

(19) **DANMARK**



Patent- og
Varemærkestyrelsen

(12)

Oversættelse af europæisk patentskrift

(10) **DK/EP 3515212 T3**

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- (51) Int.Cl.: **A 23 L 33/155 (2016.01)** **A 23 L 33/17 (2016.01)** **A 61 K 31/59 (2006.01)**
A 61 K 31/593 (2006.01) **A 61 K 38/17 (2006.01)** **C 07 K 14/435 (2006.01)**
- (45) Oversættelsen bekendtgjort den: **2024-08-26**
- (80) Dato for Den Europæiske Patentmyndigheds bekendtgørelse om meddelelse af patentet: **2024-07-24**
- (86) Europæisk ansøgning nr.: **17788336.0**
- (86) Europæisk indleveringsdag: **2017-09-25**
- (87) Den europæiske ansøgnings publiceringsdag: **2019-07-31**
- (86) International ansøgning nr.: **US2017053279**
- (87) Internationalt publikationsnr.: **WO2018058050**
- (30) Prioritet: **2016-09-23 US 201662398669 P**
- (84) Designerede stater: **AL AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HR HU IE IS IT LI LT LU LV MC MK MT NL NO PL PT RO RS SE SI SK SM TR**
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- (54) Benævnelse: **APOAEQUORIN OG D-VITAMIN INDEHOLDENDE SAMMENSÆTNINGER OG FREMGANGSMÅDER TIL ANVENDELSE DERAFT**
- (56) Fremdragne publikationer:
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DESCRIPTION

Description

FIELD OF THE INVENTION

[0001] This invention relates generally to compositions useful for the maintenance of calcium homeostasis. In particular, this invention is directed to apoaeguorin and vitamin D-containing compositions useful in preventing and/or alleviating diseases or symptoms associated with calcium imbalance and vitamin D deficiency.

BACKGROUND OF THE INVENTION

[0002] Calcium is the fifth most abundant element in the human body and occurs mainly in the bone. More than 99% of the calcium in the body is stored in the skeleton, which constantly exchanges its supply with the remaining 1 % dissolved in body fluids and soft tissue, such as the blood. The control of this exchange is largely dictated by the endocrine system which senses the concentration of ionized calcium in the plasma and directs calcium exchange to maintain this critical balance. Only a small fraction of the 1% of calcium in interstitial fluids and soft tissues is ionized and soluble. The remaining calcium in fluids and tissues is bound to proteins, particularly calcium-binding proteins (CaBPs). CaBPs are known to function in the maintenance of calcium homeostasis.

[0003] As the body requires specific concentrations of calcium ions to carry out requisite physiological processes, the maintenance of calcium homeostasis is of critical importance for bodily health. Proper ionic calcium concentrations in plasma and body fluids are understood by the medical community to be critical in bodily functions, including, but not limited to, neuronal excitability, muscle contraction, membrane permeability, cell division, hormone secretion and bone mineralization. A disruption in calcium homeostasis, i.e., a calcium imbalance, is associated with many diseases, syndromes and conditions, including, but not limited to, cancer, heart disease and neurodegenerative disease.

[0004] In the past, calcium channel antagonists, which block the flow of calcium between cell interiors and interstitial fluid, have been widely-prescribed as pharmaceutical agents useful in the prevention of calcium-related disorders including hypertension, angina, asthma, migraines and neural deterioration. For example, nimidopine has been found to improve clinical symptomatology and cognitive functions in dementia by alleviating a calcium imbalance which causes neural deterioration. However, many of these calcium channel antagonists have

unwanted side effects including, but not limited to, malaise, fluid retention, heartburn, erratic heart rate, dizziness, upset stomach and, in rare cases, fainting, fever and excessive bleeding.

[0005] Despite these advances, there is still a need for new and alternative therapeutics which alleviate or prevent calcium imbalance. In particular, pharmaceutical or nutraceutical compositions which have reduced side effects as compared to prior agents are desired and, if discovered, would meet a long felt need in the medical and nutritional health communities.

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"Jellyfish Protein Shows Potential to Help With Memory", INTERNET CITATION, 15 October 2008, pages 1-2, XP002528809, Retrieved from the Internet: URL:<http://www.reuters.com/article/pressRelease/idUS219580+15-Oct-2008+PRN20081015> [retrieved on 2009-05-19]; US 2014/294874 A1;

J. A. DETERT ET AL: "Pretreatment with Apoaequorin Protects Hippocampal CA1 Neurons from Oxygen-Glucose Deprivation", PLOS ONE, vol. 8, no. 11, 11 November 2013, page e79002, XP055425497, DOI: 10.1371/journal.pone.0079002 and CN 1 752 101 A describe the use of apoaequorin for the treatment of different diseases;

H. TANIURA ET AL: "Chronic vitamin D3 treatment protects against neurotoxicity by glutamate in association with upregulation of vitamin D receptor mRNA expression in cultured rat cortical neurons", JOURNAL OF NEUROSCIENCE RESEARCH., vol. 83, no. 7, 1 January 2006, pages 1179-1189, XP055426534, US, ISSN: 0360-4012, DOI: 10.1002/jnr.20824 and

KAJTA MET AL: "Neuroprotection by co-treatment and post-treating with calcitriol following the ischemic and excitotoxic insult in vivo and in vitro", NEUROCHEMISTRY INTERNATIONAL, vol. 55, no. 5, 1 October 2009, pages 265-274, XP026251694, ISSN: 0197-0186, DOI: 10.1016/J.NEUINT.2009.03.010 [retrieved on 2009-03-26] describe the use of vitamin D. The second document discloses that Vitamin D is protective against neuronal death caused by hypoxia.

SUMMARY OF THE INVENTION

[0006] The present invention, which is defined by the claims, provides compositions which are advantageous in the alleviation and/or prevention of symptoms or disorders associated with calcium imbalance and vitamin D deficiency. Such compositions include apoaequorin and vitamin D in combination with acceptable carriers for administration to a subject by a variety of routes.

