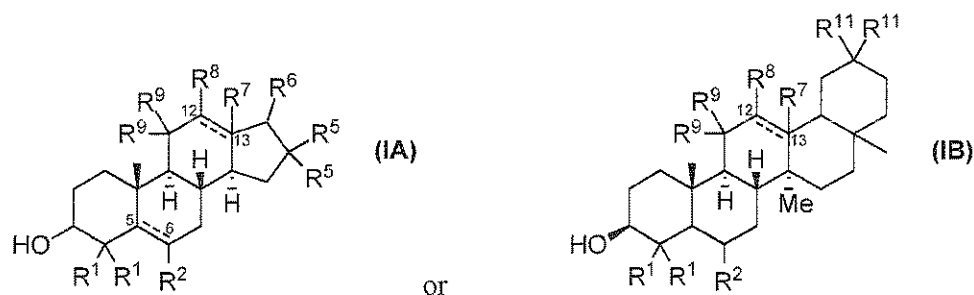
the dashed line between carbons 12 and 13 is an optional double bond, with the proviso that R^7 is not present when the double bond between carbons 12 and 13 is present, and R^7 is H or Me when the double bond between carbons 12 and 13 is not present;

R^8 is H or OH;

both R^9 together form an oxo (=O) or both R^9 are hydrogen; and

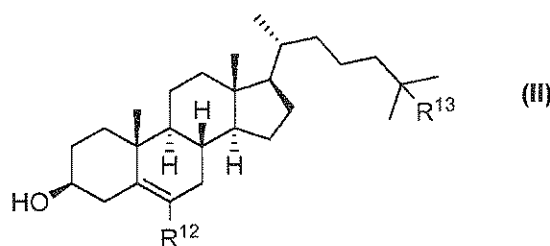
R^{10} is CO_2H or linear or branched C_1 - C_6 alkyl.

[0039] For example, the compound of the inventive method or composition is a compound of formula IA or formula IB:



wherein each R^{11} is independently alkyl, CO_2H , or CO_2 alkyl.

[0040] In various embodiments, the compound has a structure of formula II:

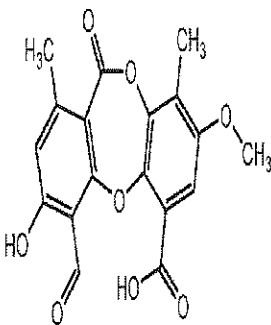
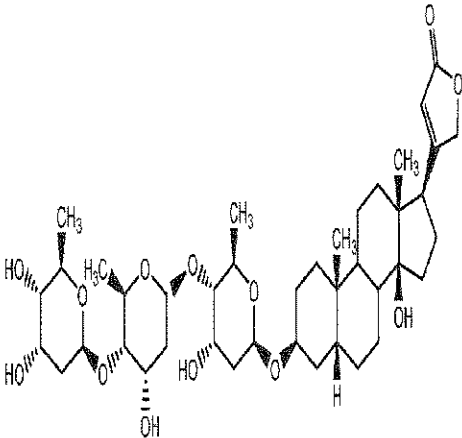
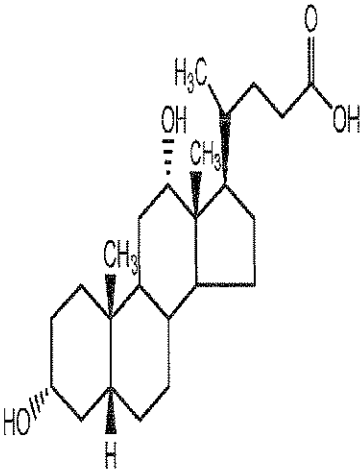


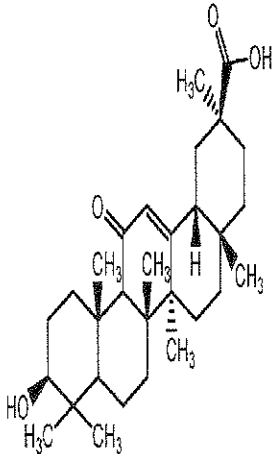
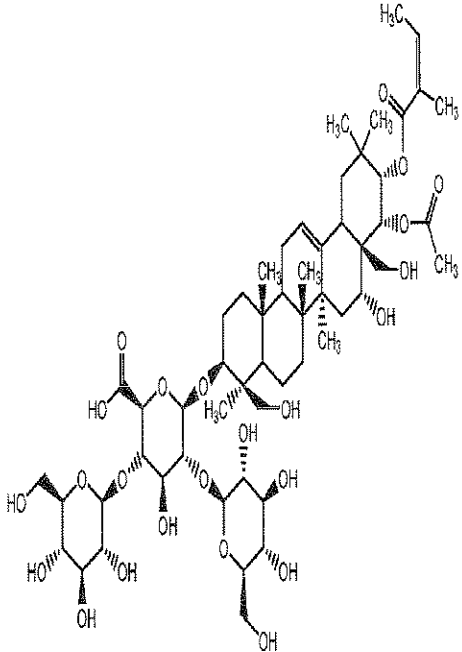
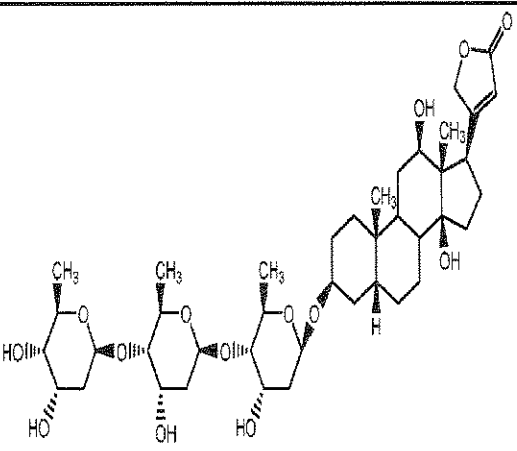
wherein R^{12} is H or OH and R^{13} is H or OH. In some embodiments, the compound is 5-cholesten-3 β ,25-diol.

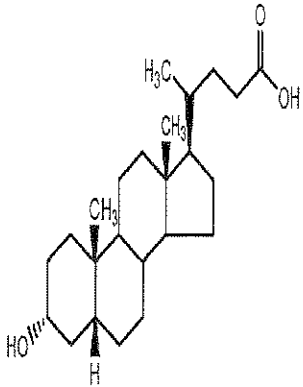
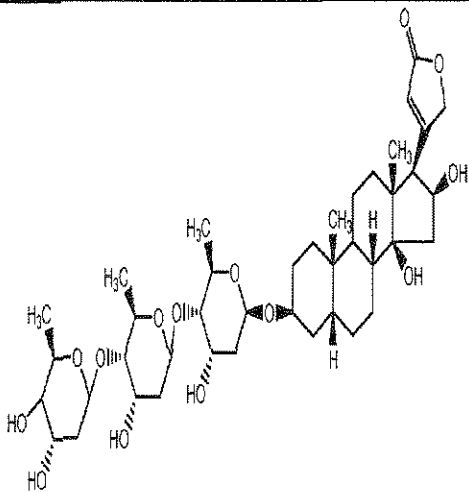
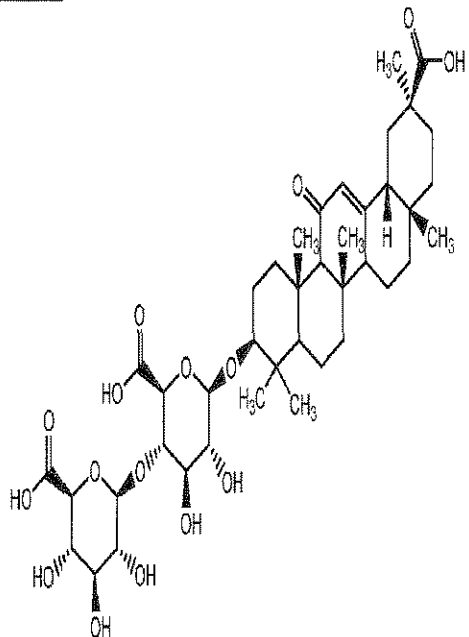
[0041] Alternatively, the compound is a compound listed in Table 1.

