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DESCRIPTION

1. FIELD

[0001] Provided herein are formulations and dosage forms of (+)-2-[1-(3-ethoxy-4-methoxy-phenyl)-2-methanesulfonyl-ethyl]-4-acetylaminoisoindoline-1,3-dione. Methods of using the formulations and dosage forms are also provided herein.

2. BACKGROUND

[0002] Drug substances are usually administered as part of a formulation in combination with one or more other agents that serve varied and specialized pharmaceutical functions. Dosage forms of various types may be made through selective use of pharmaceutical excipients. As pharmaceutical excipients have various functions and contribute to the pharmaceutical formulations in many different ways, e.g., solubilization, dilution, thickening, stabilization, preservation, coloring, flavoring, etc. The properties that are commonly considered when formulating an active drug substance include bioavailability, ease of manufacture, ease of administration, and stability of the dosage form. Due to the varying properties of the active drug substance to be formulated, dosage forms typically require pharmaceutical excipients that are uniquely tailored to the active drug substance in order to achieve advantageous physical and pharmaceutical properties.

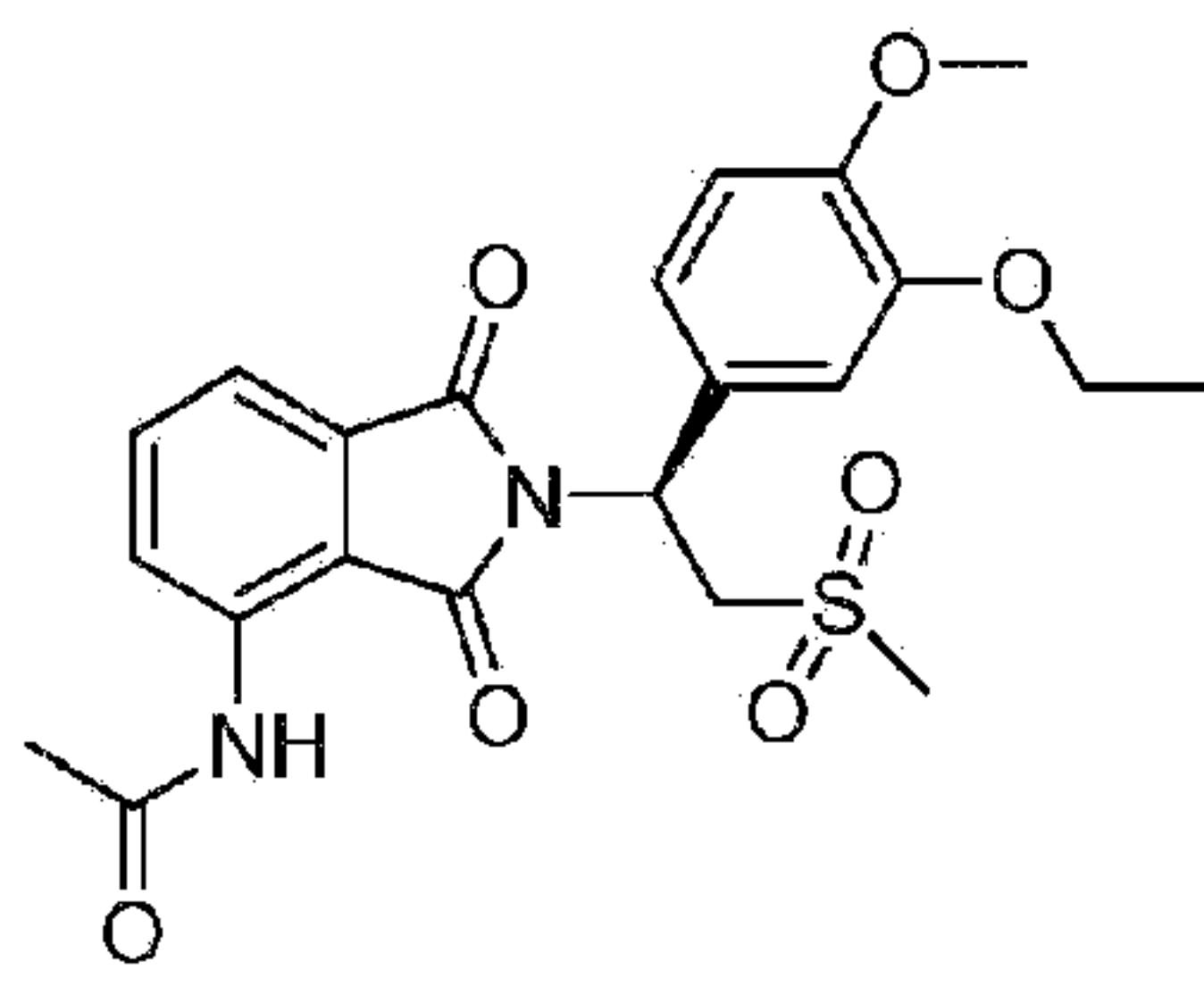
[0003] (+)-2-[1-(3-ethoxy-4-methoxy-phenyl)-2-methanesulfonyl-ethyl]-4-acetylaminoisoindoline-1,3-dione ("Compound A") is a compound with anti-inflammatory activity in clinical development for the treatment of a variety of chronic inflammatory conditions. Pharmacologically, Compound A blocks the degradation of cyclic adenosine monophosphate (cAMP) via inhibition of the phosphodiesterase type IV (PDE4) enzyme, resulting in an increase in cAMP in PDE4-expressing cells including monocytes, T cells, and neutrophils. Enzyme assay data using purified PDE4 enzyme from U937 human monocytic cells indicate that Compound A has a PDE4 IC₅₀ of about 74 nM. Compound A and methods for its synthesis are described, e.g., in U.S. Patent No. 6,962,940.

[0004] Due to its diversified pharmacological properties, Compound A is useful in treating, preventing, and/or managing various diseases or disorders. Thus, a need exists as to dosage forms of Compound A having advantageous physical and pharmaceutical properties. Reference can be made to US2007/155791 and WO01/04195.

3. SUMMARY

[0005] Provided herein is a pharmaceutical composition comprising:

1. (A) compound A:



or a pharmaceutically acceptable salt thereof in an amount of 5% to 25% by weight of the total composition; a filler; a disintegrant; and a lubricant, wherein the filler comprises lactose monohydrate in an amount of 50% to 65% by weight of the total composition; and

2. (B) a coating formulation comprising polyvinyl alcohol.

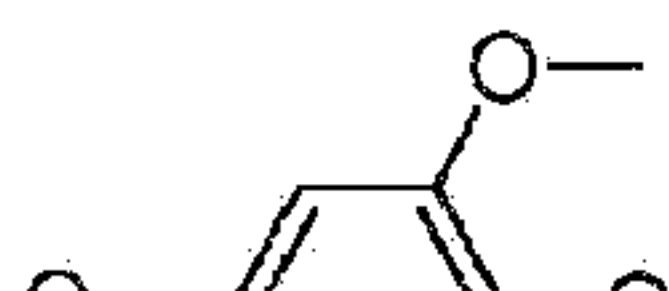
[0006] Also provided herein is the pharmaceutical composition provided herein for use in a method of treating, preventing or managing a disease or disorder in a patient, the method comprising administering to the patient the pharmaceutical composition, wherein the disease or disorder is psoriasis, arthritis, dermatitis, acne, dermatomyositis, ulcerative colitis, Behcet's disease, Crohn's disease, sarcoidosis, uveitis, rosacea or lichen planus.

