DIPTERINYL CALCIUM PENTAHYDRATE (DCP) AND THERAPEUTIC METHODS BASED THEREON

Inventors: Phillip Moheno, La Jolla, CA (US); Wolfgang Pfeiderer, Konstanz (DE)

Correspondence Address:
WILSON, SONSINI, GOODRICH & ROSATI
650 PAGE MILL ROAD
PALO ALTO, CA 94304-1050 (US)

Appl. No.: 12/643,752
Filed: Dec. 21, 2009

Related U.S. Application Data
Division of application No. 11/981,398, filed on Oct. 30, 2007, now Pat. No. 7,662,820.

Provided herein is dipterinyl calcium pentahydrate (DCP) and therapeutic methods based thereon. Also provided herein is the compound dipterinyl calcium pentahydrate (DCP) or an analog or polymorph thereof.
Figure 2. Structure of DCP as determined by single crystal x-ray diffraction
Figure 3. Nude mice treated with DCP showed no significant weight loss
Figure 4: Determination of optimum dose in nude mice
Day 46 Tumor Growth of MDA-MB-231 Human Breast Cancer Xenographs in Nude Mice

Figure 5: Comparison of DCP to other forms of calcium pterin after 46 days of treatment
Figure 6: Comparison of DCP to other forms of calcium pterin after 57 days of treatment
Figure 7: Treatment/Control (T/C) values in nude mice w/ MDA-MB-231 after 11 days of treatment
Figure 8: Treatment/Control (T/C) values in nude mice w/ MDA-MB-231 after 36 days of treatment
Figure 9: Treatment/Control (T/C) values in nude mice w/ MDA-MB-231 after 47 days of treatment
DIPTERINYL CALCIUM PENTAHYDRATE (DCP) AND THERAPEUTIC METHODS BASED THEREON

RELATED APPLICATIONS

[0001] This application claims priority to U.S. Provisional Application No. 60/863,547, filed on Oct. 30, 2006, which is incorporated by reference in its entirety.

SUMMARY OF THE INVENTION

[0002] One embodiment provides dipterinyl calcium pentahydrate (DCP) and therapeutic methods based thereon. Another embodiment provides the compound dipterinyl calcium pentahydrate (DCP) or an analog or polymorph thereof. Another embodiment provides a method of synthesizing dipterinyl calcium pentahydrate (DCP) comprising: dissolving pterin in an aqueous solution of NaOH, adding CaCl₂·2H₂O to the solution with stirring at a pH of about 11, continuing stirring for about 1 day, and collecting the precipitate as DCP. One embodiment provides a method of modulating tryptophan production comprising administering to a subject an effective amount of DCP. Another embodiment provides a method of modulating tryptophan production comprising administering to a subject an effective amount of DCP suspension. Additional embodiments provide a method of modulating tryptophan degradation comprising administering to a subject an effective amount of DCP. Another embodiment provides method of modulating tryptophan degradation comprising administering to a subject an effective amount of DCP suspension. Another embodiment provides a method of modulating neopterin production comprising administering to a subject an effective amount of DCP. An additional embodiment provides a method of modulating neopterin production comprising administering to a subject an effective amount of DCP suspension.

[0003] Yet another embodiment provides a method of modulating IFN-γ production comprising administering to a subject an effective amount of DCP. An additional embodiment provides a method of modulating IFN-γ production comprising administering to a subject an effective amount of DCP suspension. One embodiment provides a method of modulating the activity of the enzyme IDO comprising administering to a subject an effective amount of DCP. Another embodiment provides a method of modulating the activity of the enzyme IDO comprising administering to a subject an effective amount of DCP suspension. One embodiment provides a method of modulating oxidants production comprising administering to a subject an effective amount of DCP. Another embodiment provides a method of modulating oxidants production comprising administering to a subject an effective amount of DCP suspension. Yet another embodiment provides a method of modulating free radical production comprising administering to a subject an effective amount of DCP. An additional embodiment provides a method of modulating free radical production comprising administering to a subject an effective amount of DCP suspension.

[0004] One embodiment provides a method of modulating release of reactive oxygen species (ROS) comprising administering to a subject an effective amount of DCP. Another embodiment provides a method of modulating release of reactive oxygen species (ROS) comprising administering to a subject an effective amount of DCP suspension. An additional embodiment provides a method of modulating release of reactive oxygen species (ROS) comprising administering to a subject an effective amount of DCP. Another embodiment provides a method of modulating release of reactive oxygen species (ROS) comprising administering to a subject an effective amount of DCP suspension. An additional embodiment provides a method of modulating the expression of downstream genes for a cytokine, a chemokine, an adhesion molecule, growth factor, enzyme and/or immune receptor comprising administering to a subject an effective amount of DCP. An additional embodiment provides a method of modulating the expression of downstream genes for a cytokine, a chemokine, an adhesion molecule, growth factor, enzyme and/or immune receptor comprising administering to a subject an effective amount of DCP suspension.