[0007] Accordingly, the present invention is directed to compositions for use in treating a symptom or disorder associated with at least one of calcium imbalance and vitamin D deficiency, wherein the symptom or disorder is related to cell death following ischemia

comprising: (a) an effective amount of apoaeguorin; (b) an effective amount of vitamin D; and (c) a pharmaceutically acceptable carrier wherein: (a) the therapeutically effective amount of apoaeguorin in the unit dosage is about 10 mg to about 20 mg; and (b) the therapeutically effective amount of vitamin D in the unit dosage is about 25 mcg to about 75 mcg. In certain embodiments, nutraceutical compositions include, in addition to apoaeguorin and vitamin D, at least one other component recognized as providing nutraceutical benefit such as, for example, an immune boosting agent, anti-inflammatory agent, anti-oxidant agent, anti-viral agent, or a mixture thereof. Apoaeguorin and vitamin D compositions in certain embodiments are provided in a unit dosage form selected from a tablet, a capsule, a solution, a suspension, a syrup, a beverage, an oral or ophthalmic formulation or an injection.

[0008] Further described (not part of the present invention) is a method for treating a symptom or disorder associated with calcium imbalance and vitamin D deficiency, comprising administering to a subject in need of such treatment an effective amount of apoaeguorin and vitamin D.

[0009] Methods described herein are useful in treating a wide variety of symptoms or disorders associated with calcium imbalance and vitamin D deficiency, including but not limited to sleep quality, energy quality, mood quality, pain, memory quality. In certain examples, the calcium imbalance and vitamin D deficiency is physiologically-related to neuronal excitability, muscle contraction, membrane permeability, cell division, hormone secretion, bone mineralization, or cell death following ischemia. In such methods, apoaeguorin and vitamin D are preferably administered to the subject in the form of a nutraceutical composition.

[0010] Further described is the use of apoaeguorin and vitamin D for the manufacture of a nutraceutical composition for treating a symptom or disorder associated with calcium imbalance and vitamin D deficiency in a subject administered the nutraceutical composition. Exemplary symptoms or disorders treated by such compositions include those associated with sleep, energy, mood, pain, or memory.

[0011] The present invention further contemplates apoaeguorin and vitamin D for use in treating a symptom or disorder associated with calcium imbalance and vitamin D deficiency in a subject, including those symptoms or disorders associated with, e.g., sleep, energy, mood, pain, or memory in a subject.

[0012] The present invention provides various advantages over prior compositions and methods in that it provides for the general improvement of a subject's mental and physical health.

[0013] Other objects, features and advantages of the present invention will become apparent after review of the specification and claims.

BRIEF DESCRIPTION OF THE DRAWINGS

[0014]

Fig. 1 provides a graph showing the percent change from baseline of scores from areas: overall quality of sleep, energy, mood, pain and general health vs. days 0 through 90.

Fig. 2 depicts a graph showing data in which apoeaquorin (10mg) was taken daily by 56 participants. The participants were evaluated from eight days to 30 days. The memory study showed a statistically significant improvement in memory after 30 days ($p < .05$). 57% of participants had improvement in general memory, 51% in retaining information, 84% in remembering driving directions and 66% in word recall. N=56; 66% female, 34% male, mean age = 56 years; range 20-78 years.

Fig. 3 provides a graph showing the percent change, from baseline, of scores from standardized cognitive battery questionnaire vs. day 0 through 90.

Fig. 4 depicts neuroprotection by AQ (apoeaquorin) and Vitamin D in rat hippocampal brain slices.

DETAILED DESCRIPTION OF THE INVENTION

I. IN GENERAL

[0015] Before the present materials and methods are described, it is understood that this invention is not limited to the particular methodology, and materials described, as these may vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments only, and is not intended to limit the scope of the present invention which will be limited only by the appended claims.

[0016] It must be noted that as used herein and in the appended claims, the singular forms "a", "an", and "the" include plural reference unless the context clearly dictates otherwise. As well, the terms "a" (or "an"), "one or more" and "at least one" can be used interchangeably herein. It is also to be noted that the terms "comprising", "including", and "having" can be used interchangeably.

[0017] Unless defined otherwise, all technical and scientific terms used herein have the same meanings as commonly understood by one of ordinary skill in the art to which this invention belongs. Although any methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention, the preferred methods and materials are now described.

II. THE INVENTION

[0018] Aequorin is a photo-protein originally isolated from luminescent jellyfish and other marine organisms. The aequorin complex comprises a 22,285-dalton apoaequorin protein, molecular oxygen and the luminophore coelenterazine. When three Ca^{2+} ions bind to this complex, coelenterazine is oxidized to coelenteramide, with a concomitant release of carbon dioxide and blue light. Aequorin is not exported or secreted by cells, nor is it compartmentalized or sequestered within cells. Accordingly, aequorin measurements have been used to detect Ca^{2+} changes that occur over relatively long periods. In several experimental systems, aequorin's luminescence was detectable many hours to days after cell loading. It is further known that aequorin also does not disrupt cell functions or embryo development.

[0019] Because of its Ca^{2+} -dependent luminescence, the aequorin complex has been extensively used as an intracellular Ca^{2+} indicator. *Aequorea victoria* aequorin has been specifically used to: (1) analyze the secretion response of single adrenal chromaffin cells to nicotinic cholinergic agonists; (2) clarify the role of Ca^{2+} release in heart muscle damage; (3) demonstrate the massive release of Ca^{2+} during fertilization; (4) study the regulation of the sarcoplasmic reticulum Ca^{2+} pump expression in developing chick myoblasts; and (5) calibrate micropipets with injection volumes of as little as three picoliters.