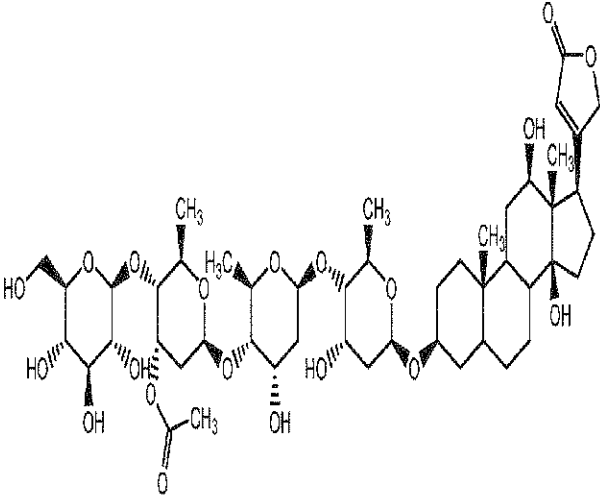
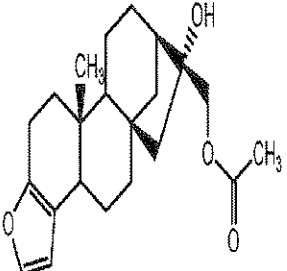
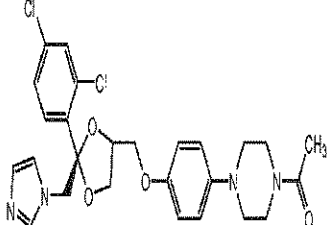
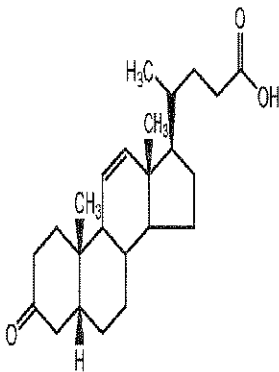
[0042] Any prodrug or pharmaceutically acceptable salt of the above compounds are contemplated to be within the scope of the invention.

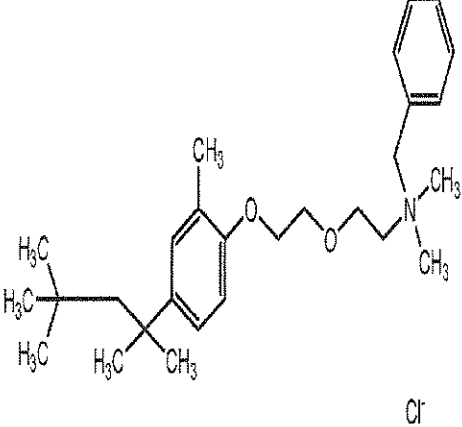
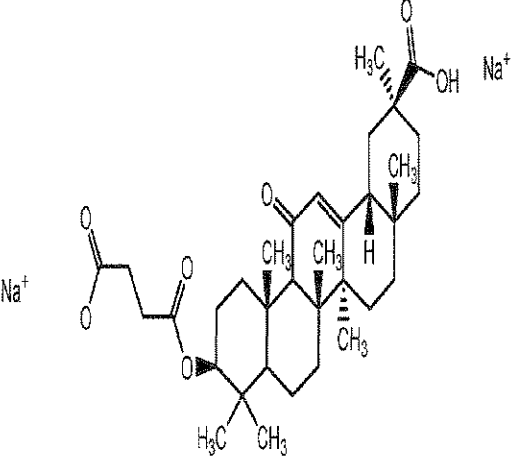
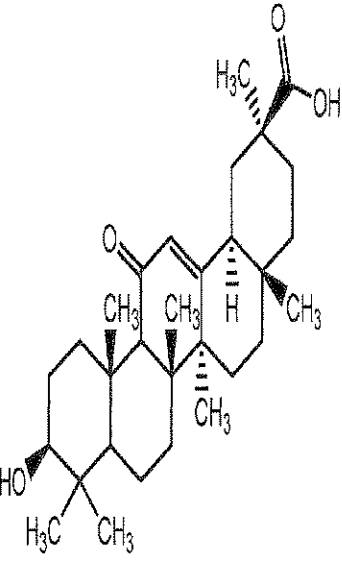
Table 1

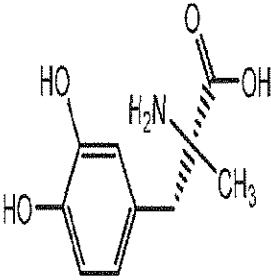
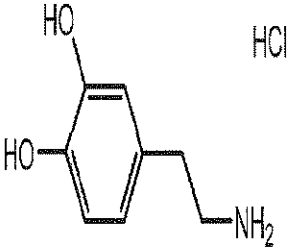
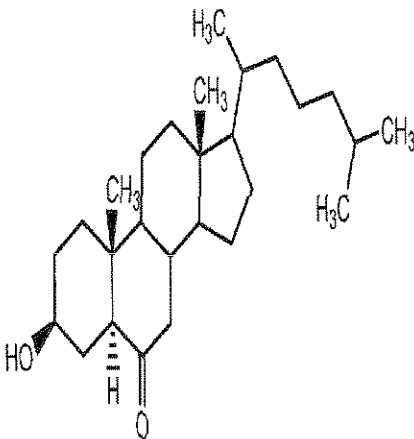
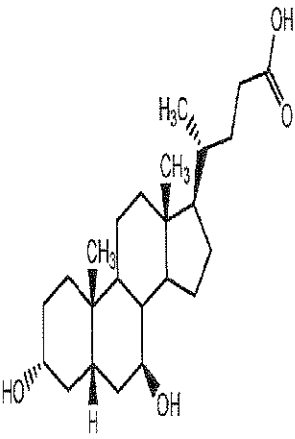
STRUCTURE	CCG ID	NAME	IUPAC
	38361	PSOROMIC ACID	15-formyl-14-hydroxy-6-methoxy-7,12-dimethyl-10-oxo-2,9-dioxatricyclo[9.4.0.0 ^{3,8}]pentadeca-1(11),3(8),4,6,12,14-hexaene-4-carboxylic acid
	38362	DIGITOXIN	4-[(2S,5S,7R,11S,14R,15R)-5-[[[(2R,4S,5S,6R)-5-[(2S,4S,5S,6R)-5-[[[(2S,4S,5S,6R)-4,5-dihydroxy-6-methyloxan-2-yl]oxy]-4-hydroxy-6-methyloxan-2-yl]oxy]-4-hydroxy-6-methyloxan-2-yl]oxy]-11-hydroxy-2,15-dimethyltetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]heptadecan-14-yl]-2,5-dihydrofuran-2-one
	38631	DEOXYCHOLIC ACID	(4R)-4-[(2S,5R,7R,14R,15R,16S)-5,16-dihydroxy-2,15-dimethyltetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]heptadecan-14-yl]pentanoic acid