[0005] One embodiment provides a method of reducing or suppressing allogeneic immune tolerance comprising administering to a subject an effective amount of DCP. Another embodiment provides a method of reducing or suppressing allogeneic immune tolerance comprising administering to a subject an effective amount of DCP suspension. Another embodiment provides a method of modulating immune resistance in human solid tumors comprising administering to a subject an effective amount of DCP. Another embodiment provides a method of modulating immune resistance in human solid tumors comprising administering to a subject an effective amount of DCP suspension. Another embodiment provides a method of modulating antiproliferative activity comprising administering to a subject an effective amount of DCP. Another embodiment provides a method of modulating antiproliferative activity comprising administering to a subject an effective amount of DCP suspension.

[0006] One embodiment provides a compound of formula (I):

M(pterin)(H₂O)y

[0007] wherein:

[0008] M is a bivalent metal ion selected from the group consisting of Ca²⁺, Cu²⁺, Mg²⁺, Zn²⁺, Cr³⁺, Mn²⁺, Fe²⁺, Mo(VI), Zn²⁺, Sr²⁺, Ba²⁺, Ra²⁺, Ru⁴⁺, Rh³⁺, Pd²⁺, Cd²⁺, Sn⁴⁺, W⁶⁺, Re⁷⁺, Os⁷⁺, Ir⁷⁺, Pt⁷⁺, Si⁷⁺; and Sm²⁺;

[0009] y is an integer from 1 to 8;

[0010] y is an integer from 1 to 8.

[0011] Another embodiment provides a method of inhibiting tumor cells in an animal comprising the administration of a therapeutically effective amount of a compound of formula (I):

M(pterin)(H₂O)y

[0012] for treating an illness or disease.
[0012] wherein:

$$M(n	ext{termin}_i)\cdot H_2 O$$

[0013] M is a bivalent metal ion selected from the group consisting of Ca$^{2+}$, Cu$^{2+}$, Mg$^{2+}$, V$^{2+}$, Cr$^{2+}$, Mn$^{2+}$, Fe$^{2+}$, Mo$^{2+}$, Zn$^{2+}$, Sr$^{2+}$, Ba$^{2+}$, Ra$^{2+}$, Ru$^{2+}$, Rh$^{2+}$, Pd$^{2+}$, Cd$^{2+}$, Sn$^{2+}$, W$^{2+}$, Re$^{2+}$, Os$^{2+}$, Ir$^{2+}$, Pt$^{2+}$, Si$^{2+}$, and Sm$^{2+}$;

[0014] X is an integer from 1 to 8; and

[0015] y is an integer of from 1 to 8.

[0016] One embodiment provides a method of treating a viral infection comprising the administration of a therapeutically effective amount of a compound of formula (I):

$$M(n	ext{termin}_i)\cdot H_2 O$$

wherein:

[0017] M is a bivalent metal ion selected from the group consisting of Ca$^{2+}$, Cu$^{2+}$, Mg$^{2+}$, V$^{2+}$, Cr$^{2+}$, Mn$^{2+}$, Fe$^{2+}$, Mo$^{2+}$, Zn$^{2+}$, Sr$^{2+}$, Ba$^{2+}$, Ra$^{2+}$, Ru$^{2+}$, Rh$^{2+}$, Pd$^{2+}$, Cd$^{2+}$, Sn$^{2+}$, W$^{2+}$, Re$^{2+}$, Os$^{2+}$, Ir$^{2+}$, Pt$^{2+}$, Si$^{2+}$, and Sm$^{2+}$;

[0018] X is an integer from 1 to 8; and

[0019] y is an integer of from 1 to 8.

[0020] Another embodiment provides a pharmaceutical composition comprising a pharmaceutically acceptable excipient and a compound of formula (I):

$$M(n	ext{termin}_i)\cdot H_2 O$$

wherein:

[0021] M is a bivalent metal ion selected from the group consisting of Ca$^{2+}$, Cu$^{2+}$, Mg$^{2+}$, V$^{2+}$, Cr$^{2+}$, Mn$^{2+}$, Fe$^{2+}$, Mo$^{2+}$, Zn$^{2+}$, Sr$^{2+}$, Ba$^{2+}$, Ra$^{2+}$, Ru$^{2+}$, Rh$^{2+}$, Pd$^{2+}$, Cd$^{2+}$, Sn$^{2+}$, W$^{2+}$, Re$^{2+}$, Os$^{2+}$, Ir$^{2+}$, Pt$^{2+}$, Si$^{2+}$, and Sm$^{2+}$;

[0022] X is an integer from 1 to 8; and

[0023] y is an integer of from 1 to 8.

Incorporation by Reference

[0025] All publications and patent applications mentioned in this specification are herein incorporated by reference to the same extent as if each individual publication or patent application was specifically and individually indicated to be incorporated by reference.