[0020] Apoaequorin has an approximate molecular weight of 22 kDa. Apoaequorin can be used to regenerate aequorin by reducing the disulfide bond in apoaequorin. The calcium-loaded apoaequorin retains the same compact scaffold and overall folding pattern as unreacted photoproteins containing a bound substrate.

[0021] Conventional purification of aequorin from the jellyfish *Aequorea victoria* requires laborious extraction procedures and sometimes yields preparations that are substantially heterogeneous or that are toxic to the organisms under study. Two tons of jellyfish typically yield approximately 125mg of the purified photoprotein. In contrast, recombinant aequorin is preferably produced by purifying apoaequorin from genetically engineered *Escherichia coli*, followed by reconstitution of the aequorin complex *in vitro* with pure coelenterazine. Apoaequorin useful in the present invention has been described and is commercially-obtainable through purification schemes and/or syntheses known to those of skill in the art. S. Inouye, S. Zenno, Y. Sakaki, and F. Tsuji. High level expression and Purification of apoaequorin. (1991) Protein Expression and Purification 2,122-126.

[0022] Vitamin D is a group of fat-soluble secosteroids responsible for increasing intestinal absorption of calcium, iron, magnesium, phosphate, and zinc. Vitamin D is produced by the body in response to the skin being exposed to ultraviolet rays from sunlight. It is also found in naturally occurring foods such as fish, fish liver oils, egg yolks, and in fortified dairy and grain

products. In dietary supplements, the two most common compound forms of vitamin D are vitamin D3 (cholecalciferol) and vitamin D2 (ergocalciferol).

[0023] Vitamin D is a fat soluble vitamin that is biologically inert and must undergo two hydroxylations in the body for activation. The first occurs in the liver and converts vitamin D to 25-hydroxyvitamin D [25(OH)D], also known as calcifediol. The second occurs primarily in the kidney and forms the physiologically active 1,25-dihydroxyvitamin D [1,25(OH)₂D], also known as calcitriol. The active form of vitamin D, calcitriol, circulates as a hormone in the blood, regulating the concentration of calcium and phosphate in the bloodstream and promoting the healthy growth and remodeling of bone.

[0024] Vitamin D promotes calcium absorption and maintains adequate serum calcium and phosphate concentrations to enable normal mineralization of bone and to prevent hypocalcemic tetany. It is also used for bone growth and bone remodeling by osteoblasts and osteoclasts. It has also been shown to play a role in modulation of cell growth, neuromuscular and immune function, and reduction of inflammation.

[0025] Described herein is the administration of apoaeguorin and vitamin D-containing compositions to a subject in order to correct or maintain the calcium balance and vitamin D levels in that subject. Vitamin D deficiency may contribute to calcium imbalances. The maintenance of ionic calcium concentrations in plasma and body fluids is understood to be critical to a wide variety of bodily functions, including, but not limited to neuronal excitability, muscle contraction, membrane permeability, cell division, hormone secretion, bone mineralization, or the prevention of cell death following ischemia. Disruption in calcium homeostasis, i.e., a calcium imbalance, is understood to cause and/or correlate with many diseases, syndromes and conditions. Such diseases, syndromes and conditions include those associated with sleep quality, energy quality, mood quality, and memory quality and pain perception. The study of CaBPs has led to their recognition as protective factors acting in the maintenance of proper ionic calcium levels.

[0026] The maintenance of vitamin D levels is understood to be critical to calcium absorption, modulation of cell growth, neuromuscular and immune function, and reduction of inflammation. Vitamin D deficiency is most associated with rickets, a disease in which the bone tissues does not properly mineralize, leading to soft bones and skeletal deformities. Guidelines from the Institute of Medicine provides that the recommended dietary allowance (RDA) of vitamin D is 600 international units (IU) for adults ages 1-70, and 800 IU for adults older than age 70 to optimize bone health.

[0027] In certain embodiments, the methods described herein comprise administering apoaeguorin in combination with vitamin D for treating calcium imbalance, for delaying the progression of calcium imbalance, for preventing the onset of calcium imbalance, for preventing and/or treating the recurrence of calcium imbalance, and for treating vitamin D deficiency. Further described are methods which comprise administering apoaeguorin and vitamin D in combination with one or more additional agents having known therapeutic or

nutraceutical value. Particularly preferred applications of apoaquorin and vitamin D are in treating one or more symptoms and disorders related to quality of sleep, energy, mood, and memory and pain perception.

[0028] As used herein, the term "treating" includes preventative as well as disorder remittent treatment. As used herein, the terms "reducing", "alleviating", "suppressing" and "inhibiting" have their commonly understood meaning of lessening or decreasing. As used herein, the term "progression" means increasing in scope or severity, advancing, growing or becoming worse. As used herein, the term "recurrence" means the return of a disease after a remission.

[0029] As used herein, the term "administering" refers to bringing a patient, tissue, organ or cell in contact with apoaquorin and vitamin D. As used herein, administration can be accomplished *in vitro*, i.e., in a test tube, or *in vivo*, i.e., in cells or tissues of living organisms, for example, humans. In preferred embodiments, the present invention encompasses administering the compositions useful in the present invention to a patient or subject. A "patient" or "subject", used equivalently herein, refers to a mammal, preferably a human, that either: (1) has a calcium imbalance-related disorder and/or vitamin D deficiency remediable or treatable by administration of apoaquorin and vitamin D; or (2) is susceptible to a calcium imbalance-related disorder and/or vitamin D deficiency that is preventable by administering apoaquorin and vitamin D.