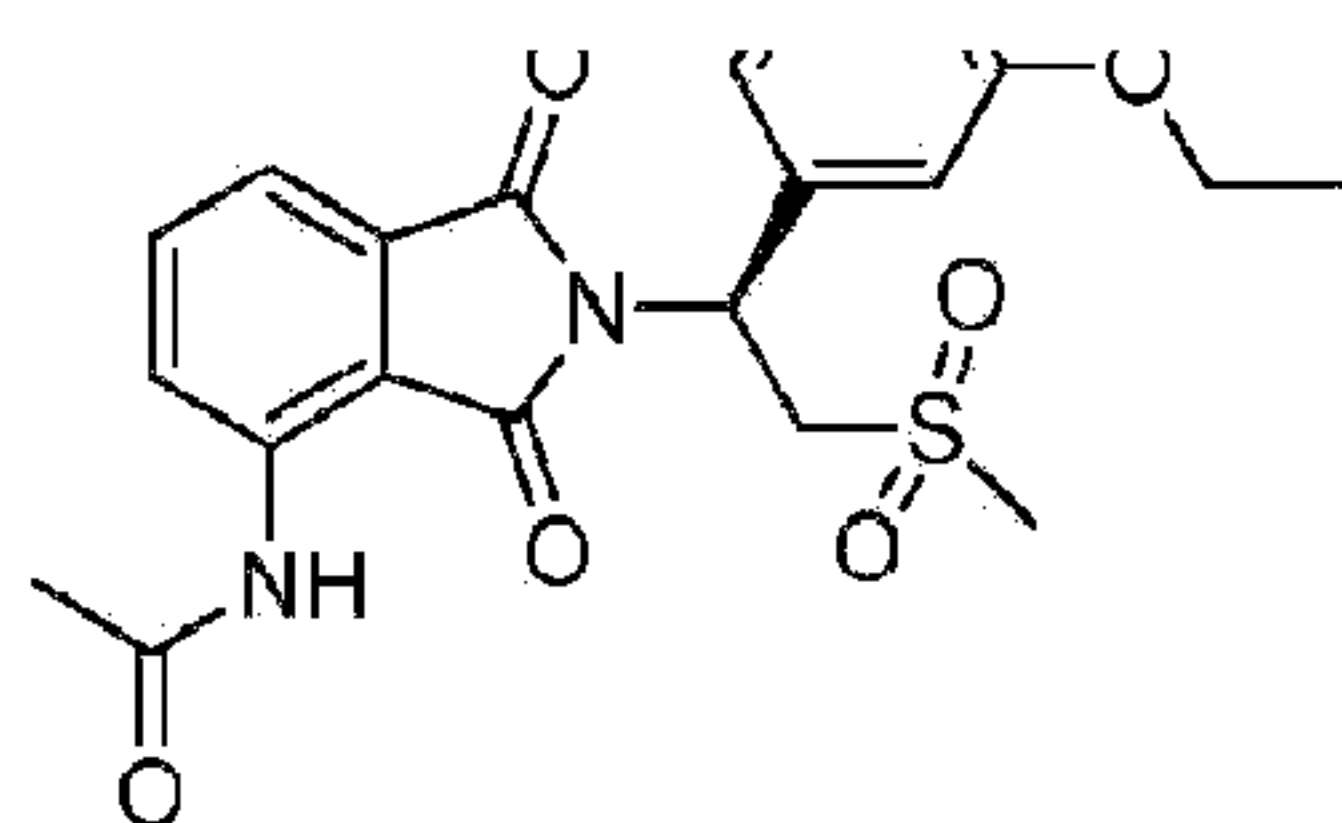
[0007] Also described herein are methods of treating, managing, or preventing diseases and conditions such as, but not limited to, cancer, pain, Macular Degeneration, a skin disease, a pulmonary disorder, an asbestos-related disorder, a parasitic disease, an immunodeficiency disorder, a CNS disorder, CNS injury, atherosclerosis, a sleep disorder, hemoglobinopathy, anemia, an inflammatory disease, an autoimmune disease, a viral disease, a genetic disease, an allergic disease, a bacterial disease, an ocular neovascular disease, a choroidal neovascular disease, a retina neovascular disease, and rubeosis, using Compound A, or a pharmaceutically acceptable stereoisomer, prodrug, salt, solvate, hydrate, or clathrate thereof, in the dosage forms described herein.

4. DETAILED DESCRIPTION

4.1 Definitions

[0008] As used herein, the term "Compound A" refers to enantiomerically pure 2-[1-(3-ethoxy-4-methoxy-phenyl)-2-methanesulfonyl-ethyl]-4-acetylaminoisoindoline-1,3-dione. Without being limited by theory, Compound A is believed to be (+)-2-[1-(3-ethoxy-4-methoxy-phenyl)-2-methanesulfonyl-ethyl]-4-acetylaminoisoindoline-1,3-dione, which has the following structure:





Compound A

[0009] As used herein and unless otherwise indicated, a composition that is "substantially free" of a compound means that the composition contains less than about 20 percent by weight, more preferably less than about 10 percent by weight, even more preferably less than about 5 percent by weight, and most preferably less than about 3 percent by weight of the compound.

[0010] As used herein and unless otherwise indicated, the term "stereomerically pure" means a composition that comprises one stereoisomer of a compound and is substantially free of other stereoisomers of that compound. For example, a stereomerically pure composition of a compound having one chiral center will be substantially free of the opposite enantiomer of the compound. A stereomerically pure composition of a compound having two chiral centers will be substantially free of other diastereomers of the compound. A typical stereomerically pure compound comprises greater than about 80 percent by weight of one stereoisomer of the compound and less than about 20 percent by weight of other stereoisomers of the compound, more preferably greater than about 90 percent by weight of one stereoisomer of the compound and less than about 10 percent by weight of the other stereoisomers of the compound, even more preferably greater than about 95 percent by weight of one stereoisomer of the compound and less than about 5 percent by weight of the other stereoisomers of the compound, and most preferably greater than about 97 percent by weight of one stereoisomer of the compound and less than about 3 percent by weight of the other stereoisomers of the compound.

[0011] As used herein and unless otherwise indicated, the term "enantiomerically pure" means a stereomerically pure composition of a compound having one chiral center.

[0012] As used herein, unless otherwise specified, the term "pharmaceutically acceptable salt(s)" includes, but is not limited to, salts of acidic or basic moieties of a compound provided herein. Basic moieties are capable of forming a wide variety of salts with various inorganic and organic acids. The acids that can be used to prepare pharmaceutically acceptable acid addition salts of such basic compounds are those that form non-toxic acid addition salts, *i.e.*, salts containing pharmacologically acceptable anions. Suitable organic acids include, but are not limited to, maleic, fumaric, benzoic, ascorbic, succinic, acetic, formic, oxalic, propionic, tartaric, salicylic, citric, gluconic, lactic, mandelic, cinnamic, oleic, tannic, aspartic, stearic, palmitic, glycolic, glutamic, gluconic, glucaronic, saccharic, isonicotinic, methanesulfonic, ethanesulfonic, p-toluenesulfonic, benzenesulfonic acids, or pamoic (*i.e.*, 1,1'-methylene-bis-(2-hydroxy-3-naphthoate) acids. Suitable inorganic acids include, but are not limited to, hydrochloric, hydrobromic, hydroiodic, sulfuric, phosphoric, or nitric acids. Compounds that include an amine moiety can form pharmaceutically acceptable salts with various amino acids,

in addition to the acids mentioned above. Chemical moieties that are acidic in nature are capable of forming base salts with various pharmacologically acceptable cations. Examples of such salts are alkali metal or alkaline earth metal salts and, particularly, calcium, magnesium, sodium, lithium, zinc, potassium, or iron salts.

[0013] As used herein, and unless otherwise specified, the term "solvate" means a compound provided herein or a salt thereof, that further includes a stoichiometric or non-stoichiometric amount of solvent bound by non-covalent intermolecular forces. Where the solvent is water, the solvate is a hydrate.

[0014] As used herein and unless otherwise indicated, the term "prodrug" means a derivative of a compound that can hydrolyze, oxidize, or otherwise react under biological conditions (*in vitro* or *in vivo*) to provide the compound. Examples of prodrugs include, but are not limited to, derivatives of thalidomide that include biohydrolyzable moieties such as biohydrolyzable amides, biohydrolyzable esters, biohydrolyzable carbamates, biohydrolyzable carbonates, biohydrolyzable ureides, and biohydrolyzable phosphate analogues. Other examples of prodrugs include derivatives of thalidomide that include -NO, -NO₂, -ONO, or -ONO₂ moieties.

[0015] As used herein and unless otherwise indicated, the terms "biohydrolyzable carbamate," "biohydrolyzable carbonate," "biohydrolyzable ureide," "biohydrolyzable phosphate" mean a carbamate, carbonate, ureide, or phosphate, respectively, of a compound that either: 1) does not interfere with the biological activity of the compound but can confer upon that compound advantageous properties *in vivo*, such as uptake, duration of action, or onset of action; or 2) is biologically inactive but is converted *in vivo* to the biologically active compound. Examples of biohydrolyzable carbamates include, but are not limited to, lower alkylamines, substituted ethylenediamines, aminoacids, hydroxyalkylamines, heterocyclic and heteroaromatic amines, and polyether amines.

[0016] As used herein and unless otherwise indicated, the term "biohydrolyzable ester" means an ester of a compound that either: 1) does not interfere with the biological activity of the compound but can confer upon that compound advantageous properties *in vivo*, such as uptake, duration of action, or onset of action; or 2) is biologically inactive but is converted *in vivo* to the biologically active compound. Examples of biohydrolyzable esters include, but are not limited to, lower alkyl esters, alkoxyacyloxy esters, alkyl acylamino alkyl esters, and choline esters.

[0017] As used herein and unless otherwise indicated, the term "biohydrolyzable amide" means an amide of a compound that either: 1) does not interfere with the biological activity of the compound but can confer upon that compound advantageous properties *in vivo*, such as uptake, duration of action, or onset of action; or 2) is biologically inactive but is converted *in vivo* to the biologically active compound. Examples of biohydrolyzable amides include, but are not limited to, lower alkyl amides, α -amino acid amides, alkoxyacyl amides, and alkylaminoalkylcarbonyl amides.

[0018] As used herein, and unless otherwise specified, the terms "treat," "treating" and "treatment" contemplate an action that occurs while a patient is suffering from the specified disease or disorder, which reduces the severity of the disease or disorder, or retards or slows the progression of the disease or disorder.

[0019] As used herein, and unless otherwise specified, the terms "prevent," "preventing" and "prevention" refer to the prevention of the onset, recurrence or spread of a disease or disorder, or of one or more symptoms thereof. The terms "prevent," "preventing" and "prevention" contemplate an action that occurs before a patient begins to suffer from the specified disease or disorder, which inhibits or reduces the severity of the disease or disorder.

[0020] As used herein, and unless otherwise indicated, the terms "manage," "managing" and "management" encompass preventing the recurrence of the specified disease or disorder in a patient who has already suffered from the disease or disorder, and/or lengthening the time that a patient who has suffered from the disease or disorder remains in remission. The terms encompass modulating the threshold, development and/or duration of the disease or disorder, or changing the way that a patient responds to the disease or disorder.