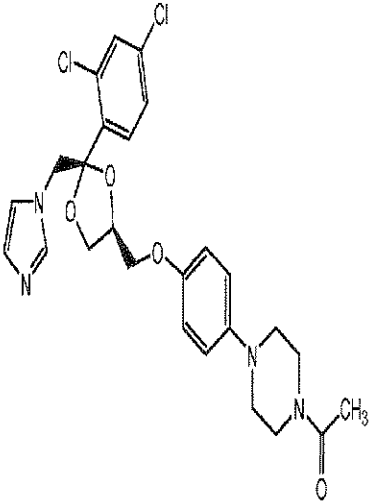
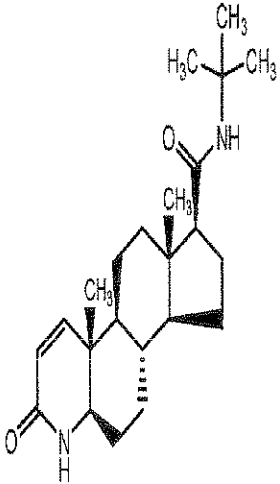
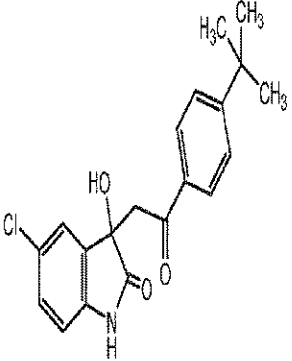
STRUCTURE	CCG ID	NAME	IUPAC
	38703	ENOXOLONE	(2S,4aS,6aS,6bR,10S,12aS,14bR)-10-hydroxy-2,4a,6a,6b,9,9,12a-heptamethyl-13-oxo-1,2,3,4,4a,5,6,6a,6b,7,8,8a,9,10,11,12,12a,12b,13,14b-icosahydricene-2-carboxylic acid
	38719	beta-ESCIN	(2S,3S,4S,5R,6R)-6-([(3S,4S,6aR,6bS,8R,8aR,9R,10S,14bR)-9-(acetyloxy)-8-hydroxy-4,8a-bis(hydroxymethyl)-4,6a,6b,11,11,14b-hexamethyl-10-[(2Z)-2-methylbut-2-enoyl]oxy)-1,2,3,4,4a,5,6,6a,6b,7,8,8a,9,10,11,12,12a,14,14a,14b-icosahydricen-3-yl]oxy)-4-hydroxy-3,5-bis([(2S,3R,4S,5S,6R)-3,4,5-trihydroxy-6-(hydroxymethyl)oxan-2-yl]oxy))oxane-2-carboxylic acid
	38826	DIGOXIN	4-[(2S,5S,7R,11S,14R,15S,16R)-5-([(2R,4S,5S,6R)-5-([(2S,4S,5S,6R)-5-([(2S,4S,5S,6R)-4,5-dihydroxy-6-methyloxan-2-yl]oxy)-4-hydroxy-6-methyloxan-2-yl]oxy)-4-hydroxy-6-methyloxan-2-yl]oxy)-11,16-dihydroxy-2,15-dimethyltetracyclo[8.7.0.0^(2,7).0^(11,15)]heptadecan-14-yl]-2,5-dihydrofuran-2-one

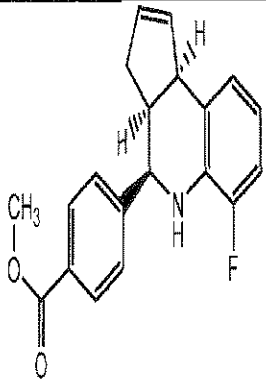
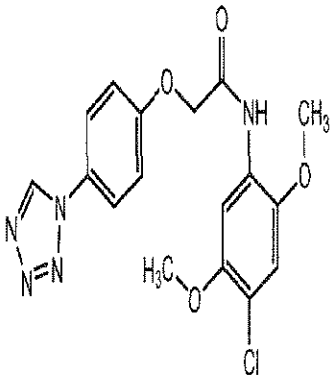
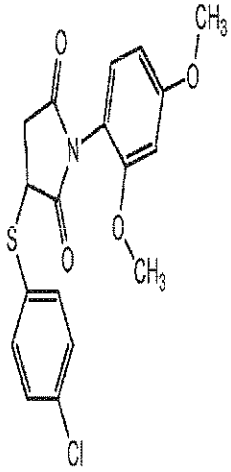
STRUCTURE	CCG ID	NAME	IUPAC
	38850	LITHOCHOLIC ACID	(4R)-4-[(2S,5R,7R,14R,15R)-5-hydroxy-2,15-dimethyltetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]]heptadecan-14-yl]pentanoic acid
	38852	GITOXIN	4-[(2S,5S,7R,10R,11S,13S,14R,15R)-5-([(2R,4S,5S,6R)-5-([(4S,5S,6R)-5-([(4S,6R)-4,5-dihydroxy-6-methyloxan-2-yl]oxy)-4-hydroxy-6-methyloxan-2-yl]oxy)-4-hydroxy-6-methyloxan-2-yl]oxy)-11,13-dihydroxy-2,15-dimethyltetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]]heptadecan-14-yl]-2,5-dihydrofuran-2-one
	38857	GLYCYRRHIZIC ACID	(2S,3S,4R,5R,6R)-6-([(3S,6aS,8aS,11S,12aR,14bS)-11-carboxy-4,4,6a,8a,11,14b-hexamethyl-14-oxo-1,2,3,4,4a,5,6,6a,6b,7,8,8a,9,10,11,12,12a,14,14a,14b-icosahydricen-3-yl]oxy)-3-([(2R,3R,4S,5S,6S)-6-carboxy-3,4,5-trihydroxyoxan-2-yl]oxy)-4,5-dihydroxyoxane-2-carboxylic acid