BRIEF DESCRIPTION OF THE DRAWINGS

[0026] FIGS. 1A and 1B (with SEMs): Twenty-nine athymic nude (nu/nu) female mice, ages 3-4 weeks, were inoculated with 10×106 MDA-MB-231 cancer cells subcutaneously into the right flank of each mouse. When tumors reached 3-5 mm in size, twenty-five of the mice were divided into five treatment groups of five mice each. Four mice were assigned as controls. Four mice with outlying tumor sizes or non-tumor takes were subsequently excluded. Experimental groups were treated by oral gavage once daily with the indicated test suspensions or solution. The control group was untreated.

[0027] FIG. 2: Structure of DCP as determined by single crystal X-ray diffraction.

[0028] FIG. 3: Nude mice treated with DCP showed no significant weight loss.

[0029] FIG. 4: Determination of optimum dose in nude mice.

[0030] FIG. 5: Comparison of DCP to other forms of calcium pterin after 46 days of treatment.

[0031] FIG. 6: Comparison of DCP to other forms of calcium pterin after 57 days of treatment.

[0032] FIG. 7: Treatment/Control (T/C) values in nude mice w/MDA-MB-231 after 11 days of treatment.

[0033] FIG. 8: Treatment/Control (T/C) values in nude mice w/MDA-MB-231 after 36 days of treatment.

[0034] FIG. 9: Treatment/Control (T/C) values in nude mice w/MDA-MB-231 after 47 days of treatment.

DETAILED DESCRIPTION OF THE INVENTION

Glossary

[0035] To more readily facilitate an understanding of the invention and its embodiments, the meanings of terms used herein will become apparent from the context of this specification in view of common usage of various terms and the explicit definitions of other terms provided in the glossary below or in the ensuing description.

[0036] As used herein, the terms “comprising,” “including,” and “such as” are used in their open, non-limiting sense.

[0037] The use of the term “about” in the present disclosure means “approximately,” and illustratively, the use of the term “about” indicates that values slightly outside the cited values may also be effective and safe, and such dosages are also encompassed by the scope of the present claims.

[0038] “Binders” impart cohesive qualities and include, e.g., algic acid and salts thereof; cellulose derivatives such as carboxymethylcellulose, methylcellulose (e.g., Methocel®), hydroxypropylcellulose, hydroxyethylcellulose, hydroxypropylcellulose (e.g., Klucel®), ethylcellulose (e.g., Ethocel®), and microcrystalline cellulose (e.g., Avicel®); microcrystalline dextrose; amylose; magnesium aluminum silicate; polysaccharide acids; bentonites; gelatin, polyvinylpyrrolidone/vinyl acetate copolymer; crosspolydione; povidone; starch; pregelatinized starch; tragacanth; dextrin, a sugar, such as sucrose (e.g., Dipsac®, glucose, dextrose, maltose, manniot, sorbitol, xylitol (e.g., Xylitab®), and lactose; a natural or synthetic gum such as acacia, tragacanth, ghatti gum, mucilage of isapol husks, polyvinylpyrrolidone (e.g., Polyvidone® CL, Kollidon® CL, Polysplasdone® XI-10), larch araboalactan, Veegum®, polyethylene glycol, waxes, sodium alginate, and the like.


[0040] The term “controlled release” includes any non-immediate release formulation, including but not limited to enteric-coated formulations and sustained release, delayed-release and pulsatile release formulations.
The term “delayed-release” includes any non-immediate release formulation, including but not limited to, film-coated formulations, enteric-coated formulations, encapsulated formulations, sustained release formulations and pulsatile release formulations.

“Diffusion facilitators” and “dispersing agents” include materials that control the diffusion of an aqueous fluid through a coating. Exemplary diffusion facilitators/dispersing agents include, e.g., hydrophilic polymers, electrolytes, Tween® 60 or 80, PEG and the like. Combinations of one or more erosion facilitator with one or more diffusion facilitator can also be used in the present invention.

“Diluents” increase bulk of the composition to facilitate compaction. Such compounds include, e.g., lactose; starch; mannitol; sorbitol; dextrose; microcrystalline cellulose such as Avicel®; dibasic calcium phosphate; dicalcium phosphate dihydrate; tricalcium phosphate; calcium phosphate; anhydrous lactose; spray-dried lactose; pregelatinized starch; compressible sugar, such as Di-Pac® (Amstar); mannitol; hydroxypropylmethylcellulose; sucrose-based diluents; confectioner’s sugar; monobasic calcium sulfate monohydrate; calcium sulfate dihydrate; calcium lactate tribhydrate; dextrates; hydrolyzed cereal solids; amylose; powdered cellulose; calcium carbonate; glycine; kaolin; mannitol; sodium chloride; inositol; bentonite; and the like.

“Filling agents” include compounds such as lactose, calcium carbonate, calcium phosphate, dibasic calcium phosphate, calcium sulfate, microcrystalline cellulose, cellulose powder, dextrose; dextrates; dextran, starches, pregelatinized starch, sucrose, xylitol, lactitol, mannitol, sorbitol, sodium chloride, polyethylene glycol, and the like. The terms “therapeutically effective amount” and “effective amount” in relation to the amount of active pharmaceutical ingredient mean, consistent with considerations known in the art, the amount of active pharmaceutical ingredient effective to elicit a pharmacologic effect or therapeutic effect without undue adverse side effects.