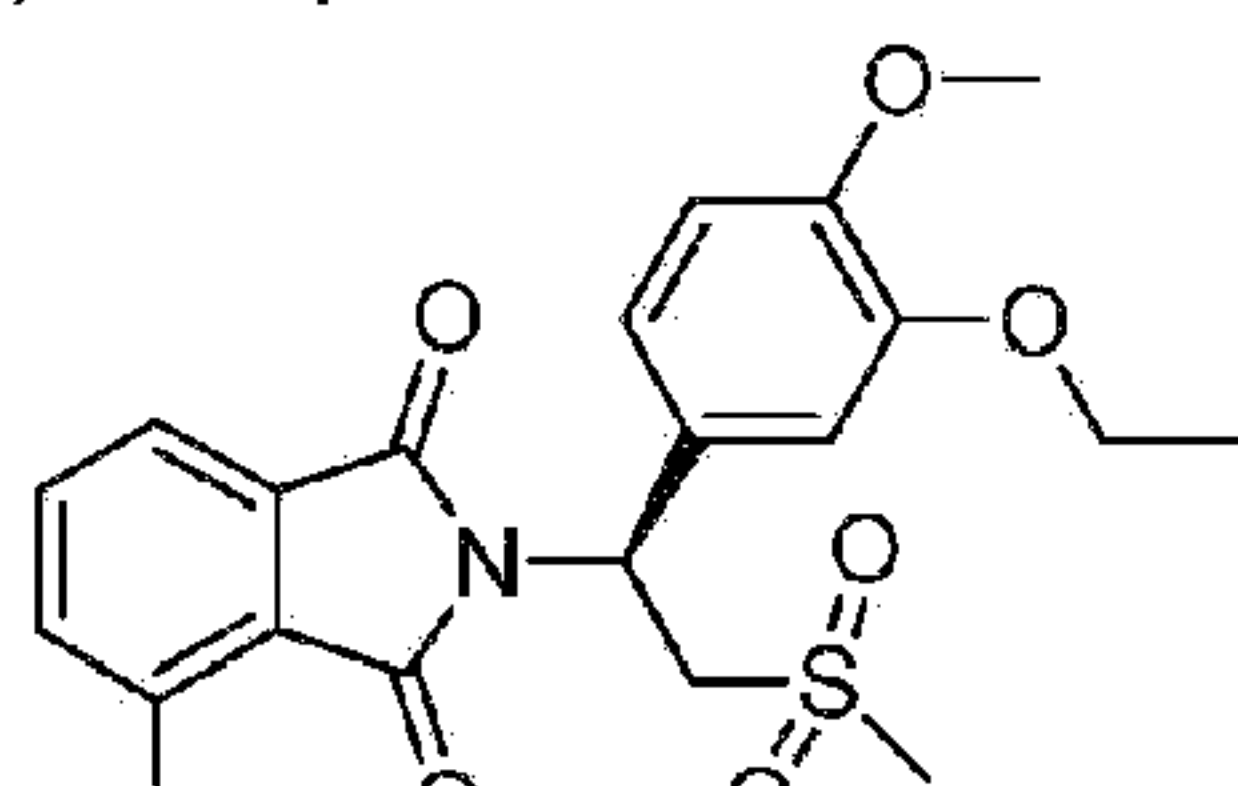
[0021] As used herein, and unless otherwise specified, the term "about," when used in connection with doses, amounts, or weight percent of ingredients of a composition or a dosage form, means dose, amount, or weight percent that is recognized by those of ordinary skill in the art to provide a pharmacological effect equivalent to that obtained from the specified dose, amount, or weight percent is encompassed. Specifically, the term "about" contemplates a dose, amount, or weight percent within 30%, 25%, 20%, 15%, 10%, 5%, 1%, 0.5%, or 0.25% of the specified dose, amount, or weight percent is encompassed.

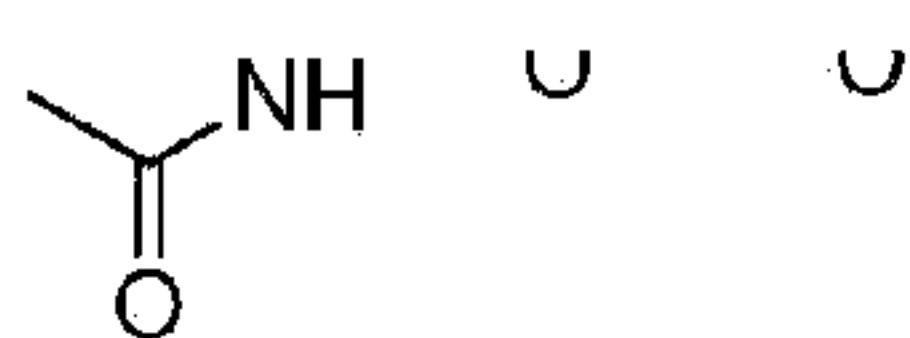
[0022] As used herein, and unless otherwise specified, the term "stable," when used in connection with a formulation or a dosage form, means that the active ingredient of the formulation or dosage form remains solubilized for a specified amount of time and does not significantly degrade or aggregate or become otherwise modified (e.g., as determined, for example, by physical methods such as visual inspection or analytical methods such as HPLC).

4.2 Formulations and Dosage Forms

[0023] Provided herein are pharmaceutical formulations and dosage forms of (+)-2-[1-(3-ethoxy-4-methoxy-phenyl)-2-methanesulfonyl-ethyl]-4-acetylaminoisoindoline-1,3-dione (Compound A) comprising:

1. (A) compound A:





(A)

- or a pharmaceutically acceptable salt thereof in an amount of 5% to 25% by weight of the total composition; a filler; a disintegrant; and a lubricant, wherein the filler comprises lactose monohydrate in an amount of 50% to 65% by weight of the total composition; and
2. (B) a coating formulation comprising polyvinyl alcohol.

[0024] In some embodiments, the dosage forms are suitable for oral administration to a patient. In other embodiments, the formulations and dosage forms provided herein exhibit advantageous physical and/or pharmacological properties. Such properties include, but are not limited to, fast disintegration, low friability, ease of assay, content uniformity, flow properties for manufacture, dissolution and bioavailability, and/or stability. Also described herein are kits comprising pharmaceutical formulations and dosage forms provided herein.

[0025] Also provided herein is the pharmaceutical composition provided herein for use in a method of treating, preventing or managing a disease or disorder in a patient, the method comprising administering to the patient the pharmaceutical composition, wherein the disease or disorder is psoriasis, arthritis, dermatitis, acne, dermatomyositis, ulcerative colitis, Behcet's disease, Crohn's disease, sarcoidosis, uveitis, rosacea or lichen planus.

[0026] In some embodiments, the formulations and dosage forms provided herein are suitable for oral administration, in particular, in the form of a tablet. As provided herein, the formulations and dosage forms comprise a core containing the active ingredient and non-functional film coating.

[0027] In one embodiment, provided herein is a core formulation comprising Compound A. In one embodiment, the core formulation comprises Compound A in an amount of about 5% to about 25%, about 5% to about 20%, about 10% to about 15%, or about 5% to 15% by the weight of the total core composition. In one embodiment, the core formulation comprises Compound A in an amount of about 5%, 10%, 15%, 20% or 25% by the weight of the total core composition. In a specific embodiment, the core formulation comprises Compound A in an amount of about 10% by the weight of the total core composition.

[0028] In one embodiment, the core formulation comprising Compound A further comprises one or more fillers, disintegrants and/or lubricants. As provided herein, the filler is lactose monohydrate (e.g., Fast-Flo®). In one embodiment, the core formulation comprises lactose in an amount of about 50% to about 65% by weight of the total core composition. In one embodiment, the core formulation comprises lactose in an amount of about 50% or 60% by weight of the total core composition. In a specific embodiment, the core formulation comprises lactose in an amount of about 60% by weight of the total core composition.

[0029] In one embodiment, the filler is additionally cellulose. In a specific embodiment, the filler is microcrystalline cellulose (e.g., Avicel®). In one embodiment, the core formulation comprises cellulose in an amount of about 10% to about 50%, about 15% to about 40%, about 20% to about 30%, or about 25% to about 30% by weight of the total core composition. In one embodiment, the core formulation comprises cellulose in an amount of about 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45% and 50% by weight of the total core composition. In a specific embodiment, the core formulation comprises cellulose in an amount of about 26.25% by weight of the total core composition.

[0030] In one embodiment, formulations or dosage forms provided herein may contain two or more fillers.

[0031] In one embodiment, the disintegrant is croscarmellose. In a specific embodiment, the disintegrant is croscarmellose sodium (e.g., Ac-di-sol®). In one embodiment, the core formulation comprises croscarmellose in an amount of about 0.1% to about 10%, about 0.5% to about 8%, about 1% to about 5%, or about 2% to about 8% by weight of the total core composition. In one embodiment, the core formulation comprises croscarmellose in an amount of about 0.1%, 1%, 2%, 3%, 4%, 5%, 6%, 7% or 8% by weight of the total core composition. In a specific embodiment, the core formulation comprises croscarmellose in an amount of about 3% by weight of the total core composition.