[0030] As used herein, the terms "effective amount" and "therapeutically effective amount" refer to the quantity of active agents sufficient to yield a desired therapeutic response without undue adverse side effects such as toxicity, irritation, or allergic response. The specific "effective amount" will, obviously, vary with such factors as the particular condition being treated, the physical condition of the patient, the type of animal being treated, the duration of the treatment, the nature of concurrent therapy (if any), and the specific formulations employed and the structure of the compounds or its derivatives. In this case, an amount would be deemed therapeutically effective if it resulted in one or more of the following: (1) the prevention of a calcium imbalance-related disorder and/or vitamin D deficiency; and (2) the reversal or stabilization of a calcium imbalance-related disorder and/or vitamin D deficiency. The optimum effective amounts can be readily determined by one of ordinary skill in the art using routine experimentation.

[0031] In certain compositions for oral administration to subjects, apoaquorin is formulated with at least one acceptable carrier at a dosage of approximately 10 to 50 mg/dose, a dose preferably in capsule form, with recommended dosage for a subject being approximately 20 mg/day. In certain preferred compositions for oral administration to subjects, vitamin D (in the form of D3 cholecalciferol) is formulated in combination with apoaquorin at a dosage of approximately 25-75 mcg/dose, with recommended dosage for a subject being approximately 50 mcg/day.

[0032] Compositions according to the present invention include liquids or lyophilized or otherwise dried formulations and include diluents of various buffer content (e.g., Tris-HCl,

acetate, phosphate), pH and ionic strength, additives such as albumin or gelatin to prevent absorption to surfaces, detergents (e.g., Tween 20, Tween 80, Pluronic F68, bile acid salts), solubilizing agents (e.g., glycerol, polyethylene glycerol), anti-oxidants (e.g., ascorbic acid, sodium metabisulfite), preservatives (e.g., Thimerosal, benzyl alcohol, parabens), bulking substances or tonicity modifiers (e.g., lactose, mannitol), covalent attachment of polymers such as polyethylene glycol to the protein, complexation with metal ions, or incorporation of the material into or onto particulate preparations of polymeric compounds such as polylactic acid, polyglycolic acid, or hydrogels, or onto liposomes, microemulsions, micelles, lamellar or multilamellar vesicles, erythrocyte ghosts or spheroplasts. Such compositions will influence the physical state, solubility, stability, rate of *in vivo* release, and rate of *in vivo* clearance. Controlled or sustained release compositions include formulation in lipophilic depots (e.g., fatty acids, waxes, oils).

[0033] Further described are methods of administering particulate compositions coated with polymers (e.g., poloxamers or poloxamines). Other embodiments of the compositions incorporate particulate forms protective coatings, protease inhibitors or permeation enhancers for various routes of administration, including parenteral, pulmonary, nasal and oral. In certain embodiments, the composition is administered parenterally, paracancerally, transmucosally, intramuscularly, intravenously, intradermally, subcutaneously, intraperitoneally, intraventricularly, intracranially or intratumorally.

[0034] Further, as used herein, "pharmaceutically acceptable carriers" are well known to those skilled in the art and include, but are not limited to, 0.0141M and preferably 0.05M phosphate buffer or 0.9% saline. Additionally, such pharmaceutically acceptable carriers may be aqueous or non-aqueous solutions, suspensions and emulsions. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oils such as olive oil and injectable organic esters such as ethyl oleate. Aqueous carriers include water, alcoholic/aqueous solutions, emulsions or suspensions, including saline and buffered media.

[0035] Parenteral vehicles include sodium chloride solution, Ringer's dextrose, dextrose and sodium chloride, lactated Ringer's and fixed oils. Intravenous vehicles include fluid and nutrient replenishers, electrolyte replenishers such as those based on Ringer's dextrose, and the like. Preservatives and other additives may also be present, such as, for example, antimicrobials, antioxidants, collating agents, inert gases and the like.

[0036] Controlled or sustained release compositions administrable according to the invention include formulation in lipophilic depots (e.g., fatty acids, waxes, oils). Also comprehended by the invention are particulate compositions coated with polymers (e.g., poloxamers or poloxamines) and the compound coupled to antibodies directed against tissue-specific receptors, ligands or antigens or coupled to ligands of tissue-specific receptors.

[0037] Other embodiments of the compositions administered according to the invention incorporate particulate forms, protective coatings, protease inhibitors or permeation enhancers for various routes of administration, including parenteral, pulmonary, nasal, ophthalmic and

oral.

[0038] Chemical entities modified by the covalent attachment of water-soluble polymers such as polyethylene glycol, copolymers of polyethylene glycol and polypropylene glycol, carboxymethyl cellulose, dextran, polyvinyl alcohol, polyvinylpyrrolidone or polyproline are known to exhibit substantially longer half-lives in blood following intravenous injection than do the corresponding unmodified compounds. Such modifications may also increase the chemical entities solubility in aqueous solution, eliminate aggregation, enhance the physical and chemical stability of the compound, and greatly reduce the immunogenicity and reactivity of the compound. As a result, the desired *in vivo* biological activity may be achieved by the administration of such polymer-entity adducts less frequently or in lower doses than with the unmodified entity.

[0039] In yet another method described herein, the composition can be delivered in a controlled release system. For example, the agent may be administered using intravenous infusion, an implantable osmotic pump, a transdermal patch, liposomes, or other modes of administration. In one embodiment, a pump may be used. In another embodiment, polymeric materials can be used. In yet another embodiment, a controlled release system can be placed in proximity to the therapeutic target, i.e., the brain, thus requiring only a fraction of the systemic dose.

[0040] The composition can comprise apoequorin and vitamin D, or can further include a pharmaceutically acceptable carrier, and can be in solid or liquid form such as tablets, powders, capsules, pellets, solutions, suspensions, elixirs, syrups, beverages, emulsions, gels, creams, ophthalmic formulations, or suppositories, including rectal and urethral suppositories. Pharmaceutically acceptable carriers also include gums, starches, sugars, cellulosic materials, and mixtures thereof. The composition containing apoequorin and vitamin D can be administered to a patient by, for example, subcutaneous implantation of a pellet. In a further embodiment, a pellet provides for controlled release of apoequorin and/or vitamin D over a period of time. The composition can also be administered by intravenous, intra-arterial, intramuscular injection of a liquid, oral administration of a liquid or solid, or by topical application. Administration can also be accomplished by use of a rectal suppository or a urethral suppository.