An “enteric-coating” is a substance that remains substantially intact in the stomach but dissolves and releases at least some of the drug once reaching the small intestine. Generally, the enteric-coating comprises a polymeric material that prevents release in the low pH environment of the stomach but that ionizes at a slightly higher pH, typically a pH of 4 or 5, and thus dissolves sufficiently in the small intestines to gradually release the active agent therein.

The term “immediate release” is intended to refer to any formulation in which all or part of the active pharmaceutical ingredient is in solution either before administration or immediately (i.e., within about 30 minutes) after administration. For example, with an “immediate release” formulation, oral administration results in immediate release of the agent from the composition into gastric fluid. For delayed-release formulations, the opposite is generally true, the rate of release of drug from the dosage form is the rate-limiting step in the delivery of the drug to the target area.

“Lubricants” are compounds which prevent, reduce or inhibit adhesion or friction of materials. Exemplary lubricants include, e.g., stearic acid; calcium hydroxide; talc; sodium stearyl fumarate; a hydrocarbon such as mineral oil, or hydrogenated vegetable oil such as hydrogenated soybean oil (Sterolins®); higher fatty acids and their alkali-metal and alkaline earth metal salts, such as aluminum, calcium, magnesium, zinc, stearic acid, sodium stearates, glycerol, talc; waxes, Stearowet®, boric acid, sodium benzoate, sodium acetate, sodium chloride, leucine, a polyethylene glycol or a methoxypolyethylene glycol such as Carbopol™, sodium oleate, glycercyl behenate, polyethylene glycol, magnesium or sodium laurel sulfate, colloidal silica such as Syloid™, Carb-O-Sil®, a starch such as corn starch, silicone oil, a surfactant, and the like.

The term “pharmaceutically acceptable” is used adjectively herein to mean that the modified noun is appropriate for use in a pharmaceutical product.

“Solubilizers” include compounds such as citric acid, succinic acid, fumaric acid, malic acid, tartaric acid, maleic acid, glu taric acid, sodium bicarbonate, sodium carbonate and the like.

“Stabilizers” include compounds such as any anti-oxidation agents, buffers, acids, and the like.

“Suspended agents” or “thickening agents” include compounds such as polyvinylpyrrolidone, e.g., polyvinylpyrrolidone K12, polyvinylpyrrolidone K17, polyvinylpyrrolidone K25, or polyvinylpyrrolidone K30; polyethylene glycol, e.g., the polyethylene glycol can have a molecular weight of about 300 to about 6000, or about 3350 to about 4000, or about 700 to about 5400; sodium carboxymethylcellulose; methylcellulose; hydroxypropylmethylcellulose; polysorbate-80; hydroxyethylcellulose; sodium alginate; gums, such as, e.g., gum tragacanth and gum acacia; guar gum; xanthans, including xanthan gum; sugars; celluloses, such as, e.g., sodium carboxymethylcellulose, methylcellulose, sodium carboxymethylcellulose, hydroxypropylmethylcellulose, hydroxyethylcellulose; polysorbate-80; sodium alginate; polyethoxylated sorbitan monolaureate; polyethoxylated sorbitan monolaurate; povidone and the like.

“Surfactants” include compounds such as sodium laurel sulfate, sorbitan monooctenyl; polyoxyethylene sorbitan monoleate, polysorbates, poloxamers, bile salts, glyceryl monostearate, copolymers of ethylene oxide and propylene oxide, e.g., Pluronic® (BASF); and the like.

As used herein, the terms “suspension” and “solution” are interchangeable with each other and generally mean a solution and/or suspension of the substituted benzimidazole in an aqueous medium.

The term “sustained release” is used in its conventional sense to refer to a drug formulation that provides for gradual release of a drug over an extended period of time, and, may sometimes, although not necessarily, result in substantially constant blood levels of a drug over an extended time period.

The term “treat” or “treatment” as used herein refers to any treatment of a disorder or disease associated with gastrointestinal disorder, and includes, but is not limited to, preventing the disorder or disease from occurring in a mammal which may be predisposed to the disorder or disease, but has not yet been diagnosed as having the disorder or disease; inhibiting the disorder or disease, for example, arresting the development of the disorder or disease; relieving the disorder or disease, for example, causing regression of the disorder or disease; or relieving the condition caused by the disease or disorder, for example, stopping the symptoms of the disease or disorder.