[0032] In one embodiment, the lubricant is magnesium stearate. In one embodiment, the core formulation comprises magnesium stearate in an amount of about 0.1% to about 5%, about 0.25% to about 5%, about 0.3% to about 2%, about 0.5% to about 1%, or about 0.5% to about 2% by weight of the total core composition. In one embodiment, the core formulation comprises magnesium stearate in an amount of about 0.1%, 0.2%, 0.3%, 0.4%, 0.5%, 0.6%, 0.7%, 0.8%, 0.9%, 1%, 2%, 3%, 4% or 5% by weight of the total core composition. In a specific embodiment, the core formulation comprises magnesium stearate in an amount of about %by weight of the total core composition.

[0033] In one embodiment, provided herein is a core formulation comprising: Compound A at an amount of about 10% by weight of the total core composition; lactose monohydrate at an amount of about 60% by weight of the total core composition; microcrystalline cellulose at an amount of about 26.25% by weight of the total core composition; croscarmellose at an amount of about 3% by weight of the total core composition; and magnesium stearate at an amount of about 0.75% by weight of the total core composition.

[0034] Without being limited by a particular theory, color change (e.g., fading) is frequently observed in tablet formulations. This phenomenon, known as blooming effect, is caused when lower molecular weight material used in the formulation diffuses to the surface under higher temperature and humidity conditions. For example, excipients such as medium chain triglycerides, which are commonly used as plasticizers in film coatings can migrate or diffuse to the tablet surface causing the color change. However, the phenomenon is not always observed when lower molecular weight materials are used in formulations, *i.e.*, presence of lower

molecular weight materials may or may not cause blooming effect. It was discovered that interaction of a particular lower molecular weight material with other ingredients of the formulation, as well as the actual amount of the lower molecular weight material and other ingredients, are critical in assessing whether a particular formulation would exhibit instability in terms of change of color or appearance.

[0035] Accordingly, in certain embodiments, the formulations or dosage forms provided herein do not comprise a lower molecular weight excipient that may diffuse to the surface of the composition. In other embodiments, the oral formulations or dosage forms provided herein comprise a lower molecular weight excipient, but at an amount that does not trigger blooming effect. In other embodiments, the oral formulations or dosage forms provided herein comprise a lower molecular weight excipient in the presence of other excipients in a way that blooming effect is not observed in connection with the resulting formulations or dosage forms. Consequently, formulations and dosage forms provided herein exhibit improved stability, particularly in terms of change of color or appearance.

[0036] In all embodiments described herein, the pharmaceutical composition provided herein comprise formulations for coating tablets comprising polyvinyl alcohol. In one embodiment, such formulations do not comprise medium chain triglycerides. In such an embodiment, the formulations may contain other various excipients such as, but not limited to, coating agents, binders, lubricants, stabilizing agents, plasticizers, adhesives, glidants, and/or diluents. In some embodiments, the coating formulations provided herein optionally contain coloring agents. These excipients are well-known in the art.

[0037] In one embodiment, provided herein are coating formulations that do not comprise medium chain triglycerides.

[0038] In one embodiment wherein coating formulations do not contain medium chain triglycerides, the excipient is polydextrose. In a specific embodiment, polydextrose is polydextrose FCC. In one embodiment, polydextrose is present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 30% by weight of the total coating formulation. In another embodiment, polydextrose is present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, polydextrose is present at an amount of about 26% by weight of the total coating formulation.

[0039] In one aspect of this embodiment, the excipient is hypromellose. In a specific embodiment, the excipient is hypromellose 15cP. In one embodiment, hypromellose is present at an amount of about 10% to about 60%, about 15% to about 50%, about 25% to about 40%, or about 30% to about 35% by weight of the total coating formulation. In another embodiment, hypromellose is present at an amount of about 5%, 10%, 15%, 20%, 25%, 30% or 35% by weight of the total coating formulation. In a specific embodiment, hypromellose is present at an amount of about 31% by weight of the total coating formulation.

[0040] In another aspect of this embodiment, the excipient is talc. In one embodiment, talc is present at an amount of about 0.1% to about 25%, about 1% to about 20%, about 3% to about 15%, or about 5% to about 10% by weight of the total coating formulation. In another embodiment, talc is present at an amount of about 1%, 2%, 3%, 4%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, talc is present at an amount of about 7% by weight of the total coating formulation.

[0041] In another aspect of this embodiment, the excipient is maltodextrin. In one embodiment, maltodextrin is present at an amount of about 0.1% to about 25%, about 1% to about 20%, about 3% to about 15%, or about 5% to about 10% by weight of the total coating formulation. In another embodiment, maltodextrin is present at an amount of about 1%, 2%, 3%, 4%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, maltodextrin is present at an amount of about 5% by weight of the total coating formulation.

[0042] In another aspect of this embodiment, the excipients are one or more coloring agents, which can be useful in distinguishing dosage forms containing different amounts of active ingredient. Examples of coloring agents include, but are not limited to, iron oxides (e.g., red, yellow and black) and titanium dioxide. Appropriate coloring agents may be mixed to obtain a desired color of the coating formulation. In some embodiments, two or more coloring agents can be used in coating formulations. In one embodiment, the coloring agents comprise titanium dioxide and red iron oxide. In another embodiment, the coloring agents comprise titanium dioxide, red iron oxide, and yellow iron oxide. another embodiment, the coloring agents comprise titanium dioxide, red iron oxide, yellow iron oxide, and black iron oxide.

[0043] In another aspect of this embodiment, the coloring agents are present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 35% by weight of the total coating formulation. In another embodiment, the coloring agents are present at an amount of about 5%, 10%, 15%, 20%, 25%, 30% or 35% by weight of the total coating formulation. In a specific embodiment, the coloring agents are present at an amount of about 31% by weight of the total coating formulation.

[0044] Described herein is a coating formulation comprising: polydextrose at an amount of about 26% by weight of the total coating formulation; hypromellose at an amount of about 31% by weight of the total coating formulation; talc at an amount of about 7% by weight of the total coating formulation; maltodextrin at an amount of about 5% by weight of the total coating formulation; and a mixture of coloring agents at an amount of about 31% by weight of the total coating formulation.

[0045] In another embodiment wherein coating formulations do not contain medium chain triglycerides, the excipient is polydextrose. In a specific embodiment, polydescctrose is polydextrose FCC. In one embodiment, polydextrose is present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 30% by weight of the total coating formulation. In another embodiment, polydextrose is present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In

a specific embodiment, polydextrose is present at an amount of about 26% by weight of the total coating formulation.

[0046] In one aspect of this embodiment, the excipient is hypromellose. In a specific embodiment, the excipient is hypromellose 15cP. In one embodiment, hypromellose is present at an amount of about 10% to about 60%, about 15% to about 50%, about 25% to about 40%, or about 30% to about 35% by weight of the total coating formulation. In another embodiment, hypromellose is present at an amount of about 5%, 10%, 15%, 20%, 25%, 30% or 35% by weight of the total coating formulation. In a specific embodiment, hypromellose is present at an amount of about 31% by weight of the total coating formulation.

[0047] In another aspect of this embodiment, the excipient is talc. In one embodiment, talc is present at an amount of about 0.1% to about 25%, about 1% to about 20%, about 3% to about 15%, or about 5% to about 10% by weight of the total coating formulation. In another embodiment, talc is present at an amount of about 1%, 2%, 3%, 4%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, talc is present at an amount of about 7% by weight of the total coating formulation.

[0048] In another aspect of this embodiment, the excipient is maltodextrin. In one embodiment, maltodextrin is present at an amount of about 0.1% to about 25%, about 1% to about 20%, about 3% to about 15%, or about 5% to about 10% by weight of the total coating formulation. In another embodiment, maltodextrin is present at an amount of about 1%, 2%, 3%, 4%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, maltodextrin is present at an amount of about 5% by weight of the total coating formulation.

[0049] In another aspect of this embodiment, the excipient is triacetin. In one embodiment, triacetin is present at an amount of about 0.1% to about 20%, about 0.5% to about 25%, about 1% to about 10%, or about 1% to about 5% by weight of the total coating formulation. In another embodiment, triacetin is present at an amount of about 1%, 2%, 3%, 4%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, triacetin is present at an amount of about 4% by weight of the total coating formulation.

[0050] In another aspect of this embodiment, the coloring agents are present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 30% by weight of the total coating formulation. In another embodiment, the coloring agents are present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, the coloring agents are present at an amount of about 27% by weight of the total coating formulation.

[0051] Described herein is a coating formulation comprising: polydextrose at an amount of about 26% by weight of the total coating formulation; hypromellose at an amount of about 31% by weight of the total coating formulation; talc at an amount of about 7% by weight of the total coating formulation; maltodextrin at an amount of about 5% by weight of the total coating formulation; triacetin at an amount of about 4% by weight of the total coating formulation; and a

mixture of coloring agents at an amount of about 27% by weight of the total coating formulation.

[0052] In an embodiment wherein coating formulations do not contain medium chain triglycerides, the excipient is polyvinyl alcohol. In one embodiment, polyvinyl alcohol is present at an amount of about 20% to about 75%, about 25% to about 65%, about 30% to about 55%, or about 35% to about 45% by weight of the total coating formulation. In another embodiment, polyvinyl alcohol is present at an amount of about 10%, 20%, 30%, 40%, 50% or 60% by weight of the total coating formulation. In a specific embodiment, polyvinyl alcohol is present at an amount of about 40% by weight of the total coating formulation.