[0041] The compositions described herein can be prepared by known dissolving, mixing, granulating, or tablet-forming processes. For oral administration, apoequorin or its physiologically-tolerated derivatives such as salts, esters, N-oxides, and the like and/or vitamin D or its physiologically-tolerated derivatives such as salts, esters, N-oxides, and the like are mixed with additives customary for this purpose, such as vehicles, stabilizers, or inert diluents, and converted by customary methods into suitable forms for administration, such as tablets, coated tablets, hard or soft gelatin capsules, aqueous, alcoholic or oily solutions.

[0042] Examples of suitable inert vehicles are conventional tablet bases such as lactose, sucrose, or cornstarch in combination with binders such as acacia, cornstarch, gelatin, with

disintegrating agents such as cornstarch, potato starch, alginic acid, or with a lubricant such as stearic acid or magnesium stearate.

[0043] Examples of suitable oily vehicles or solvents are vegetable or animal oils such as sunflower oil or fish-liver oil. Compositions can be effected both as dry and as wet granules. For parenteral administration (subcutaneous, intravenous, intraarterial, or intramuscular injection), the chemical entity or its physiologically tolerated derivatives such as salts, esters, N-oxides, and the like are converted into a solution, suspension, or expulsion, if desired with the substances customary and suitable for this purpose, for example, solubilizers or other auxiliaries.

[0044] Examples are sterile liquids such as water and oils, with or without the addition of a surfactant and other pharmaceutically acceptable adjuvants. Illustrative oils are those of petroleum, animal, vegetable, or synthetic origin, for example, peanut oil, soybean oil or mineral oil. In general, water, saline, aqueous dextrose and related sugar solutions, and glycols such as propylene glycols or polyethylene glycol are preferred liquid carriers, particularly for injectable solutions.

[0045] The preparation of compositions which contain an active component is well understood in the art. Such compositions may be prepared as aerosols delivered to the nasopharynx or as injectables, either as a liquid solutions or suspensions; however, solid forms suitable for solution in, or suspension in, liquid prior to injection can also be prepared. The composition can also be emulsified. The active therapeutic ingredient is often mixed with excipients which are pharmaceutically acceptable and compatible with the active ingredient. Suitable excipients include, for example, water, saline, dextrose, glycerol, ethanol, or the like or any combination thereof. In addition, the composition can contain minor amounts of auxiliary substances such as wetting or emulsifying agents, pH buffering agents which enhance the effectiveness of the active ingredient.

[0046] An active component can be formulated into the composition as neutralized pharmaceutically acceptable salt forms. Pharmaceutically acceptable salts include the acid addition salts, which are formed with inorganic acids such as, for example, hydrochloric, or phosphoric acids, or such organic acids as acetic, tartaric, mandelic, and the like. Salts formed from the free carboxyl groups can also be derived from inorganic bases such as, for example, sodium, potassium, ammonium, calcium, or ferric hydroxides, and such organic bases as isopropylamine, trimethylamine, 2-ethylamino ethanol, histidine, procaine, and the like.

[0047] For topical administration to body surfaces using, for example, creams, gels, drops, and the like apoequorin or its physiologically-tolerated derivatives and/or vitamin D or its physiologically-tolerated derivatives are prepared and applied as solutions, suspensions, or emulsions in a physiologically acceptable diluent with or without a pharmaceutical carrier.

[0048] In another method described herein, the active component can be delivered in a vesicle, in particular, a liposome (see Langer, Science 249:1527-1533 (1990); Treat et al.,

Liposomes in the Therapy of Infectious Disease and Cancer, Lopez-Berestein and Fidler (eds.), Liss, N.Y., pp.353-365 (1989).

[0049] Salts of apoeaquerin and/or vitamin D are preferably pharmaceutically acceptable salts. Other salts may, however, be useful in the preparation of the compositions according to the invention or of their pharmaceutically acceptable salts. Suitable pharmaceutically acceptable salts include acid addition salts which may, for example, be formed by mixing a solution of apoeaquerin and/or vitamin D with a solution of a pharmaceutically acceptable acid such as hydrochloric acid, sulphuric acid, methanesulphonic acid, fumaric acid, maleic acid, succinic acid, acetic acid, benzoic acid, oxalic acid, citric acid, tartaric acid, carbonic acid or phosphoric acid.

[0050] In addition, apoeaquerin and vitamin D-containing compositions described herein may be provided in the form of nutraceutical compositions where apoeaquerin and vitamin D prevent the onset of or reduce or stabilize various deleterious calcium imbalance-related disorders and vitamin D deficiency. The term "nutraceutical" or "nutraceutical composition", for the purpose of this specification, refers to a food item, or a part of a food item, that offers medical health benefits, including prevention and/or treatment of disease. A nutraceutical composition according to the present invention may contain only apoeaquerin and vitamin D as active ingredients, or alternatively, may further comprise, in admixture with dietary supplements including vitamins, co-enzymes, minerals, herbs, amino acids and the like which supplement the diet by increasing the total intake of that substance.

[0051] Therefore, described herein are methods of providing nutraceutical benefits to a patient comprising the step of administering to the patient a nutraceutical composition containing apoeaquerin and vitamin D. Such compositions generally include a "nutraceutically-acceptable carrier" which, as referred to herein, is any carrier suitable for oral delivery including aforementioned pharmaceutically-acceptable carriers suitable for the oral route. In certain embodiments, nutraceutical compositions according to the invention comprise dietary supplements which, defined on a functional basis, include immune boosting agents, anti-inflammatory agents, anti-oxidant agents, anti-viral agents, or mixtures thereof.