Perin has been a point of interest in the biomedical research community for some time. Xanthopterin was found to inhibit sarcoma growth in mice over 60 years ago (Lewisohn et al, Proc. Soc. Exp. Biol. Med. (1944) 56, 144-145). Isoxanthopterin was also shown to inhibit tumor growth (Kokolis et al, Z. Naturforsch. (1972) B27, 292-295). The

Further investigations by Moheno demonstrated the importance of selecting an immunocompetent mouse strain for the evaluation of anti-tumor efficacy of pterin and related analogs (Moheno, Int. J. Pharm. (2004), 271, 293-300). In this study, a suspension of calcium pterin (known as CaPterin) was found to possess significant antitumor efficacy against MDA-MB-231 human breast xenografts in nude mice, as well as highly significant activity against spontaneous mammary gland tumors in C3H/HeN-MTV+ mice, based upon National Cancer Institute standards. Immunomodulatory action for CaPterin was deduced by comparing the antitumor efficacy of CaPterin in four different mouse/tumor systems: i.e., the two cited above, as well as Balb/c mice with EMT6 xenografts and SCID mice with MDA-MB-231 xenografts. Comparison of results obtained by testing CaPterin in either nude or SCID mice (severely compromised immunodeficient) implanted with MDA-MB-231 human cancer cells showed a significant antitumor response in the nude and no response in the SCIDs. This comparison argues for B-cell immunological involvement in the mechanism of CaPterin antitumor activity since nude mice possess B-cell capability while SCID mice do not. This comparison also indicates that there is no measurable direct cancer cell toxicity from the CaPterin. Results showing no CaPterin antitumor efficacy against EMT6 tumor cells implanted in Balb/c mice also suggest an antitumor mechanism involving B-cells, since transforming growth factor beta (TGF-beta), produced by EMT6 cells, is known to cause B-cell apoptosis. These results indicate that CaPterin’s antitumor mechanism involves antibody-dependent cellular cytotoxicity (ADCC) mediated, for example, by natural killer (NK) cells, interleukin-2.

Further study of the immunomodulatory properties of CaPterin was performed by Moheno and co-workers (Winkler et al., Immunobiology (2006) 211, 779-84). They found that CaPterin was able to suppress both the activity of IDO, the degradation of tryptophan and the production of neopterin in PHA- and Con A-stimulated PBMC in a dose-dependent manner. In PHA- and Con A-stimulated PBMC, the production of IFN-γ is increased and induces the degradation of tryptophan and the production of neopterin. Accelerated tryptophan degradation and high IDO expression levels have been associated with poor prognosis in cancer patients.

Provided herein is diterpinyi calcium pentahydrate (DCP), which is suitable as an antitumor agent. Antitumor dose-response data are presented for diterpinyi calcium pentahydrate (DCP) at two dosages.

Therapeutically effective amounts of diterpinyi calcium pentahydrate may be administered as an aqueous suspension. As well, it is contemplated to administer DCP as the active ingredient in a pharmaceutical composition. Accordingly, provided herein are pharmaceutical compositions, which include therapeutically effective amounts of diterpinyi calcium pentahydrate and one or more pharmaceutically acceptable carriers, diluents, or excipients. The carrier(s), diluent(s) or excipient(s) must be acceptable in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof. According to another aspect of the invention there is also provided a process for the preparation of a pharmaceutical formulation including admixing diterpinyi calcium pentahydrate with one or more pharmaceutically acceptable carriers, diluents or excipients.

Pharmaceutical formulations adapted for oral administration may be presented as discrete units such as capsules or tablets; powders or granules; solutions or suspensions in aqueous or non-aqueous liquids; edible foams or whips; or oil-in-water liquid emulsions or water-in-oil liquid emulsions.

For instance, powders are prepared by comminuting the compound to a suitable fine size and mixing with a similarly comminuted pharmaceutical carrier such as an edible carbohydrate, as, for example, starch or mannitol. Flavoring, preservative, dispersing and coloring agents can also be present.

Capsules are made by preparing a powder mixture as described above, and filling formed gelatin sheaths. Glidants and lubricants such as colloidal silica, talc, magnesium stearate, calcium stearate or solid polyethylene glycol can be added to the powder mixture before the filling operation. A disintegrating or solubilizing agent such as agar-agar, calcium carbonate or sodium carbonate can also be added to improve the availability of the medicament when the capsule is ingested.

Moreover, when desired or necessary, suitable binders, lubricants, disintegrating agents and coloring agents can also be incorporated into the mixture. Suitable binders include starch, gelatin, natural sugars such as glucose or beta-lactose, corn sweeteners, natural and synthetic gums such as acesia, tragacanth or sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes and the like. Lubricants used in these dosage forms include sodium oleate, sodium stearate, magnesium stearate, sodium benzoate, sodium acetate, sodium chloride and the like. Disintegrators include, without limitation, starch, methyl cellulose, agar, bentonite, xanthan gum and the like. Tablets are formulated, for example, by preparing a powder mixture, granulating or slugging, adding a lubricant and disintegrant and pressing into tablets. A powder mixture is prepared by mixing the compound, suitably comminuted, with a diluent or base as described above, and optionally, with a binder such as carboxymethylcellulose, an alginate, gelatin, or polyvinyl pyrrolidone, a solution retardant such as paraffin, a resorption accelerator such as a quaternary salt and/or an absorption agent such as bentonite, kaolin or dicalcium phosphate. The powder mixture can be granulated by wetting with a binder such as syrup, starch paste, acacia mucilage or solutions of
cellulosic or polymeric materials and forcing through a screen. As an alternative to granulating, the powder mixture can be run through the tablet machine and the result is imperfectly formed slugs broken into granules. The granules can be lubricated to prevent sticking to the tablet forming dies by means of the addition of stearic acid, a stearate salt, tale or mineral oil. The lubricated mixture is then compressed into tablets. The compounds of the present invention can also be combined with a free flowing inert carrier and compressed into tablets directly without going through the granulating or slugging steps. A clear or opaque protective coating consisting of a sealing coat of shellac, a coating of sugar or polymeric material and a polish coating of wax can be provided. Dye-stuffs can be added to these coatings to distinguish different unit dosages.