[0053] In one aspect of this embodiment, the excipient is polyethylene glycol. In one specific embodiment, the excipient is polyethylene glycol 3350. In one embodiment, polyethylene glycol is present at an amount of about 5% to about 50%, about 10% to about 40%, about 15% to about 30%, or about 20% to about 25% by weight of the total coating formulation. In another embodiment, polyethylene glycol is present at an amount of about 10%, 20%, 30%, 40% or 50% by weight of the total coating formulation. In a specific embodiment, polyethylene glycol is present at an amount of about 20% (e.g., 20.2%) by weight of the total coating formulation.

[0054] In another aspect of this embodiment, the excipient is talc. In one embodiment, talc is present at an amount of about 1% to about 30%, about 3% to about 25%, about 5% to about 20%, or about 10% to about 15% by weight of the total coating formulation. In another embodiment, talc is present at an amount of about 1%, 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, talc is present at an amount of about 15% (e.g., 14.8%) by weight of the total coating formulation.

[0055] In another aspect of this embodiment, the excipients are one or more coloring agents. In one embodiment, the coloring agents are present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 30% by weight of the total coating formulation. In another embodiment, the coloring agents are present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, the coloring agents are present at an amount of about 25% by weight of the total coating formulation.

[0056] In a specific embodiment, provided herein is a coating formulation comprising: polyvinyl alcohol at an amount of about 40% by weight of the total coating formulation; polyethylene glycol at an amount of about 20% by weight of the total coating formulation; talc at an amount of about 15% by weight of the total coating formulation; and a mixture of coloring agents at an amount of about 25% by weight of the total coating formulation.

[0057] In another embodiment wherein coating formulations do not contain medium chain triglycerides, the excipient is lactose. In one embodiment, lactose is present at an amount of about 10% to about 60%, about 20% to about 50%, about 25% to about 40%, or about 30% to about 35% by weight of the total coating formulation. In another embodiment, lactose is

present at an amount of about 5%, 15%, 25%, 35%, 45%, 55% or 65% by weight of the total coating formulation. In a specific embodiment, polyvinyl alcohol is present at an amount of about 33% by weight of the total coating formulation.

[0058] In one aspect of this embodiment, the excipient is hypromellose. In a specific embodiment, the excipient is hypromellose 6cP. In one embodiment, hypromellose is present at an amount of about 10% to about 60%, about 15% to about 50%, about 25% to about 40%, or about 30% to about 35% by weight of the total coating formulation. In another embodiment, hypromellose is present at an amount of about 5%, 10%, 15%, 20%, 25%, 30% or 35% by weight of the total coating formulation. In a specific embodiment, hypromellose is present at an amount of about 31% by weight of the total coating formulation.

[0059] In another aspect of this embodiment, the excipient is polyethylene glycol. In one specific embodiment, the excipient is polyethylene glycol 3350. In one embodiment, polyethylene glycol is present at an amount of about 0.1% to about 20%, about 0.3% to about 10%, about 0.5% to about 15%, or about 1% to about 5% by weight of the total coating formulation. In another embodiment, polyethylene glycol is present at an amount of about 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, polyethylene glycol is present at an amount of about 5% by weight of the total coating formulation.

[0060] In another aspect of this embodiment, the excipient is triacetin. In one embodiment, triacetin is present at an amount of about 0.1% to about 20%, about 0.5% to about 25%, about 1% to about 10%, or about 1% to about 5% by weight of the total coating formulation. In another embodiment, triacetin is present at an amount of about 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, triacetin is present at an amount of about 4% by weight of the total coating formulation.

[0061] In another aspect of this embodiment, the excipients are one or more coloring agents. In one embodiment, the coloring agents are present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 30% by weight of the total coating formulation. In another embodiment, the coloring agents are present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, the coloring agents are present at an amount of about 27% by weight of the total coating formulation.

[0062] Described herein is a coating formulation comprising: lactose at an amount of about 33% by weight of the total coating formulation; hypromellose at an amount of about 31% by weight of the total coating formulation; polyethylene glycol at an amount of about 5% by weight of the total coating formulation; triacetin at an amount of about 4% by weight of the total coating formulation; and a mixture of coloring agents at an amount of about 27% of the total weight of the coating formulation.

[0063] In other embodiments, provided herein are coating formulations that contain medium

chain triglycerides and other excipients, yet do not cause blooming effect, e.g., changes in color or appearance under storage.

[0064] In one embodiment where coating formulations contain medium chain triglycerides, medium chain triglycerides are present at an amount of about 0.1 to about 15%, about 0.5 to about 10%, about 1% to about 5%, or about 1% to 3% by weight of the total coating formulation. In another embodiment, medium chain triglycerides are present at an amount of about 0.1%, 0.5%, 1%, 1.5%, 2%, 2.5%, 3%, 3.5%, 4%, 4.5% or 5% by weight of the total coating formulation. In a specific embodiment, medium chain triglycerides are present at an amount of about 2% by weight of the total coating formulation.

[0065] In one aspect of this embodiment, another excipient is polydextrose. In one embodiment, polydextrose is present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 30% by weight of the total coating formulation. In another embodiment, polydextrose is present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, polydextrose is present at an amount of about 26% by weight of the total coating formulation.

[0066] In another aspect of this embodiment, another excipient is hypromellose. In one specific embodiment, hypromellose is hypromellose 15cP. In one embodiment, hypromellose is present at an amount of about 10% to about 60%, about 15% to about 50%, about 25% to about 40%, or about 30% to about 35% by weight of the total coating formulation. In another embodiment, hypromellose is present at an amount of about 5%, 10%, 15%, 20%, 25%, 30% or 35% by weight of the total coating formulation. In a specific embodiment, hypromellose is present at an amount of about 31% by weight of the total coating formulation.

[0067] In another aspect of this embodiment, another excipient is talc. In one embodiment, talc is present at an amount of about 1% to about 20%, about 3% to about 15%, about 5% to about 10%, or about 5% to about 10% by weight of the total coating formulation. In another embodiment, talc is present at an amount of about 1%, 5%, 10%, 15% or 20% by weight of the total coating formulation. In a specific embodiment, talc is present at an amount of about 7% by weight of the total coating formulation.

[0068] In another aspect of this embodiment, another excipient is maltodextrin. In one embodiment, maltodextrin is present at an amount of about 0.1% to about 20%, about 0.5% to about 25%, about 1% to about 10%, or about 1% to about 5% by weight of the total coating formulation. In another embodiment, maltodextrin is present at an amount of about 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, maltodextrin is present at an amount of about 5% by weight of the total coating formulation.

[0069] In another aspect of this embodiment, the excipients are one or more coloring agents. In one embodiment, the coloring agents are present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 30% by weight of

the total coating formulation. In another embodiment, the coloring agents are present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, the coloring agents are present at an amount of about 29% by weight of the total coating formulation.

[0070] Described herein is a coating formulation comprising: polydextrose at an amount of about 26% by weight of the total coating formulation; hypromellose at an amount of about 31% by weight of the total coating formulation; talc at an amount of about 7% by weight of the total coating formulation; maltodextrin at an amount of about 5% by weight of the total coating formulation; medium chain triglycerides at an amount of about 2% by weight of the total coating formulation; and a mixture of coloring agents at an amount of about 29% of the total weight of the coating formulation.

[0071] In another embodiment where coating formulations contain medium chain triglycerides, medium chain triglycerides are present at an amount of about 0.1 to about 15%, about 0.5 to about 10%, about 1% to about 5%, or about 1% to 3% by weight of the total coating formulation. In another embodiment, medium chain triglycerides are present at an amount of about 0.1%, 0.5%, 1%, 1.5%, 2%, 2.5%, 3%, 3.5%, 4%, 4.5% or 5% by weight of the total coating formulation. In a specific embodiment, medium chain triglycerides are present at an amount of about 4% by weight of the total coating formulation.

[0072] In one aspect of this embodiment, another excipient is polydextrose. In one embodiment, polydextrose is present at an amount of about 1% to about 40%, about 5% to about 30%, about 10% to about 20%, or about 10% to about 15% by weight of the total coating formulation. In another embodiment, polydextrose is present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, polydextrose is present at an amount of about 13% by weight of the total coating formulation.

[0073] In another aspect of this embodiment, another excipient is hypromellose. In one specific embodiment, hypromellose is hypromellose 15cP. In one embodiment, hypromellose is present at an amount of about 10% to about 65%, about 20% to about 60%, about 25% to about 50%, or about 35% to about 45% by weight of the total coating formulation. In another embodiment, hypromellose is present at an amount of about 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55% or 60% by weight of the total coating formulation. In a specific embodiment, hypromellose is present at an amount of about 44% by weight of the total coating formulation.

[0074] In another aspect of this embodiment, another excipient is talc. In one embodiment, talc is present at an amount of about 1% to about 20%, about 3% to about 15%, about 5% to about 10%, or about 5% to about 10% by weight of the total coating formulation. In another embodiment, talc is present at an amount of about 1%, 5%, 10%, 15% or 20% by weight of the total coating formulation. In a specific embodiment, talc is present at an amount of about 7% by weight of the total coating formulation.