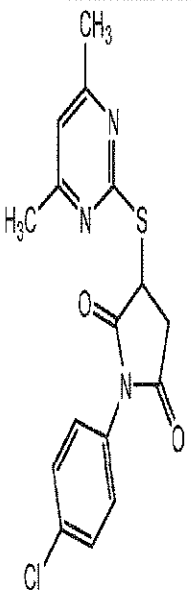
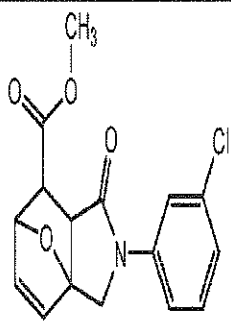
STRUCTURE	CCG ID	NAME	IUPAC
	38866	LANATOSIDE C	(2R,3R,4S,6S)-6-[[[(2R,3S,4S,6S)-6-[[[(2R,3S,4S,6R)-6-[[[(2S,5S,11S,14R,15S,16R)-11,16-dihydroxy-2,15-dimethyl-14-(5-oxo-2,5-dihydrofuran-3-yl)tetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]]heptadecan-5-yl]oxy}-4-hydroxy-2-methyloxan-3-yl]oxy}-4-hydroxy-2-methyloxan-3-yl]oxy}-2-methyl-3-[[[(2S,3R,4S,5S,6R)-3,4,5-trihydroxy-6-(hydroxymethyl)oxan-2-yl]oxy]oxan-4-yl acetate
	38894	CAFESTOL ACETATE	[(1S,12R,16R,17R)-17-hydroxy-12-methyl-8-oxapentacyclo[14.2.1.0 ^{1,13} .0 ^{4,12} .0 ^{5,9}]]nonadeca-5(9),6-dien-17-yl]methyl acetate
	38938	KETOCONAZOLE	1-[4-(4-[[[(2R)-2-(2,4-dichlorophenyl)-2-(1H-imidazol-1-ylmethyl)-1,3-dioxolan-4-yl]methoxy]phenyl)piperazin-1-yl]ethan-1-one
	39166	CHOL-11-ENIC ACID	(4R)-4-[(2S,7R,14R,15R)-2,15-dimethyl-5-oxotetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]]heptadec-16-en-14-yl]pentanoic acid

STRUCTURE	CCG ID	NAME	IUPAC
 Cl⁻	39322	METHYLBENZETHONIUM CHLORIDE	benzyl(dimethyl(2-{2-[2-methyl-4-(2,4,4-trimethylpentan-2-yl)phenoxy]ethoxy}ethyl)azanium chloride
 Na⁺	39682	CARBENOXOLONE SODIUM	disodium (2S,4aS,6aS,6bR,10S,12aS,14bR)-10-[(3-carboxylatopropanoyl)oxy]-2,4a,6a,6b,9,9,12a-heptamethyl-13-oxo-1,2,3,4,4a,5,6,6a,6b,7,8,8a,9,10,11,12,12a,12b,13,14b-icosahydricene-2-carboxylate
	39958	18alpha-GLYCYRRHETINIC ACID	(2S,4aS,6aS,6bR,10S,12aS,14bS)-10-hydroxy-2,4a,6a,6b,9,9,12a-heptamethyl-13-oxo-1,2,3,4,4a,5,6,6a,6b,7,8,8a,9,10,11,12,12a,12b,13,14b-icosahydricene-2-carboxylic acid

STRUCTURE	CCG ID	NAME	IUPAC
	40120	METHYLDOPA	2-amino-3-(3,4-dihydroxyphenyl)-2-methylpropanoic acid
	40211	DOPAMINE HYDROCHLORIDE	4-(2-aminoethyl)benzene-1,2-diol hydrochloride
	40270	5alpha-CHOLESTAN- 3beta-OL-6-ONE	(2R,5S,7S,15R)-5-hydroxy-2,15-dimethyl-14-(6-methylheptan-2-yl)tetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]heptadecan-8-one
	40274	URSODIOL	(4R)-4-[(2S,5R,7S,9S,14R,15R)-5,9-dihydroxy-2,15-dimethyltetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]heptadecan-14-yl]pentanoic acid

STRUCTURE	CCG ID	NAME	IUPAC
	100815	CPD000058460_KETO CONAZOLE	1-[4-(4-[[[(2R,4S)-2-(2,4-dichlorophenyl)-2-(1H-imidazol-1-ylmethyl)-1,3-dioxolan-4-yl]methoxy]phenyl]piperazin-1-yl]ethan-1-one
	100937	CPD000466304_ FINASTERIDE	(1S,2R,7R,10S,11S,14S,15S)-N-tert-butyl-2,15-dimethyl-5-oxo-6-azatetracyclo[8.7.0.0 ^{2,7} .0 ^{11,15}]]heptadec-3-ene-14-carboxamide
	22428		3-[2-(4-tert-butylphenyl)-2-oxoethyl]-5-chloro-3-hydroxy-2,3-dihydro-1H-indol-2-one

STRUCTURE	CCG ID	NAME	IUPAC
	22775		methyl 4-[(2R,6S,7R)-10-fluoro-8-azatricyclo[7.4.0.0 ^{2,6}]]trideca-1(13),3,9,11-tetraen-7-yl]benzoate
	108668		N-(4-chloro-2,5-dimethoxyphenyl)-2-[4-(1H-1,2,3,4-tetrazol-1-yl)phenoxy]acetamide
	109888		3-[(4-chlorophenyl)sulfanyl]-1-(2,4-dimethoxyphenyl)pyrrolidine-2,5-dione

STRUCTURE	CCG ID	NAME	IUPAC
	112216		1-(4-chlorophenyl)-3-[(4,6-dimethylpyrimidin-2-yl)sulfanyl]pyrrolidine-2,5-dione
	116781		methyl 3-(3-chlorophenyl)-4-oxo-10-oxa-3-azatricyclo[5.2.1.0 ^{1,5}]dec-8-ene-6-carboxylate

Pharmaceutical compositions

[0043] The invention further comprises a composition comprising the compound of any one of structural formulae I, IA, IB, or II, or a compound listed in Table 1, and a pharmaceutically acceptable ophthalmic carrier, e.g., a pharmaceutically acceptable excipient, carrier, binder, and/or diluent. Optionally, the composition includes a free acid, free base, salt (e.g., an acid or base addition salt), hydrate or prodrug of the compound of any one of structural formula I, IA, IB, or II, or a compound listed in Table 1. The prodrug is a material that includes the compound in any one of structural formulae I, IA, IB, or II, or a compound listed in Table 1, covalently bound to a carrier moiety. The carrier moiety can be released from the compound in any one of structural formulae I, IA, IB, or II, or a compound listed in Table 1, *in vitro* or *in vivo* to yield compound in any one of structural formula I, IA, IB, or II, or a compound listed in Table 1. Prodrug forms are well known in the art as exemplified in Sloan, K. B., *Prodrugs*, M. Dekker, New York, 1992; and Testa, B. and Mayer, J. M., *Hydrolysis in drug and prodrug metabolism: chemistry, biochemistry, and enzymology*, Wiley-VCH, Zurich, 2003.

[0044] In various embodiments, the composition comprises the compound of any one of structural formulae I, IA, IB, or II, or a compound listed in Table 1, formulated as eye drops, injectable solutions or eye ointments. These pharmaceutical compositions can be formulated by admixing, diluting or dissolving the compound, optionally, with appropriate pharmaceutical additives such as excipients, disintegrators, binders, lubricants, diluents, buffers, antiseptics, moistening agents, emulsifiers, dispersing agents, stabilizing agents and dissolving aids in accordance with conventional methods and formulating in a conventional manner depending upon the dosage form. For example, eye drops can be formulated by dissolving the compound in sterilized water in which a surface active agent is dissolved and optionally adding appropriate pharmaceutical additives such as a preservative, a stabilizing agent, a buffer, an antioxidant and a viscosity improver. In some embodiments, the composition includes a cyclodextrin. In a specific embodiment, the cyclodextrin is (2-hydroxypropyl)- β -cyclodextrin.

[0045] Physiologically acceptable buffers include, but are not limited to, a phosphate buffer or a Tris-HCl buffer (comprising tris(hydroxymethyl)aminomethane and HCl). For example, a Tris-HCl buffer having pH of 7.4 comprises 3 g/l of

tris(hydroxymethyl)aminomethane and 0.76 g/l of HCl. In yet another aspect, the buffer is 10x phosphate buffer saline ("PBS") or 5x PBS solution.