[0052] Immune boosters and/or anti-viral agents are useful for accelerating wound-healing and improved immune function; and they include extracts from the coneflowers, or herbs of the genus *Echinacea*, extracts from herbs of the genus *Sambuca*, and Goldenseal extracts. Herbs of the genus *Astragalus* are also effective immune boosters in either their natural or processed forms. *Astragalus* stimulates development of stem cells in the marrow and lymph tissue active immune cells. Zinc and its bioactive salts, such as zinc gluconate and zinc acetate, also act as immune boosters in the treatment of the common cold.

[0053] Antioxidants include the natural, sulfur-containing amino acid allicin, which acts to increase the level of antioxidant enzymes in the blood. Herbs or herbal extracts, such as garlic, which contain allicin, are also effective antioxidants. The catechins, and the extracts of herbs such as green tea containing catechins, are also effective antioxidants. Extracts of the genus

Astragalus also show antioxidant activity. The bioflavonoids, such as quercetin, hesperidin, rutin, and mixtures thereof, are also effective as antioxidants. The primary beneficial role of the bioflavonoids may be in protecting vitamin C from oxidation in the body. This makes more vitamin C, or ascorbic acid, available for use by the body.

[0054] Bioflavonoids such as quercetin are also effective anti-inflammatory agents, and may be used as such in the inventive compositions. Anti-inflammatory herbal supplements and anti-inflammatory compounds derived from plants or herbs may also be used as anti-inflammatory agents in the inventive composition. These include bromelain, a proteolytic enzyme found in pineapple; teas and extracts of stinging nettle; turmeric, extracts of turmeric, or curcumin, a yellow pigment isolated from turmeric.

[0055] Another supplement which may be used in the present invention is ginger, derived from herbs of the genus *Zingiber*. This has been found to possess cardiogenic activity due to compounds such as gingerol and the related compound shogaol as well as providing benefits in the treatment of dizziness, and vestibular disorders. Ginger is also effective in the treatment of nausea and other stomach disorders.

[0056] Supplements which assist in rebuilding soft tissue structures, particularly in rebuilding cartilage, are useful in compositions for treating the pain of arthritis and other joint disorders. Glucosamine, glucosamine sulfate, chondroitin may be derived from a variety of sources such as Elk Velvet Antler. Marine lipid complexes, omega 3 fatty acid complexes, and fish oil are also known to be useful in treating pain associated with arthritis.

[0057] Supplements useful in treating migraine headaches include feverfew and *Ginkgo biloba*. The main active ingredient in feverfew is the sesquiterpene lactone parthenolide, which inhibits the secretions of prostaglandins which in turn cause pain through vasospastic activity in the blood vessels. Feverfew also exhibits anti-inflammatory properties. Fish oil, owing to its platelet-stabilizing and antivasospastic actions, may also be useful in treating migraine headaches. The herb *Ginkgo biloba* also assists in treatment of migraines by stabilizing arteries and improving blood circulation.

[0058] Although some of the supplements listed above have been described as to their pharmacological effects, other supplements may also be utilized in the present invention and their effects are well documented in the scientific literature.

[0059] The invention will be more fully understood upon consideration of the following nonlimiting Examples describing and disclosing the chemicals, instruments, statistical analysis and methodologies which are reported in the publications which might be used in connection with the invention.

EXAMPLES:

Example 1. Administration of apoeaquorin over a ninety (90) day time course results in improved quality of life for test subjects.

[0060] The present analysis, an open-label study, of 32 patients over a 90 day period shows an increase in overall quality of sleep, energy, mood, pain, general health. Changes in performance were measured via a standardized battery of questions. These included assessments of qualitative cognitive test, a sleep index, a headache index and a Quality of Life questionnaire. The study shows improved performance. No participants discontinued the study due to an adverse event.

[0061] The results illustrated in Fig. 1 show the percent change from baseline of scores from the areas mentioned; we have excluded the memory scores for another graph. The analysis here is shown as marked on the graph as 1, 2, 3, 4 and 5 vs. days 0 through 90. The graph shows an increase in overall quality of sleep, energy, mood, pain and general health. The baseline was known from a pre-study phase.

Example 2. Administration of apoeaquorin over a thirty (30) day time course results in improved quality of life for test subjects.

[0062] The present study was an open-label study for 56 participants over a 30 day period. Changes in performance were measured via a memory screening tool. As illustrated in Fig. 2, the study showed improved memory performance as early as eight days but with statistically greater improvement at day 30. No participants discontinued the study due to an adverse event.

Example 3. Administration of apoeaquorin over a ninety (90) day time course results in improved cognition for test subjects.

[0063] The present analysis, for an open-label study of 32 patients shows an increase in cognitive ability. Changes in performance were measured via a standardized cognitive battery. The study showed improved cognition as early as eight days but with statistically greater improvement at day 30, as well as 60-90. No participants discontinued the study due to an adverse event. The results shown in Fig. 3 demonstrate the significant percent increase from baseline of scores in cognitive ability. Note: Greater than 51% of participants had an increase in cognitive ability.

Example 4. Administration of apoeaquorin and vitamin D-containing composition.

[0064] An apoeaquorin and vitamin D-containing composition is administered to a patient in capsule form containing an admixture of apoeaquorin and vitamin D3. The composition contains 50 mcg of vitamin D3 (in the form of D3 cholecalciferol) and 20 mg of apoeaquorin. The nutraceutical composition is carried in a nutraceutically-acceptable vegetable capsule (vegetable cellulose, water). The composition further contains microcrystalline cellulose, sugar, and small amounts of acacia (gum Arabic), casein peptones, corn starch, lactose, magnesium stearate (vegetable source), medium chain triglycerides (vegetable oil), salt, soy peptones, DL- α -tacopheral, tricalcium phosphate, and water. One capsule taken daily provides a recommended dosage. Such dosage contains approximately 250% of the recommended daily allowance of vitamin D. Vitamin D suitable for use in the invention is available from BASF and sold as "Dry Vitamin D3 100 GFP/HP". The single dose may alternatively be formulated to contain other amounts of apoeaquorin, including single dosages containing 10 mg or 40 mg of apoeaquorin.