[0075] In another aspect of this embodiment, another excipient is maltodextrin. In one

embodiment, maltodextrin is present at an amount of about 0.1% to about 20%, about 0.5% to about 25%, about 1% to about 10%, or about 1% to about 5% by weight of the total coating formulation. In another embodiment, maltodextrin is present at an amount of about 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9% or 10% by weight of the total coating formulation. In a specific embodiment, maltodextrin is present at an amount of about 5% by weight of the total coating formulation.

[0076] In another aspect of this embodiment, the excipients are one or more coloring agents. In one embodiment, the coloring agents are present at an amount of about 10% to about 60%, about 15% to about 50%, about 20% to about 40%, or about 25% to about 30% by weight of the total coating formulation. In another embodiment, the coloring agents are present at an amount of about 5%, 10%, 15%, 20%, 25% or 30% by weight of the total coating formulation. In a specific embodiment, the coloring agents are present at an amount of about 27% by weight of the total coating formulation.

[0077] Described herein is a coating formulation comprising: polydextrose at an amount of about 13% by weight of the total coating formulation; hypromellose at an amount of about 44% by weight of the total coating formulation; talc at an amount of about 7% by weight of the total coating formulation; maltodextrin at an amount of about 5% by weight of the total coating formulation; medium chain triglycerides at an amount of about 4% by weight of the total coating formulation; and a mixture of coloring agents at an amount of about 27% of the total weight of the coating formulation.

[0078] Any combination between core formulations and coating formulations provided herein can be used.

[0079] In one embodiment, provided herein is a formulation comprising Compound A or a pharmaceutically acceptable salt thereof; a filler; a disintegrant; and a lubricant, wherein compound A is present at an amount of about 5% to about 25% by weight of the total composition. For example, the filler is lactose and is present at an amount of about 20% to 85% by weight of the total composition. In another aspect of this embodiment, the composition further comprises cellulose as a filler and is present at an amount of about 10% to about 50% by weight of the total composition. In another aspect of this embodiment, the disintegrant is croscarmellose and is present at an amount of about 2% to about 8% by weight of the total composition. In another aspect of this embodiment, the lubricant is magnesium stearate and is present at an amount of about 0.25% to about 5% of the total composition.

[0080] In one embodiment, the pharmaceutical composition is coated with a coating formulation, wherein the coating formulation comprises one or more excipients, wherein the excipient is polyvinyl alcohol and present at an amount of about 35% to about 45% by weight of the total coating formulation. In another embodiment the excipient is polyethylene glycol and is present at an amount of about 20% to about 25% by weight of the total coating formulation. In another embodiment the excipient is talc and is present at an amount of about 10% to about 15% by weight of the total coating formulation. In another embodiment, the excipients are one

or more coloring agents present at an amount of about 25% to about 30% by weight of the total coating formulation.

[0081] In a specific embodiment, provided herein is a formulation comprising (1) Compound A or a pharmaceutically acceptable salt thereof; at an amount of about 5% to about 25% by weight of the total composition; (2) lactose monohydrate at an amount of about 20% to 85% by weight of the total composition; (3) cellulose at an amount of about 10% to about 50% by weight of the total composition; (4) croscarmellose at an amount of about 2% to about 8% by weight of the total composition; and (5) magnesium stearate at an amount of about 0.25% to about 5% of the total composition.

[0082] In a specific embodiment, the pharmaceutical composition is coated with a coating formulation, wherein the coating formulation comprises: (1) polyvinyl alcohol at an amount of about 35% to about 45% by weight of the total coating formulation; and/or (2) polyethylene glycol at an amount of about 20% to about 25% by weight of the total coating formulation; and/or (3) talc at an amount of about 10% to about 15% by weight of the total coating formulation; and/or (4) one or more coloring agents at an amount of about 25% to about 30% by weight of the total coating formulation.

[0083] In a specific embodiment, provided herein is a formulation of Compound A, wherein the core formulation comprises the following:

Compound A at an amount of about 10% by weight of the total core composition; lactose monohydrate at an amount of about 60% by weight of the total core composition; microcrystalline cellulose at an amount of about 26.25% by weight of the total core composition; croscarmellose at an amount of about 3% by weight of the total core composition; and magnesium stearate at an amount of about 0.75% by weight of the total core composition; and

the coating formulation comprises the following:

polyvinyl alcohol at an amount of about 40% by weight of the total coating formulation; polyethylene glycol at an amount of about 20% by weight of the total coating formulation; talc at an amount of about 15% by weight of the total coating formulation; and a mixture of coloring agents at an amount of about 25% by weight of the total coating formulation.

[0084] Pharmaceutical compositions and formulations provided herein can be presented as discrete dosage forms. Although a preferred oral dosage unit form is in the form of a tablet, other forms of dosage forms can also be employed. For example, it is possible to use the core formulation in connection with other forms of dosage forms such as a capsule or a caplet. In some embodiments, the formulation is in the form of a tablet.

[0085] In some embodiments, because it is typical to obtain Compound A, or a pharmaceutically acceptable prodrug, salt, solvate, or clathrate thereof, at a purity of less than 100%, the formulations and dosage forms provided herein may be defined as compositions, formulations, or dosage forms that comprise Compound A, or a pharmaceutically acceptable prodrug, salt, solvate, or clathrate thereof, at an amount that provides the potency of a

specified amount of 100% pure Compound A.

[0086] In certain embodiments, provided herein are anhydrous pharmaceutical compositions and dosage forms including an active ingredient, since water can facilitate the degradation of some compounds. For example, the addition of water (e.g., 5 percent) is widely accepted in the pharmaceutical arts as a means of simulating shelf-life, *i.e.*, long-term storage in order to determine characteristics such as shelf-life or the stability of formulations over time. See, e.g., Jens T. Carstensen, Drug Stability: Principles & Practice, 2d. Ed., Marcel Dekker, NY, NY, 1995, pp. 379-80. In effect, water and heat accelerate decomposition. Thus, the effect of water on a formulation can be of great significance since moisture and/or humidity are commonly encountered during manufacture, handling, packaging, storage, shipment, and use of formulations.

[0087] An anhydrous pharmaceutical composition should be prepared and stored such that the anhydrous nature is maintained. Accordingly, in some embodiments, anhydrous compositions are packaged using materials known to prevent exposure to water such that they can be included in suitable formulary kits. Examples of suitable packaging include, but are not limited to, hermetically sealed foils, plastic or the like, unit dose containers, blister packs, and strip packs.

[0088] In this regard, also described herein is a method of preparing a solid pharmaceutical formulation including an active ingredient through admixing the active ingredient and an excipient under anhydrous or low moisture/humidity conditions, wherein the ingredients are substantially free of water. The method can further include packaging the anhydrous or non-hygroscopic solid formulation under low moisture conditions. By using such conditions, the risk of contact with water is reduced and the degradation of the active ingredient can be prevented or substantially reduced.

4.1.1. Second Active Agents

[0089] In certain embodiments, provided herein are compositions and dosage forms of Compound A, or a pharmaceutically acceptable prodrug, salt, solvate, or clathrate thereof, which may further comprise one or more secondary active ingredients. Certain combinations may work synergistically in the treatment of particular types of diseases or disorders, and conditions and symptoms associated with such diseases or disorders. Compound A, or a pharmaceutically acceptable prodrug, salt, solvate, or clathrate thereof, can also work to alleviate adverse effects associated with certain second active agents, and *vice versa*.

[0090] Specific second active compounds that can be contained in the formulations and dosage forms provided herein vary depending on the specific indication to be treated, prevented or managed.

[0091] For instance, for the treatment, prevention or management of cancer, second active

agents include, but are not limited to: semaxanib; cyclosporin; etanercept; doxycycline; bortezomib; acivicin; aclarubicin; acodazole hydrochloride; acronine; adozelesin; aldesleukin; altretamine; ambomycin; ametantrone acetate; amsacrine; anastrozole; anthramycin; asparaginase; asperlin; azacitidine; azetepa; azotomycin; batimastat; benzodepa; bicalutamide; bisantrene hydrochloride; bisnafide dimesylate; bizelesin; bleomycin sulfate; brequinar sodium; bropirimine; busulfan; cactinomycin; calusterone; caracemide; carbetimer; carboplatin; carmustine; carubicin hydrochloride; carzelesin; cedefingol; celecoxib; chlorambucil; cirolemycin; cisplatin; cladribine; crisnatol mesylate; cyclophosphamide; cytarabine; dacarbazine; dactinomycin; daunorubicin hydrochloride; decitabine; dexormaplatin; dezaguanine; dezaguanine mesylate; diaziquone; docetaxel; doxorubicin; doxorubicin hydrochloride; droloxifene; droloxifene citrate; dromostanolone propionate; duazomycin; edatrexate; eflornithine hydrochloride; elsamitucin; enloplatin; enpromate; epipropidine; epirubicin hydrochloride; erbulozole; esorubicin hydrochloride; estramustine; estramustine phosphate sodium; etanidazole; etoposide; etoposide phosphate; etoprine; fadrozole hydrochloride; fazarabine; fenretinide; floxuridine; fludarabine phosphate; fluorouracil; flurocitabine; fosquidone; fostriecin sodium; gemcitabine; gemcitabine hydrochloride; hydroxyurea; idarubicin hydrochloride; ifosfamide; ilmofofinc; iproplatin; irinotecan; irinotecan hydrochloride; lanrcotidc acetate; lctrozolc; lcuprolidc acetate; liarozolc hydrochloride; lomctrcxol sodium; lomustinc; losoxantrone hydrochloride; masoprocol; maytansine; mechlorethamine hydrochloride; megestrol acetate; melengestrol acetate; melphalan; menogaril; mercaptopurine; methotrexate; methotrexate sodium; metoprine; meturedopa; mitindomide; mitocarcin; mitocromin; mitogillin; mitomalcin; mitomycin; mitosper; mitotane; mitoxantrone hydrochloride; mycophenolic acid; nocodazole; nogalamycin; ormaplatin; oxisuran; paclitaxel; pegaspargase; peliomycin; pentamustine; peplomycin sulfate; perfosfamide; pipobroman; piposulfan; piroxantrone hydrochloride; plicamycin; plomestane; porfimer sodium; porfiromycin; prednimustine; procarbazine hydrochloride; puromycin; puromycin hydrochloride; pyrazofurin; riboprine; safingol; safingol hydrochloride; semustine; simtrazene; sparfosate sodium; sparsomycin; spirogermanium hydrochloride; spiromustine; spiroplatin; streptonigrin; streptozocin; sulofenur; talisomycin; tecogalan sodium; taxotere; tegafur; teloxantrone hydrochloride; temoporfin; teniposide; teroxirone; testolactone; thiamiprine; thioguanine; thiotepa; tiazofurin; tirapazamine; toremifene citrate; trestolone acetate; triciribine phosphate; trimetrexate; trimetrexate glucuronate; triptorelin; tubulozole hydrochloride; uracil mustard; uredepa; vapreotide; verteporfin; vinblastine sulfate; vincristine sulfate; vindesine; vindesine sulfate; vinepidine sulfate; vinglycinatate sulfate; vinleurosine sulfate; vinorelbine tartrate; vinrosidine sulfate; vinzolidine sulfate; vorozole; zeniplatin; zinostatin; and zorubicin hydrochloride.