[0046] Other buffers include, but are not limited to, buffers based on HEPES (N-{2-hydroxyethyl}peperazine-N'-(2-ethanesulfonic acid)) having pK_a of 7.5 at 25°C and pH in the range of about 6.8-8.2; BES (N,N-bis{2-hydroxyethyl}2-aminoethanesulfonic acid) having pK_a of 7.1 at 25°C and pH in the range of about 6.4-7.8; MOPS (3-{N-morpholino}propanesulfonic acid) having pK_a of 7.2 at 25°C and pH in the range of about 6.5-7.9; TES (N-tris{hydroxymethyl}-methyl-2-aminoethanesulfonic acid) having pK_a of 7.4 at 25°C and pH in the range of about 6.8-8.2; MOBS (4-{N-morpholino}butanesulfonic acid) having pK_a of 7.6 at 25°C and pH in the range of about 6.9-8.3; DIPSO (3-(N,N-bis{2-hydroxyethyl}amino)-2-hydroxypropane)) having pK_a of 7.52 at 25°C and pH in the range of about 7-8.2; TAPSO (2-hydroxy-3{tris(hydroxymethyl)methylamino}-1-propanesulfonic acid)) having pK_a of 7.61 at 25°C and pH in the range of about 7-8.2; TAPS ((2-hydroxy-1,1-bis(hydroxymethyl)ethyl)amino)-1-propanesulfonic acid)) having pK_a of 8.4 at 25°C and pH in the range of about 7.7-9.1; TABS (N-tris(hydroxymethyl)methyl-4-aminobutanesulfonic acid) having pK_a of 8.9 at 25°C and pH in the range of about 8.2-9.6; AMPPO (N-(1,1-dimethyl-2-hydroxyethyl)-3-amino-2-hydroxypropanesulfonic acid)) having pK_a of 9.0 at 25°C and pH in the range of about 8.3-9.7; CHES (2-cyclohexylamino)ethanesulfonic acid) having pK_a of 9.5 at 25°C and pH in the range of about 8.6-10.0; CAPSO (3-(cyclohexylamino)-2-hydroxy-1-propanesulfonic acid) having pK_a of 9.6 at 25°C and pH in the range of about 8.9-10.3; and CAPS (3-(cyclohexylamino)-1-propane sulfonic acid) having pK_a of 10.4 at 25°C and pH in the range of about 9.7-11.1.

[0047] A number of effective methods for controlled release of an active agent are available. See, for example, Wagh V. D., Inamdar B., Samanta M. K., Polymers used in ocular dosage form and drug delivery systems. *Asian J Pharm* 2, 2008, 12-17 and the literature references cited therein, the contents of which are incorporated herein by reference. The use of polymers (e.g., cellulose derivatives such as hydroxypropylmethylcellulose (HPMC) and hydroxypropylcellulose (HPC), poly(acrylic acid) (PAA), polyacrylates, cyclodextrins and natural gums, polyorthoesters (POEs) and mucoadhesive polymers); semisolids such as gels, films and other inserts; resins such as ion exchange resins; iontophoretic delivery; and colloidal particles such as microspheres and nanoparticles, are specifically contemplated.

[0048] The compounds of the invention may also be provided in combination with other therapeutic agents. In some embodiments, the compounds of the invention may be co-formulated with a pain reliever, an anesthetic, artificial tears, an enzyme inhibitor, a cytokine inhibitor, an anti-inflammatory, or an antibiotic. In some embodiments, the antibiotic is an antibacterial, antiviral, antifungal, antiprotozoal agent, or a combination thereof.

[0049] In various embodiments, the compounds of the invention may also be provided in combination with an ocular therapeutic selected from the group consisting of Acular (ketorolac tromethamine ophthalmic solution) 0.5%, Acuvail (ketorolac tromethamine), AK-Con-A (naphazoline ophthalmic), Akten (lidocaine hydrochloride), Alamast, Alphagan (brimonidine), Alrex, Astepro (azelastine hydrochloride nasal spray), AzaSite (azithromycin), Bepreve (bepotastine besilate ophthalmic solution), Besivance (besifloxacin ophthalmic suspension), Betaxon, BSS Sterile Irrigating Solution, Cosopt, Durezol (difluprednate), Eylea (aflibercept), Lotemax, Lucentis (ranibizumab), Lumigan (bimatoprost ophthalmic solution), Macugen (pegaptanib), Ocuflox (ofloxacin ophthalmic solution) 0.3%, OcuHist, Ozurdex (dexamethasone), Quixin (levofloxacin), Rescula (unoprostone isopropyl ophthalmic solution) 0.15%, Restasis (cyclosporine ophthalmic emulsion), Salagen Tablets, Travatan (travoprost ophthalmic solution), Valcyte (valganciclovir HCl), Viroptic, Vistide (cidofovir), Visudyne (verteporfin for injection), Vitrasert Implant, Vitravene Injection, ZADITOR, Zioptan (tafluprost ophthalmic solution), Zirgan (ganciclovir ophthalmic gel), Zymaxid (gatifloxacin ophthalmic solution), Atropine, Flurbiprofen, Physostigmine, Azopt, Gentamicin, Pilocarpine, Bacitracin, Goniosol, Polymyxin B, Betadine, Gramicidin, Prednisolone, Betaxolol, Humorsol, Proparacaine, Betoptic, Hylartin, Propine, Brinzolamide, Hypertonic NaCl, Puralube, BSS, Indocyanine Green, Rose Bengal, Carbachol, Itraconazole, Sodium Hyaluronate, Cefazolin, Latanoprost, Suprofen, Celluvisc, Mannitol, Terramycin, Chloramphenicol, Methazolamide, Timolol, Ciloxan, Miconazole, Tobramycin, Ciprofloxacin, Miostat, Triamcinolone, Cosopt, Muro 128, Trifluridine, Demecarium, Neomycin, Tropicamide, Dexamethasone, Neptazane, Trusopt, Dipivefrin, Ocuflox, Vidarabine, Dorzolamide, Ofloxacin, Vira-A, Epinephrine, Oxytetracycline, Viroptic, Fluorescein, Phenylephrine, and Xalatan.

Screening methods and systems

[0050] In various embodiments, the invention includes a high-throughput method of screening compounds for modulation of protein thermal stability. The method comprises (a)

contacting a protein with each of a plurality of test compounds; and (b) measuring the melting transition (T_m) of the protein in the presence of each of the plurality of test compounds, wherein a compound that decreases or increases the apparent T_m by at least 2 standard deviations is a pharmacological protein chaperone. In various embodiments, a compound that decreases or increases the apparent T_m by at least 3 standard deviations is a pharmacological protein chaperone.

[0051] An agent that modulates protein thermal stability can be identified using differential scanning fluorimetry (DSF). In DSF, the melting transition (T_m) of the protein target is measured in the presence of potential ligand. DSF measures thermal unfolding of the target protein via the fluorescence of intrinsic tryptophans, or a dye, such as 1,8-anilino-naphthalene-6-sulfonic acid (ANS) or Sypro Orange. Binding of a ligand adds free energy to the folded state or key intermediates, which limits unfolding and shifts the apparent T_m . Thus, in some embodiments, the melting transition is determined using a high-throughput differential scanning fluorimetry device, such as a ThermoFluor® 384-well DSF platform, or a real-time PCT thermocycler.