Example 5. Evaluation of neuroprotection by AQ (apoeaquorin) and Vitamin D in rat hippocampal brain slices.

[0065] Preliminary results from a subset of experimental data indicate significant cell death following 5-min of oxygen glucose deprivation (OGD Fig. 4; upper left image) and minimal cell death in slices that were not subjected to 5-min OGD (Fig. 4; upper right image). Direct hippocampal injection of AQ results in fewer dead and dying cells (i.e., blue stained cells). Rats fed a diet supplemented with vitamin D (0.0125 mg/kg) for ~10 days also appear to have fewer dead or dying cells. The combination of AQ and vitamin D may further reduce the amount of cell death (Fig. 4; lower left image). Control slices (those not subjected to OGD) do not appear to vary appreciably as a function of treatment condition.

REFERENCES CITED IN THE DESCRIPTION

Cited references

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Patentkrav

1. Sammensætning til anvendelse ved behandling af et symptom eller en lidelse associeret med mindst én af calciumubalance og D-vitaminmangel, hvor

5 symptomet eller lidelsen er relateret til celledød efter iskæmi, omfattende: (a) en effektiv mængde af apoaeguorin; (b) en effektiv mængde af D-vitamin; og (c) en farmaceutisk acceptabel bærer, hvor: (a) den terapeutisk effektive mængde af apoaeguorin i enhedsdosen er ca. 10 mg til ca. 50 mg; og (b) den terapeutisk effektive mængde af D-vitamin i enhedsdosen er ca. 25 mcg til ca. 75 mcg.

10
2. Sammensætningen ifølge krav 1, hvor sammensætningen er i form af en enhedsdosis indeholdende de effektive mængder af apoaeguorin og D-vitamin.
3. Sammensætningen ifølge krav 1, hvor: (a) den effektive mængde af

15 apoaeguorin i enhedsdosen er ca. 20 mg; og (b) den effektive mængde af D-vitamin i enhedsdosen er ca. 50 mcg.
4. Sammensætningen ifølge et hvilket som helst af kravene 1-3, hvor D-vitaminet er i form af D3 cholecalciferol.

20
5. Sammensætningen ifølge et hvilket som helst af kravene 1-4, hvor enhedsdosen er en kapsel.
6. Sammensætningen ifølge et hvilket som helst af kravene 1-5, hvor

25 sammensætningen er en nutraceutisk sammensætning.