[0092] Other second agents include, but are not limited to: 20-epi-1,25 dihydroxyvitamin D3; 5-ethynyluracil; abiraterone; aclarubicin; acylfulvene; adecypenol; adozelesin; aldesleukin; ALL-TK antagonists; altretamine; ambamustine; amidox; amifostine; aminolevulinic acid; amrubicin; amsacrine; anagrelide; anastrozole; andrographolide; angiogenesis inhibitors; antagonist D; antagonist G; antarelix; anti-dorsalizing morphogenetic protein-1; antiandrogen, prostatic carcinoma; antiestrogen; antineoplaston; antisense oligonucleotides; aphidicolin glycinate; apoptosis gene modulators; apoptosis regulators; apurinic acid; ara-CDP-DL-PTBA; arginine

deaminase; asulacrine; atamestane; atrimustine; axinastatin 1; axinastatin 2; axinastatin 3; azasetron; azatoxin; azatyrosine; baccatin III derivatives; balanol; batimastat; BCR/ABL antagonists; benzochlorins; benzoylstaurosporine; beta lactam derivatives; beta-alethine; betaclamycin B; betulinic acid; bFGF inhibitor; bicalutamide; bisantrene; bisaziridinylspermine; bisnafide; bistratene A; bizelesin; breflate; bropirimine; budotitane; buthionine sulfoximine; calcipotriol; calphostin C; camptothecin derivatives; capecitabine; carboxamide-amino-triazole; carboxyamidotriazole; CaRest M3; CARN 700; cartilage derived inhibitor; carzelesin; casein kinase inhibitors (ICOS); castanospermine; cccropin B; cctorclix; chlorlins; chloroquinoxaline sulfonamide; cicaprost; cis-porphyrin; cladribine; clomifene analogues; clotrimazole; collismycin A; collismycin B; combretastatin A4; combretastatin analogue; conagenin; crambescidin 816; crisnatol; cryptophycin 8; cryptophycin A derivatives; curacin A; cyclopentantraquinones; cycloplatin; cypemycin; cytarabine ocfosphate; cytolytic factor; cytostatin; dacliximab; decitabine; dehydrodidemnin B; deslorelin; dexamethasone; dexifosfamide; dexrazoxane; dexverapamil; diaziquone; didemnin B; didox; diethylnorspermine; dihydro-5-azacytidine; dihydrotaxol, 9-; dioxamycin; diphenyl spiromustine; docetaxel; docosanol; dolasetron; doxifluridine; doxorubicin; droloxifene; dronabinol; duocarmycin SA; ebselen; ecomustine; edelfosine; edrecolomab; eflornithine; elemene; emitefur; epirubicin; epristeride; estramustine analogue; estrogen agonists; estrogen antagonists; etanidazole; etoposide phosphate; exemestane; fadrozole; fazarabine; fenretinide; filgrastim; finasteride; flavopiridol; flezelastine; fluasterone; fludarabine; fluorodaunorubicin hydrochloride; forfenimex; formestane; fostriecin; fotemustine; gadolinium texaphyrin; gallium nitrate; galocitabine; ganirelix; gelatinase inhibitors; gemcitabine; glutathione inhibitors; hepsulfam; heregulin; hexamethylene bisacetamide; hypericin; ibandronic acid; idarubicin; idoxifene; idramantone; ilmofosine; ilomastat; imatinib (Gleevec®), imiquimod; immunostimulant peptides; insulin-like growth factor-1 receptor inhibitor; interferon agonists; interferons; interleukins; iobenguane; iododoxorubicin; ipomeanol, 4-; iroplact; irsogladine; isobengazole; isohomohalicondrin B; itasetron; jasplakinolide; kahalalide F; lamellarin-N triacetate; lanreotide; leinamycin; lenograstim; lentinan sulfate; leptolstatin; letrozole; leukemia inhibiting factor; leukocyte alpha interferon; leuprolide+estrogen+progesterone; leuprorelin; levamisole; liarozole; linear polyamine analogue; lipophilic disaccharide peptide; lipophilic platinum compounds; lissoclinamide 7; lobaplatin; lombricine; lometrexol; lonidamine; losoxantrone; loxoribine; lurtotecan; lutetium texaphyrin; lysofylline; lytic peptides; maitansine; mannostatin A; marimastat; masoprocol; maspin; matrilysin inhibitors; matrix metalloproteinase inhibitors; menogaril; merbarone; meterelin; methioninase; metoclopramide; MIF inhibitor; mifepristone; miltefosine; mirimostim; mitoguazone; mitolactol; mitomycin analogues; mitonafide; mitotoxin fibroblast growth factor-saporin; mitoxantrone; mofarotene; molgramostim; Erbitux, human chorionic gonadotrophin; monophosphoryl lipid A+myobacterium cell wall sk; mopidamol; mustard anticancer agent; mycaperoxide B; mycobacterial cell wall extract; myriaporone; N-acetyldinaline; N-substituted benzamides; nafarelin; nagrestip; naloxone+pentazocine; napavin; naphterpin; nartograstim; nedaplatin; nemorubicin; neridronic acid; nilutamide; nisamycin; nitric oxide modulators; nitroxide antioxidant; nitrullyn; oblimcrsen (Genasense®); O6-benzylguanine; octreotide; okicenone; oligonucleotides; onapristone; ondansetron; ondansetron; oracin; oral cytokine inducer; ormaplatin; osaterone; oxaliplatin; oxaunomycin; paclitaxel; paclitaxel analogues; paclitaxel derivatives; palauamine; palmitoylrhizoxin; pamidronic acid; panaxytriol; panomifene;

parabactin; pazelliptine; pegaspargase; peldesine; pentosan polysulfate sodium; pentostatin; pentozole; perflubron; perfosfamide; perillyl alcohol; phenazinomycin; phenylacetate; phosphatase inhibitors; picibanil; pilocarpine hydrochloride; pirarubicin; piritrexim; placetin A; placetin B; plasminogen activator inhibitor; platinum complex; platinum compounds; platinum-triamine complex; porfimer sodium; porfiromycin; prednisone; propyl bis-acridone; prostaglandin J2; proteasome inhibitors; protein A-based immune modulator; protein kinase C inhibitor; protein kinase C inhibitors, microalgal; protein tyrosine phosphatase inhibitors; purine nucleoside phosphorylase inhibitors; purpurins; pyrazoloacridine; pyridoxylated hemoglobin polyoxyethylene conjugate; raf antagonists; raltitrexed; ramosetron; ras farnesyl protein transferase inhibitors; ras inhibitors; ras-GAP inhibitor; retelliptine demethylated; rhenium Re 186 etidronate; rhizoxin; ribozymes; RII retinamide; rohitukine; romurtide; roquinimex; rubiginone B1; ruboxyl; safingol; saintopin; SarCNU; sarcophytol A; sargramostim; Sdi 1 mimetics; semustine; senescence derived inhibitor 1; sense oligonucleotides; signal transduction inhibitors; sizofiran; sobuzoxane; sodium borocaptate; sodium phenylacetate; solverol; somatomedin binding protein; sonermin; sparfosic acid; spicamycin D; spiromustine; splenopentin; spongistatin 1; squalamine; stipiamide; stromelysin inhibitors; sulfinosine; superactive vasoactive intestinal peptide antagonist; suradista; suramin; swainsonine; tallimustine; tamoxifen methiodide; tauromustine; tazarotene; tecogalan sodium; tegafur; tellurapyrylium; telomerase inhibitors; temoporfin; teniposide; tetrachlorodecaoxide; tetrazomine; thaliblastine; thiocoraline; thrombopoietin; thrombopoietin mimetic; thymalfasin; thymopoietin receptor agonist; thymotrinan; thyroid stimulating hormone; tin ethyl etiopurpurin; tirapazamine; titanocene bichloride; topsentin; toremifene; translation inhibitors; tretinoin; triacetyluridine; triciribine; trimetrexate; triptorelin; tropisetron; turosteride; tyrosine kinase inhibitors; tyrphostins; UBC inhibitors; ubenimex; urogenital sinus-derived growth inhibitory factor; urokinase receptor antagonists; vapreotide; variolin B; velaresol; veramine; verdins; verteporfin; vinorelbine; vinxaltine; vitaxin; vorozole; zanoterone; zeniplatin; zilascorb; and zinostatin stimalamer.