[0052] In various embodiments of the screening method, the step wherein the melting transition is measured comprises the following steps: (b1) heating the protein in the presence of each of a plurality of test compounds from 50°C to 80°C, (b2) cooling the protein to 25°C, (b3) maintaining the protein at 25°C for 10 seconds, and (b4) measuring the fluorescence of the protein. In more specific embodiments, the method further comprises repeating steps (b1)-(b4) between 2 and 30 times, wherein each repeat of step (b1) is performed at an incrementally higher temperature. For example, during the first iteration, the protein is heated to 65°C in the presence of a test compound and cooled to 25°C where it is maintained for about 10 seconds before the fluorescence of the protein is measured. During the second iteration, the protein is heated to 66°C in the presence of a test compound and cooled to 25°C where it is maintained for about 10 seconds before the fluorescence of the protein is measured. This process is repeated (e.g., between 2 and 30 times) while increasing the peak temperature to which the protein is heated, e.g., in 1°C increments. In some embodiments, during step (b1), the protein is equilibrated at the peak temperature between 60 and 180 seconds. In specific embodiments of the method, the equilibrating step is 130 seconds.

[0053] In various embodiments of the screening method, the step wherein the melting transition is measured comprises gradually heating the protein in the presence of each of a

plurality of test compounds to 80°C while continuously measuring the fluorescence of the protein.

[0054] In various aspects of the screening method, the protein is an amyloid-forming protein. Generally, when the protein is an amyloid-forming protein, the desired modulator decreases the melting transition of the protein. Such a modulator stabilizes the non-amyloid form of the protein. In some embodiments, the amyloid-forming protein is selected from the group consisting of Hsp27, α A-crystallin (cataract), α B-crystallin (cataract), β B2-crystallin (cataract), β B1-crystallin (cataract), γ D-crystallin (cataract), Hsp22, Hsp20, tau, Alpha-synuclein (Parkinson's disease), IAPP (Diabetes mellitus type 2), beta-amyloid (Alzheimer's disease), PrP (Transmissible spongiform encephalopathy), Huntingtin (Huntington's Disease), Calcitonin (Medullary carcinoma of the thyroid), Atrial natriuretic factor (Isolated atrial amyloidosis), Apolipoprotein AI (Atherosclerosis), Serum amyloid A (Rheumatoid arthritis), Medin (Aortic medial amyloid), Prolactin (Prolactinomas), Transthyretin (Familial amyloid polyneuropathy), Lysozyme (Hereditary non-neuropathic systemic amyloidosis), Beta 2 microglobulin (Dialysis related amyloidosis), Gelsolin (Finnish amyloidosis), Keratoepithelin (Lattice corneal dystrophy), Cystatin (Cerebral amyloid angiopathy: Icelandic type), Immunoglobulin light chain AL (systemic AL amyloidosis), myocilin (glaucoma), and S-IBM (Sporadic Inclusion Body Myositis).

[0055] In some embodiments of the screening method, the protein is a protein underlying a loss-of-function disease. Generally, when the protein is a protein underlying a loss-of-function disease, the desired modulator increases the melting transition of the protein. Such a modulator stabilizes the mutant form of the protein such that it remains active (i.e., degradation of the protein is reduced). The protein underlying a loss-of-function disease can be, for example, mutant β -glucosidase, mutant glucosylceramidase (Gaucher's disease), mutant cystic fibrosis transmembrane receptor (cystic fibrosis), mutant hexosaminidase A (Tay-Sachs disease), mutant hexosaminidase B (Sandhoff disease), mutant β -galactosidase (Morquio syndrome), and mutant alpha-glucosidase (i.e., Pompe disease).

[0056] In some embodiments, the invention includes a high-throughput screening system, comprising: (a) an amyloid-forming protein; (b) a device capable of measuring the melting transition (T_m) of the amyloid-forming protein; and (c) a plurality of test compounds. The device capable of measuring the T_m of the amyloid-forming protein can be any device known in the art. As noted above, in some embodiments, the device is a differential scanning

fluorimetry device, such as a ThermoFluor® 384-well DSF platform. In various aspects of the screening system, the protein is selected from the group consisting of Hsp27, α A-crystallin, α B-crystallin, β B2-crystallin, β B1-crystallin, γ D-crystallin, Hsp22, Hsp20, tau, Alpha-synuclein, IAPP, beta-amyloid, PrP, Huntingtin, Calcitonin, Atrial natriuretic factor, Apolipoprotein AI, Serum amyloid A, Medin, Prolactin, Transthyretin, Lysozyme, Beta 2 microglobulin, Gelsolin, Keratoepithelin, Cystatin, Immunoglobulin light chain AL, and S-IBM.

EXAMPLES

[0057] The following are non-limiting examples of various aspects of the methods described herein. The examples are given solely for the purpose of illustration and are not to be construed as limiting the disclosure, as many variations thereof are possible.

EXAMPLE 1

[0058] This Example describes a high-throughput method of screening for compounds that stabilize an exemplary aggregation-prone protein, Hsp27.

[0059] Differential scanning fluorimetry (DSF) was utilized to identify compounds that stabilize cryAB. In DSF experiments, the melting transition (T_m) of the protein target is measured in the presence of potential ligands (Cummings et al., *J Biomol Screen* **11**, 854 (2006)). DSF measures thermal unfolding of the target protein via the fluorescence of intrinsic tryptophans, or (more commonly) a dye, such as 1,8-anilinonaphthalenesulfonic (bis-ANS) or Sypro Orange. Binding of a ligand adds free energy to the folded state or key intermediates, which limits unfolding and shifts the apparent T_m . These values are used to calculate the unfolding transition. When compounds bind to the target prior to heating, they stabilize against unfolding, shifting the ΔT_m . For the case of cryAB and the cryAB R120G mutant, compounds that reduced the apparent T_m were sought, because amyloids typically have a higher apparent stability in this platform. For example, the T_m of wild type cryAB was $64.1 \pm 0.5^\circ\text{C}$, whereas the more amyloid-prone R120G cryAB mutant had an apparent T_m of $68.3 \pm 0.2^\circ\text{C}$ (3). This difference correlated with the increased amyloid propensity of R120G. Thus, it was reasoned that the screening “hits” which reduce apparent T_m might suppress aggregation.

[0060] Using a miniaturized 384-well DSF platform (ThermoFluor®), 2,446 compounds were screened for the ability to block aggregation of an sHSP. As a model, human Hsp27 was used for the HTS because it readily forms amyloids and it retains the highly conserved crystallin domain found in all sHSPs. Hsp27 was soluble at room temperature, but it thermally unfolded and rapidly formed stable amyloids upon heating.