DRAWINGS

Drawing

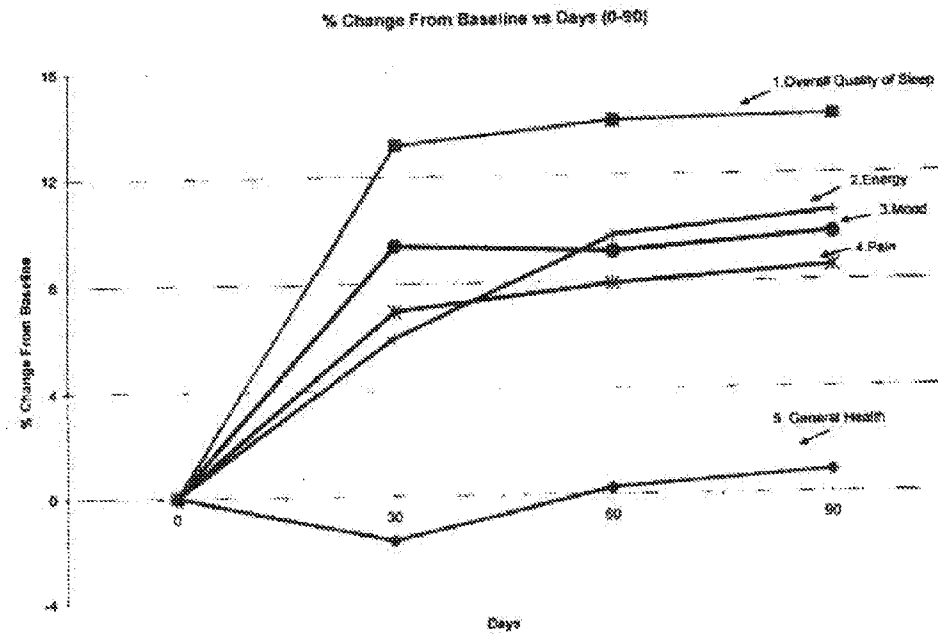


Fig. 1

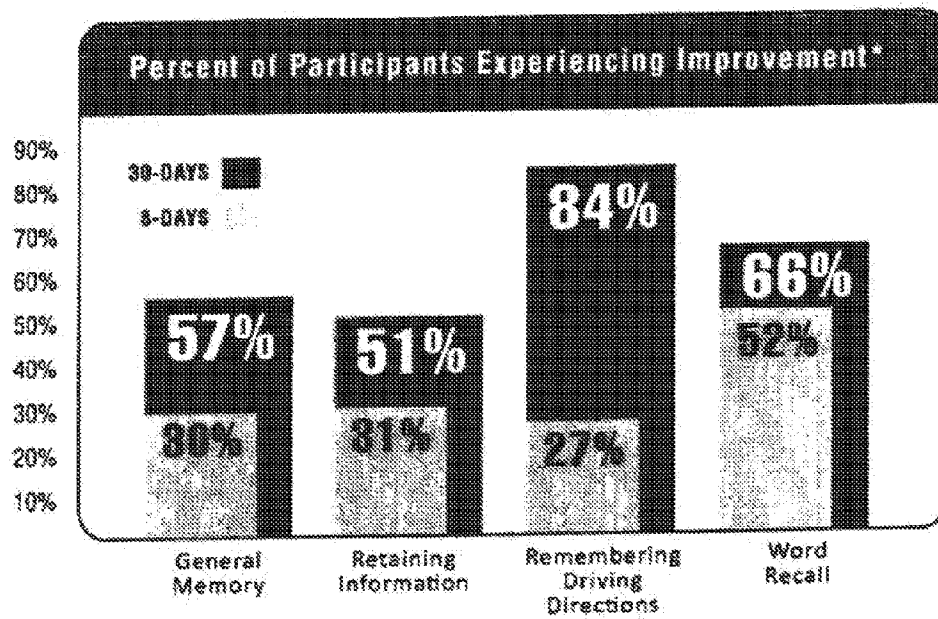


Fig. 2

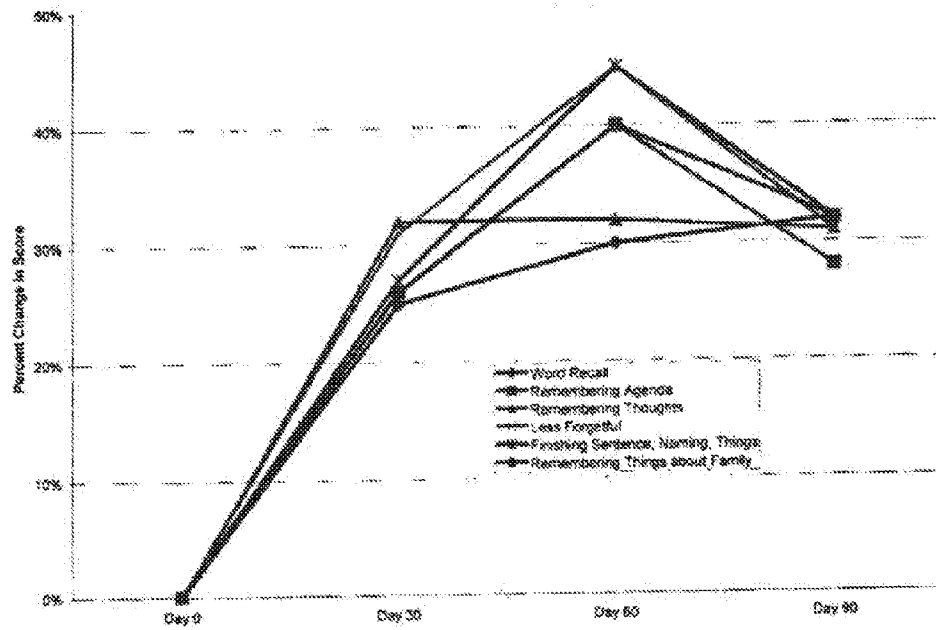
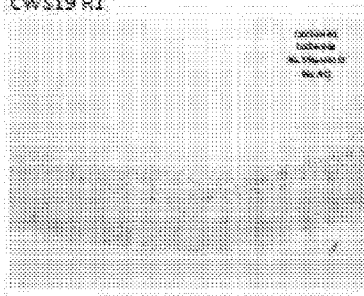
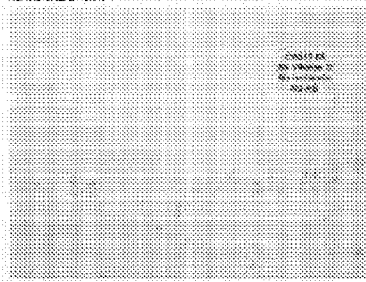


Fig. 3

Ischemia/No AQ/No VitD
CWS19 R1



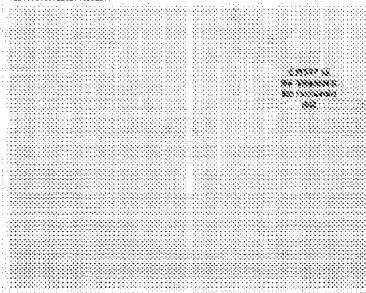
No Ischemia/No AQ/No VitD
CWS19 R4



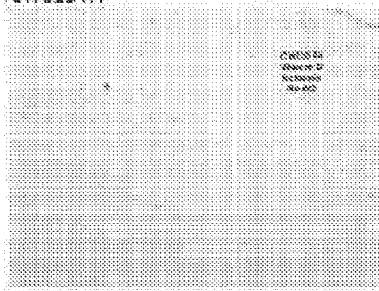
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CWS17 L4



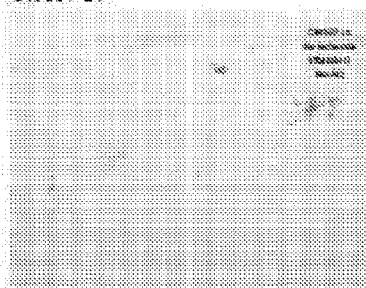
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CWS19 L2



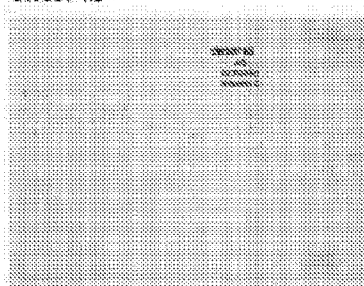
Ischemia/No AQ/VitD
CWS10 R4



No Ischemia/No AQ/VitD
CWS07 L4



Ischemia/AQ/VitD
CWS07 R5



No Ischemia/AQ/VitD
CWS07 R4

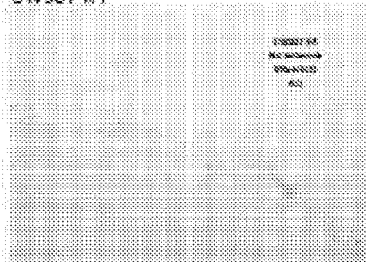


Fig. 4