[0065] Oral fluids such as solution, syrups and elixirs can be prepared in dosage unit form so that a given quantity contains a predetermined amount of the compound. Syrups can be prepared by dissolving the compound in a suitably flavored aqueous solution, while elixirs are prepared through the use of a non-toxic alcoholic vehicle. Suspensions can be formulated by dispersing the compound in a non-toxic vehicle. Solubilizers and emulsifiers such as ethoxylated isostearyl alcohols and polyoxy ethylene sorbitol ethers, preservatives, flavor additive such as peppermint oil or natural sweeteners or saccharin or other artificial sweeteners, and the like can also be added.

[0066] Where appropriate, dosage unit formulations for oral administration can be microencapsulated. The formulation can also be prepared to prolong or sustain the release as for example by coating or embedding particulate material in polymers, wax or the like.

[0067] One embodiment provides dipterinyl calcium pentahydrate (DCP) or a polymorph thereof. Another embodiment is a method of synthesizing dipterinyl calcium pentahydrate (DCP) comprising dissolving pterin in an aqueous solution of NaOH, adding CaCl2.2H2O to the solution with stirring at a pH of about 11, continuing stirring for about 1 day and collecting the precipitate as DCP.

[0068] In another embodiment is a method of modulating tryptophan production comprising administering to a subject an effective amount of DCP. In yet another embodiment is a method of modulating tryptophan production comprising administering to a subject an effective amount of DCP suspension. In another embodiment is a method of modulating tryptophan degradation comprising administering to a subject an effective amount of DCP. A further embodiment is a method of modulating tryptophan degradation comprising administering to a subject an effective amount of DCP suspension.

[0069] In one embodiment is a method of modulating neopterin production comprising administering to a subject an effective amount of DCP. In a further embodiment is a method of modulating neopterin production comprising administering to a subject an effective amount of DCP suspension.

[0070] In an additional embodiment is a method of modulating IFN-γ production comprising administering to a subject an effective amount of DCP. In yet another embodiment is a method of modulating IFN-γ production comprising administering to a subject an effective amount of DCP suspension.

[0071] In one embodiment is a method of modulating the activity of the enzyme IDO comprising administering to a subject an effective amount of DCP. In another embodiment is a method of modulating the activity of the enzyme IDO comprising administering to a subject an effective amount of DCP suspension.

[0072] In one embodiment is a method of modulating oxidants production comprising administering to a subject an effective amount of DCP. In yet another embodiment is a method of modulating oxidants production comprising administering to a subject an effective amount of DCP suspension.

[0073] In one embodiment is a method of modulating free radical production comprising administering to a subject an effective amount of DCP. In a further embodiment is a method of modulating free radical production comprising administering to a subject an effective amount of DCP suspension. In another embodiment is a method of modulating release of reactive oxygen species (ROS) comprising administering to a subject an effective amount of DCP. In yet another embodiment is a method of modulating release of reactive oxygen species (ROS) comprising administering to a subject an effective amount of DCP suspension.

[0074] In one embodiment is a method of modulating release pro-inflammatory transcription factor NF-κB comprising administering to a subject an effective amount of DCP. In another embodiment is a method of modulating release pro-inflammatory transcription factor NF-κB comprising administering to a subject an effective amount of DCP suspension.

[0075] In another embodiment is a method of modulating the expression of down-stream genes for a cytokine, a chemokine, adhesion molecule, growth factor, enzyme and/or immune receptor comprising administering to a subject an effective amount of DCP. In a further embodiment is a method of modulating the expression of down-stream genes for a cytokine, a chemokine, adhesion molecule, growth factor, enzyme and/or immune receptor comprising administering to a subject an effective amount of DCP suspension.

[0076] In another embodiment is a method of reducing inflammatory activity in a subject comprising administering to a subject an effective amount of DCP. In still another embodiment is a method of reducing inflammatory activity in a subject comprising administering to a subject an effective amount of DCP suspension.

[0077] In another embodiment is a method of reducing or suppressing expression of the enzyme IDO comprising administering to a subject an effective amount of DCP. In a further embodiment is a method of reducing or suppressing expression of the enzyme IDO comprising administering to a subject an effective amount of DCP suspension.

[0078] In another embodiment is a method of modulating human T-cell response comprising administering to a subject an effective amount of DCP. In an additional embodiment is a method of modulating human T-cell response comprising administering to a subject an effective amount of DCP suspension.