[0061] The screen was performed in 7 μ L volume, with 10 μ M Hsp27 in 50 mM NaPO₄, pH 7.4, 700 mM NaCl, 50 mM LiCl, and 100 μ M bis-ANS in Abgene black 384-well PCR plates. Reactions were covered with silicon oil to limit evaporation. Control tests suggested that the assay tolerated up to 4% DMSO but 1% was used as a final concentration. The plates were measured in up/down mode, which was empirically determined to give better signal-to-noise than the continuous ramp mode. Plates were heated from 65°C to 80°C in 1°C increments, equilibrated for 130 seconds at each high temperature, cooled to 25°C, and held for 10 seconds at 25°C prior to imaging with a single 10 second exposure for each temperature reading. Plate uniformity tests measured the T_m of Hsp27 at 72.3 ± 0.16 °C. The Z factor was calculated to vary between ~0.59 and 0.71 and the CVs were 8%. Most of the variability appeared to arise from some “edge effects”

[0062] For each well, a series of fluorescence vs. temperature points were obtained, plotted and fit to determine the T_m . An automated method was developed in MatLab that performs the individual fits and then ranks each test well on the basis of how closely the three independent fitting methods (Savitzky-Golay derivative, sigmoidal curve and parallel baseline Hill) agree.

[0063] From the primary screen, 45 compounds were identified that decreased the apparent T_m by more than three standard deviations ($-3SD$; $\sim 0.6^\circ C$) and 45 compounds increased the T_m by $+3SD$. All 90 of these compounds were explored in dose dependence experiments to confirm their activity and, from those studies, 64 had activity below 50 μ M. The 45 compounds identified that decreased the apparent T_m by more than three standard deviations were tested in 12-point dose dependence assays, providing 28 that both confirmed and shifted ΔT_m at concentrations less than 20 μ M. Interestingly, 12 of these 28 were part of a single structural class of related sterols. This scaffold was selected for further investigation.

[0064] Taken together, these studies validate the HTS method as a robust means for identifying pharmacological chaperones for amyloid-forming proteins, such as cryAB.

Indeed, the studies validate the HTS method for finding agents that modulate T_m for any protein wherein increased (e.g., loss of function proteins such as the mutant form of alpha-glucosidase responsible for Pompe Disease) or decreased stability (aggregation-prone proteins such as Huntingtin) is desired.

EXAMPLE 2

[0065] This Example describes structure-activity relationship (SAR) studies of a sterol scaffold identified in the primary screen of Example 1. As described further below, the SAR revealed a general chemical structure (Formula I) for compounds that reduced the T_m of an amyloid-forming protein, R120G cryAB, by at least 2°C.

[0066] Thirty-two sterols with chemical structures related to 17-a-hydroxy-progesterone were collected and subjected to further testing (see Table 2). DSF experiments on this focused collection were performed using R120G cryAB and two compounds that reduced T_m by at least 2°C were identified: 5a-cholestan-3b-ol-6-one and 5-cholesten-3b,25-diol. Many of the other closely-related sterols, including cholesterol, were inactive and the resulting SAR supported a rather specific molecular interaction.

Table 2

NAME	ACTIVITY (shift in thermal stability at 100 μ M)
5-cholesten-3b,25-diol	-2.0°C
5a-cholestan-3b-ol-6-one	-3.0°C
5-cholesten-3b-ol	-1.1°C
etiocholan-17b-ol-3-one	+1.7°C
17-a-hydroxy-progesterone	between +1°C and -1°C
5a-androstan-3b,17b-diol	between +1°C and -1°C
17a-hydroxy-pregnenolone	between +1°C and -1°C
5-a-androstan-3,17-dione	between +1°C and -1°C
epiandrosterone	between +1°C and -1°C
progesterone	between +1°C and -1°C
dehydroisoandrosterone	between +1°C and -1°C
D4-androsten-3,17-dione	between +1°C and -1°C
b-estradiol	between +1°C and -1°C
testosterone	between +1°C and -1°C
corticosterone	between +1°C and -1°C
cortisone	between +1°C and -1°C
estriol	between +1°C and -1°C
4-cholesten-3-one	between +1°C and -1°C
cholesterol	between +1°C and -1°C
hydrocortisone	between +1°C and -1°C
estrone	between +1°C and -1°C

NAME	ACTIVITY (shift in thermal stability at 100 μ M)
D5-pregnen-3b-ol-20-one	between +1°C and -1°C
testosterone acetate	between +1°C and -1°C
deoxycorticosterone	between +1°C and -1°C
androsterone	between +1°C and -1°C
corticosterone-21-acetate	between +1°C and -1°C
digitonin	between +1°C and -1°C
5-cholesten-3b-ol-7-one	between +1°C and -1°C
ursodeoxycholic acid	between +1°C and -1°C
6-ketocholestanol	between +1°C and -1°C
18-a-glycyrrhetic acid	between +1°C and -1°C

[0067] Further dose dependence studies with 5a-cholestan-3b-ol-6-one, 5-cholesten-3b,25-diol, and cholesterol showed that both 5a-cholestan-3b-ol-6-one and 5-cholesten-3b,25-diol were able to partially recover the wild type T_m , indicating their utility as pharmacological chaperones. To confirm the direct interaction with cryAB in a distinct experimental platform, biolayer interferometry (described in Example 1) was used to determine that 5-cholesten-3b,25-diol binds to R120G cryAB with a K_D of $10.1 \pm 4.4 \mu\text{M}$. Measuring affinity for the test protein by biolayer interferometry (Octet Red) allows for the identification of false positives. Briefly, biotinylated cryAB is immobilized to streptavidin-coated pins and equilibrium association data across a hundred-fold concentration range is analyzed to generate the apparent K_D . To control for nonspecific binding, the response of a biocytin-blocked pin is subtracted from each sensorgram.

[0068] Taken together, the results above demonstrate that compounds of Formula I function as pharmacological chaperones for R120G cryAB *in vitro* and bind with high affinity. Moreover, the results indicate that the disclosed DSF methods represent a robust high-throughput screening system for identifying novel pharmacological chaperones for non-enzymes.

EXAMPLE 3

[0069] This Example demonstrates that the compounds identified in Example 1 suppress amyloid formation. Significantly, the compounds are also able to *reverse* amyloid formation.

[0070] The DSF studies suggested that sterols suppress R120G cryAB amyloid formation. To test this idea, R120G cryAB (15 μM) was treated with 5-cholesten-3b,25-diol or cholesterol (100 μM) and its ability to aggregate was measured by electron microscopy. These studies confirmed that 5-cholesten-3b,25-diol, but not cholesterol or the solvent

control, dramatically suppressed amyloid formation. Visual inspection of the solutions supported this conclusion, because only 5-cholesten-3b,25-diol reduced the opacity of the R120G cryAB mixture. Moreover, 5-cholesten-3b,25-diol also reversed the aggregation of pre-formed R120G cryAB amyloids, suggesting that it shifts the equilibrium towards non-amyloid structures.

[0071] To explore the mechanism of action of 5-cholesten-3b,25-diol, the binding site on R120G cryAB was explored by ^{15}N HSQC NMR, and analysis of the resulting chemical shifts suggested that it binds an exposed face of the conserved crystallin domain. Specifically, 5-cholesten-3b,25-diol binds a groove across the cryAB dimer interface and makes contacts with residues near R120 and D109.

[0072] The above results support a model in which the sterol stabilizes the sHSP and reduces amyloid formation.

EXAMPLE 4

[0073] This Example demonstrates that compounds identified herein reverse the cataract phenotype *in vivo*.

[0074] The R120G cryAB knock-in mouse is a clinically-acceptable model of aging-associated and hereditary cataract. The animal model develops severe cataracts within 20 weeks (Andley et al., *PLoS One* **6**, e17671 (2011)). Excised eyes from these mice were treated with 5-cholesten-3b,25-diol (100 μM) or a saline control. 5-cholesten-3b,25-diol treatment significantly reduced the aggregation of cryAB and improved cryAB solubility.