[0079] In another embodiment is a method of reducing or suppressing allogeneic immune tolerance comprising administering to a subject an effective amount of DCP. In yet another embodiment is a method of reducing or suppressing allogeneic immune tolerance comprising administering to a subject an effective amount of DCP suspension.

[0080] In another embodiment is a method of modulating immune resistance in human solid tumors comprising administering to a subject an effective amount of DCP. In an additional embodiment is a method of modulating immune resis-
stance in human solid tumors comprising administering to a subject an effective amount of DCP suspension.

[0081] In another embodiment is a method of modulating antiproliferative activity comprising administering to a subject an effective amount of DCP. In still another embodiment is a method of modulating antiproliferative activity comprising administering to a subject an effective amount of DCP suspension.

[0082] In another aspect of the invention is the compound of formula (1): M(pterin), (H2O), wherein: M is a bivalent metal ion selected from the group consisting of Ca2+, Cu2+, Mg2+, V2+, Cr2+, Mn2+, Fe3+, Mo3+, Zn2+, Sr2+, Ba2+, Ra2+, Ru3+, Rh3+, Pd2+, Cd2+, Sn2+, W2+, Re2+, Os2+, Ir2+, Pt2+, Si2+, and Sm2+; X is an integer from 1 to 8; and y is an integer of from 1 to 8. In another embodiment is a method of inhibiting tumor cells in an animal comprising the administration of a therapeutically effective amount of a compound of formula (1): M(pterin), (H2O), wherein: M is a bivalent metal ion selected from the group consisting of Ca2+, Cu2+, Mg2+, V2+, Cr2+, Mn2+, Fe3+, Mo3+, Zn2+, Sr2+, Ba2+, Ra2+, Ru3+, Rh3+, Pd2+, Cd2+, Sn2+, W2+, Re2+, Os2+, Ir2+, Pt2+, Si2+, and Sm2+; X is an integer from 1 to 8; and y is an integer of from 1 to 8.

[0083] In another embodiment is a method of treating a viral infection comprising the administration of a therapeutically effective amount of a compound of formula (1): M(pterin), (H2O), wherein: M is a bivalent metal ion selected from the group consisting of Ca2+, Cu2+, Mg2+, V2+, Cr2+, Mn2+, Fe3+, Mo3+, Zn2+, Sr2+, Ba2+, Ra2+, Ru3+, Rh3+, Pd2+, Cd2+, Sn2+, W2+, Re2+, Os2+, Ir2+, Pt2+, Si2+, and Sm2+; X is an integer from 1 to 8; and y is an integer of from 1 to 8.

[0084] In another embodiment is a pharmaceutical composition comprising a pharmaceutically acceptable excipient and a compound of formula (1): M(pterin), (H2O), wherein: M is a bivalent metal ion selected from the group consisting of Ca2+, Cu2+, Mg2+, V2+, Cr2+, Mn2+, Fe3+, Mo3+, Zn2+, Sr2+, Ba2+, Ra2+, Ru3+, Rh3+, Pd2+, Cd2+, Sn2+, W2+, Re2+, Os2+, Ir2+, Pt2+, Si2+, and Sm2+; X is an integer from 1 to 8; and y is an integer of from 1 to 8.

EXAM P L E S

Example 1

Synthesis of Dipterinyl Calcium Pentahydrate (DCP)

[0085] Pure pterin (81.7 mg, 0.5 mmol) was dissolved in H2O (50 ml) and 0.1 N NaOH (6 ml) and CaCl2.2H2O (36.7 mg, 0.25 mmol) was added to the clear solution with stirring (pH 10.93). A yellowish precipitate was formed within a few minutes. Stirring was continued for 1 day and then the precipitate collected and dried in a vacuum desiccator to give 75 mg. The elemental analysis is consistent with (C14H10N4O2)2Ca.5H2O (MW 454.4).

Calculated: C: 31.74  H: 4.00  N: 30.85
Found: C: 31.22  H: 3.97  N: 29.83.

[0086] The comparison of the extinctions of the UV spectra of pterin and (C14H10N4O2)2Ca.5H2O taken at pH 13 show the following:

| Pterin: | 223 nm (8,700), 250 nm (21,380), 357 nm (8,510) |
| (C14H10N4O2)2Ca.5H2O: | 223 nm (14,450), 250 nm (39,810), 357 nm (13,490) |

[0087] The structure of DCP as determined by single crystal x-ray diffraction is shown in FIG. 2.

[0088] Dipterinyl calcium pentahydrate (DCP) suspensions:

A 1.1 mg/ml suspension was prepared by mixing 44 mg dipterinyl calcium pentahydrate in 40 ml distilled H2O. A 3.3 mg/ml suspension was prepared by mixing 132 mg dipterinyl calcium pentahydrate in 40 ml distilled H2O.