[0075] 5-cholesten-3b,25-diol was then formulated in a saline-cyclodextrin solution (5 mM cyclodextran; 5 mM 5-cholesten-3b,25-diol) and delivered by eye dropper three times a week for two weeks to live animals ($n = 15$). cryAB solubility and lens transparency were determined by measuring slit lens illumination in live animals and by gel permeation chromatography/light scattering from lens homogenates. This treatment significantly improved cryAB solubility, improved lens transparency and rescued cataract phenotypes in 10/15 treated R120G cryAB knock-in mice. Thus, 5-cholesten-3b,25-diol, and more generally, a compound of formula I, is a promising therapeutic for the treatment of cataract. More broadly, these studies suggest that DSF-based HTS campaigns can be used to identify pharmacological chaperones for non-enzymes, such as cryAB.

EXAMPLE 5

[0076] This Example demonstrates that compounds identified herein increase levels of soluble α -crystallin in the eye.

[0077] 5-cholesten-3b,25-diol formulated in a saline-cyclodextrin solution (5 mM cyclodextran; 5 mM 5-cholesten-3b,25-diol) was delivered by eye dropper three times a week for two weeks to live mice (n = 15). Gel filtration analysis of α -crystallin solubility was conducted by processing samples as previously described by Andley et al. (*PLoS ONE*, 6:3, 1-13 (2011)). Briefly, degree of aggregation of α -crystallin was determined using gel permeation chromatography (GPC) with light scattering and refractive index (RI) measurements. All data were analyzed using SAS version 9.3 (SAS Institute; Cary, NC). To control for differences in assay sensitivity among animals, the area under the curve for α -crystallin in the drug treated eye was compared to the area under the curve generated from the fellow untreated eye and expressed as “percent” difference in area using the untreated eye as the denominator. The percent difference between treated and untreated eyes was compared using the Wilcoxon signed rank test for the null hypothesis of no difference. The association between age and percent difference was calculated using the Spearman rank correlation coefficient. Non-parametric statistical models were used to protect from departures from normality.

[0078] The area under the curve of α -crystallin was found to be statistically significantly higher in the drug treated eyes versus the fellow untreated eyes (n=17, mean percent difference= 381.2 ± 974.6 , median = 63.3, Wilcoxon signed rank p-value = 0.001). Older mice demonstrated a higher percentage increase in α -crystallin in treated eyes relative to untreated eyes (Spearman rank correlation = 0.44, p=0.075). The data in Table 1 show a noticeable increase in median percent difference when the age is greater than 200 days.

		Age (days)						All
		66	79	184	192	200	266	
Percent difference in area between untreated and treated eyes	N	3	1	2	3	4	4	17
	Mean	177.2	98.1	61.4	22.7	38.7	1376.3	381.2
	Standard Deviation	347.9	.	2.7	68.4	25.9	1799.8	974.6
	Median	-20.2	98.1	61.4	55.8	31.8	679.0	63.3

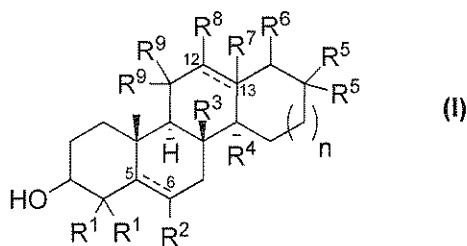
[0079] The data reveal that a compound of formula I, 5-cholesten-3 β ,25-diol, significantly increased solubility of α -crystallin *in vivo*, particularly in aged subjects, thereby targeting an underlying mechanism of cataract formation.

[0080] Every document cited herein, including any cross referenced or related patent or application, is hereby incorporated herein by reference in its entirety unless expressly excluded or otherwise limited. The citation of any document is not an admission that it is prior art with respect to any invention disclosed or claimed herein or that it alone, or in any combination with any other reference or references, teaches, suggests or discloses any such invention. Further, to the extent that any meaning or definition of a term in this document conflicts with any meaning or definition of the same term in a document incorporated by reference, the meaning or definition assigned to that term in this document shall govern.

[0081] While particular embodiments of the disclosed subject matter have been illustrated and described, it would be apparent to those skilled in the art that various other changes and modifications can be made without departing from the spirit and scope of the subject matter claimed. It is therefore intended to cover in the appended claims all such changes and modifications.

WHAT IS CLAIMED:

1. Use of a composition comprising a therapeutically effective amount of a compound of formula I for the manufacture of a medicament for treating or preventing cataract in an individual in need thereof:



wherein:

each R^1 is H or each R^1 is Me;

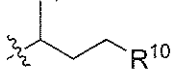
R^2 is H or OH;

the dashed line between carbons 5 and 6 indicates an optional double bond;

R^3 is H or Me;

R^4 is H or Me;

n is 0 or 1;

R^6 is  and each R^5 is independently H or Me;

the dashed line between carbons 12 and 13 is an optional double bond,

R^7 is H or Me, with the proviso that R^7 is not present when the double bond between carbons 12 and 13 is present;

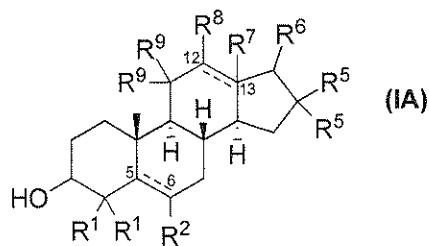
R^8 is H or OH;

both R^9 groups together form an oxo (=O) or each R^9 is hydrogen; and

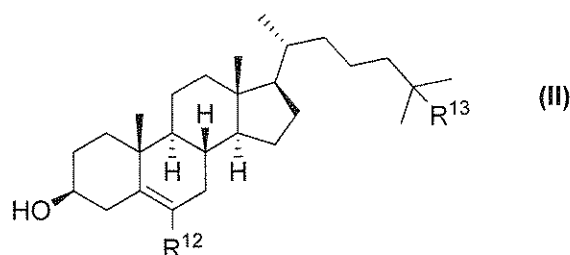
R^{10} is branched C_1 - C_6 alkyl, substituted with a hydroxyl group;

or pharmaceutically acceptable salt thereof.

2. The use according to claim 1 wherein the compound of formula I has a structure of formula IA:



3. The use according to claim 1 or 2 wherein the compound has a structure of formula II:



wherein R^{12} is H or OH and R^{13} is OH.

4. The use according to claim 3 wherein the compound is 5-cholesten-3b,25-diol.
5. Use of 5a-cholestan-3b-ol-6-one for the manufacture of a medicament for treating or preventing cataract.
6. The use according to any one of claims 1-5, wherein the medicament is to be administered topically, subconjunctivally, retrobulbarly, periocularly, subretinally, suprachoroidally, or intraocularly.
7. The use according to any one of claims 1-6, wherein (a) said cataract is an age-related cataract or a diabetic cataract or (b) wherein the individual has a hereditary form of cataract with early onset.
8. The use according to claim 7(b), wherein the individual has a R120G mutation or a D109H mutation in cryAB.

9. The use of any one of claims 1-8 wherein the composition interacts with α A-crystallin or α B-crystallin, optionally wherein the interaction is a stabilization of α A-crystallin or α B-crystallin.