Example 2

In Vivo Testing

[0089] Cell Line Propagation and Inoculation: The MDA-MB-231 human breast tumor cell lines were supplied by SR1 International (Menlo Park, Calif.) and propagated using standard in vitro cell expansion methods. Briefly, cells were grown in L-15 media from Gibco (Cat. No. 11415-064) supplemented with 2 mM L-Glutamine and 10% Fetal Bovine Serum (FBS). The cells were cultured in an incubator with 5% CO2, 37.50 C, and 80% humidity. Cells were harvested with 0.25% (w/v) Trypsin-0.05% (w/v) EDTA solution. Cells were prepared for injection by standard methods to appropriate concentrations. Animals were temporarily restrained but not anesthetized for the inoculation of the tumor cells. Animals were subcutaneously injected with the tumor cells in a 100-200 µl volume.

[0090] Animal Care: The animals were housed 4 to a cage in approved micro-isolator cages. Caging bedding and related items were autoclaved prior to use. No other species were housed in the same room(s) as the experimental animals. The rooms were well ventilated (greater than 10 air changes per hour) with 100% fresh air (no air recirculation). A 12-hour light/12-hour dark photoperiod was maintained, except when room lights were turned on during the dark cycle to accommodate study procedures. Room temperature was maintained between 16-22°C. Animal room and cage cleaning was performed according to Perry Scientific SOP (Standard Operating Procedure). Animals had ad libitum access to irradiated ProLab mouse chow. Autoclaved and chlorinated, municipal tap water was available ad libitum to each animal via water bottles. Treatment of the animals was in accordance with Perry Scientific SOP, which adhered to the regulations outlined in the USDA Animal Welfare Act (9 CFR, Parts 1, 2 and 3) and the conditions specified in The Guide for Care and Use of Laboratory Animals (ILA publication, 1996, National Academy Press). The protocol was approved by Perry Scientific’s Institutional Animal Care and Use Committee prior to initiation of the study. The study conduct was in general compliance with the US FDA Good Laboratory Practice Regulations currently in effect (21 CFR, Part 58).

[0091] Antitumor efficacy was evaluated in nude mice with MDA-MB-231 human tumor xenographs by Perry Scientific (San Diego, Calif.). Twenty-nine athymic nude were each injected subcutaneously with 10x106 MDA-MB-231 cancer cells into the right flank. When tumors reached 3.5 mm in size, the mice were divided into five treatment groups of five each and a control group of four mice. Four of these with
Tumor Outlining Measurements: Each animal was individually tracked for tumor growth by external caliper measurements of protruding tumor. Primary tumor sizes were measured using calipers and an approximate tumor volume calculated using the formula 1/2 (a x b x c), where b was the smallest of two perpendicular diameters.

Results

Fig. 1 shows that calcium pterin (1:4 mol/mol form), dipterinyl calcium pentahydrate (DCP) at both dosages tested, and calcium chloride dihydrate all significantly inhibit MDA-MB-231 xenograft growth in nude mice. These findings identify a new efficacious form of calcium pterin, dipterinyl calcium pentahydrate (DCP). Tumor size data for the control group at days 4 and 7 were missed due to a technical oversight.

Conclusion: Our results show that oral dipterinyl calcium pentahydrate inhibits MDA-MB-231 xenograft tumors in nude mice significantly greater than (1:4 mol/mol) calcium pterin (CalPterin).

Determinant of the optimum dose in nude mice is shown in Fig. 4.

Comparison of DCP to other forms of calcium pterin is shown in Figs. 5 and 6.

Comparison to other common chemotherapeutics is shown in the Figs. 7, 8, and 9.

Comparative data for Figs. 7, 8, and 9 was obtained from the following references:

Paclitaxel:


Doxorubicin:


Docetaxel:


Cisplatin:


Epirubicin:


Osphermifene:


Tamoxifen:


Toremifene:


1. (canceled)

4. A method of treating disease in a subject comprising the administration of a therapeutically effective amount of a compound of formula (I):

\[ M(\text{piperin}), (\text{H}_2\text{O}), \]

wherein:

M is a bivalent metal ion selected from the group consisting of Ca$^{2+}$, Cu$^{2+}$, Mg$^{2+}$, V$^{2+}$, Cr$^{2+}$, Mn$^{2+}$, Fe$^{3+}$, Mo$^{4+}$, Zn$^{2+}$, Sr$^{2+}$, Ba$^{2+}$, Ra$^{2+}$, Ru$^{3+}$, Rh$^{3+}$, Pd$^{2+}$, Cd$^{2+}$, Sn$^{2+}$, W$^{6+}$, Re$^{2+}$, Os$^{2+}$, Ir$^{3+}$, Pt$^{4+}$, Si$^{2+}$, and Sm$^{3+}$;

X is an integer from 1 to 8; and

y is an integer of from 1 to 8.

5. The method of claim 4 wherein the compound of formula (I) is dipipinyl calcium cyclaminate (DPCP).

6. The method of claim 5 wherein the disease is a neoplastic disease.

7. The method of claim 5 wherein the disease is an infectious disease.

8. The method of claim 6 comprising modulating tryptophan degradation.


10. The method of claim 6 comprising modulating IFN-γ production.

11. The method of claim 6 comprising modulating the activity of the enzyme IDO.


13.-21. (canceled)

* * * * *