[0093] Yet other second active agents include, but are not limited to, 2-methoxyestradiol, telomestatin, inducers of apoptosis in multiple myeloma cells (such as, for example, TRAIL), statins, scmaxanib, cyclosporin, ctancrcpt, doxycycline, bortezomib, oblimcrscn (Genasense®), remicade, docetaxel, cclccoxib, mclphalan, dexamethasone (Decadron®), steroids, gemcitabine, cisplatin, temozolomide, etoposide, cyclophosphamide, temodar, carboplatin, procarbazine, gliadel, tamoxifen, topotecan, methotrexate, Arisa®, taxol, taxotere, fluorouracil, leucovorin, irinotecan, xeloda, CPT-11, interferon alpha, pegylated interferon alpha (e.g., PEG INTRON-A), capecitabine, cisplatin, thiotepa, fludarabine, carboplatin, liposomal daunorubicin, cytarabine, doxetaxol, paclitaxel, vinblastine, IL-2, GM-CSF, dacarbazine, vinorelbine, zoledronic acid, palmitronate, biacin, busulphan, prednisone, bisphosphonate, arsenic trioxide, vincristine, doxorubicin (Doxil®), paclitaxel, ganciclovir, adriamycin, estramustine sodium phosphate (Emcyt®), sulindac, and etoposide.

[0094] In another embodiment, examples of specific second agents according to the indications to be treated, prevented, or managed can be found in the following references; U.S. patent nos. 6,281,230 and 5,635,517; U.S. publication nos. 2004/0220144, 2004/0190609,

2004/0087546, 2005/0203142, 2004/0091455, 2005/0100529, 2005/0143344, 2005/0214328, 2005/0239842, 2006/0122228, 2006/0154880 and 2006/0188475.

[0095] Examples of second active agents that may be used for the treatment, prevention and/or management of pain include, but are not limited to, conventional therapeutics used to treat or prevent pain such as antidepressants, anticonvulsants, antihypertensives, anxiolytics, calcium channel blockers, muscle relaxants, non-narcotic analgesics, opioid analgesics, anti-inflammatories, cox-2 inhibitors, immunomodulatory agents, alpha-adrenergic receptor agonists or antagonists, immunosuppressive agents, corticosteroids, hyperbaric oxygen, ketamine, other anesthetic agents, NMDA antagonists, and other therapeutics found, for example, in the *Physician's Desk Reference* 2003. Specific examples include, but are not limited to, salicylic acid acetate (Aspirin®), celecoxib (Celebrex®), Enbrel®, ketamine, gabapentin (Neurontin®), phenytoin (Dilantin®), carbamazepine (Tegretol®), oxcarbazepine (Trileptal®), valproic acid (Depakene®), morphine sulfate, hydromorphone, prednisone, griseofulvin, pentonium, alendronate, dyphenhydramide, guanethidine, ketorolac (Acular®), thyrocalcitonin, dimethylsulfoxide (DMSO), clonidine (Catapress®), bretylium, ketanserin, reserpine, droperidol, atropine, phentolamine, bupivacaine, lidocaine, acetaminophen, nortriptyline (Pamelor®), amitriptyline (Elavil®), imipramine (Tofranil®), doxepin (Sinequan®), clomipramine (Anafranil®), fluoxetine (Prozac®), sertraline (Zoloft®), naproxen, nefazodone (Serzone®), venlafaxine (Effexor®), trazodone (Desyrel®), bupropion (Wellbutrin®), mexiletine, nifedipine, propranolol, tramadol, lamotrigine, viox, ziconotide, ketamine, dextromethorphan, benzodiazepines, baclofen, tizanidine and phenoxybenzamine.

[0096] Examples of second active agents that may be used for the treatment, prevention and/or management of macular degeneration and related syndromes include, but are not limited to, a steroid, a light sensitizer, an integrin, an antioxidant, an interferon, a xanthine derivative, a growth hormone, a neurotrophic factor, a regulator of neovascularization, an anti-VEGF antibody, a prostaglandin, an antibiotic, a phytoestrogen, an anti-inflammatory compound or an antiangiogenesis compound, or a combination thereof. Specific examples include, but are not limited to, verteporfin, purlytin, an angiostatic steroid, rhuFab, interferon-2 α , pentoxifylline, tin etiopurpurin, motexafin, lucentis, lutetium, 9-fluoro-11,21-dihydroxy-16, 17-1-methylethylidenebis(oxy)pregna-1,4-diene-3,20-dione, latanoprost (see U.S. Patent No. 6,225,348), tetracycline and its derivatives, rifamycin and its derivatives, macrolides, metronidazole (U.S. Patent Nos. 6,218,369 and 6,015,803), genistein, genistin, 6'-O-Mal genistin, 6'-O-Ac genistin, daidzein, daidzin, 6'-O-Mal daidzin, 6'-O-Ac daidzin, glycitein, glycitin, 6'-O-Mal glycitin, biochanin A, formononetin (U.S. Patent No. 6,001,368), triamcinolone acetomide, dexamethasone (U.S. Patent No. 5,770,589), thalidomide, glutathione (U.S. Patent No. 5,632,984), basic fibroblast growth factor (bFGF), transforming growth factor β (TGF- β), brain-derived neurotrophic factor (BDNF), plasminogen activator factor type 2 (PAI-2), EYE101 (Eyetechnopharmaceuticals), LY333531 (Eli Lilly), Miravant, and RETISERT implant (Bausch & Lomb).

[0097] Examples of second active agents that may be used for the treatment, prevention and/or management of skin diseases include, but are not limited to, keratolytics, retinoids, α -

hydroxy acids, antibiotics, collagen, botulinum toxin, interferon, steroids, and immunomodulatory agents. Specific examples include, but are not limited to, 5-fluorouracil, masoprocol, trichloroacetic acid, salicylic acid, lactic acid, ammonium lactate, urea, tretinoin, isotretinoin, antibiotics, collagen, botulinum toxin, interferon, corticosteroid, transretinoic acid and collagens such as human placental collagen, animal placental collagen, Dermalogen, AlloDerm, Fascia, Cymetra, Autologen, Zyderm, Zylplast, Resoplast, and Isolagen.

[0098] Examples of second active agents that may be used for the treatment, prevention and/or management of pulmonary hypertension and related disorders include, but are not limited to, anticoagulants, diuretics, cardiac glycosides, calcium channel blockers, vasodilators, prostacyclin analogues, endothelin antagonists, phosphodiesterase inhibitors (e.g., PDE V inhibitors), endopeptidase inhibitors, lipid lowering agents, thromboxane inhibitors, and other therapeutics known to reduce pulmonary artery pressure. Specific examples include, but are not limited to, warfarin (Coumadin®), a diuretic, a cardiac glycoside, digoxin-oxygen, diltiazem, nifedipine, a vasodilator such as prostacyclin (e.g., prostaglandin 12 (PGI₂), epoprostenol (EPO, Floran®), treprostinil (Remodulin®), nitric oxide (NO), bosentan (Tracleer®), amlodipine, epoprostenol (Floran®), treprostinil (Remodulin®), prostacyclin, tadalafil (Cialis®), simvastatin (Zocor®), omapatrilat (Vanlev®), irbesartan (Avapro®), pravastatin (Pravachol®), digoxin, L-arginine, iloprost, betaprost, and sildenafil (Viagra®).

[0099] Examples of second active agents that may be used for the treatment, prevention and/or management of asbestos-related disorders include, but are not limited to, anthracycline, platinum, alkylating agent, oblimersen (Genasense®), cisplatin, cyclophosphamide, temodar, carboplatin, procarbazine, gliadel, tamoxifen, topotecan, methotrexate, taxotere, irinotecan, capecitabine, cisplatin, thiotepa, fludarabine, carboplatin, liposomal daunorubicin, cytarabine, doxorubicin (Doxil®), paclitaxel, vinblastine, IL-2, GM-CSF, dacarbazine, vinorelbine, zoledronic acid, palmitronate, biaxin, busulphan, prednisone, bisphosphonate, arsenic trioxide, vincristine, doxorubicin (Doxil®), paclitaxel, ganciclovir, adriamycin, bleomycin, hyaluronidase, mitomycin C, mepacrine, thiotepa, tetracycline and gemcitabine.

[0100] Examples of second active agents that may be used for the treatment, prevention and/or management of parasitic diseases include, but are not limited to, chloroquine, quinine, quinidine, pyrimethamine, sulfadiazine, doxycycline, clindamycin, mefloquine, halofantrine, primaquine, hydroxychloroquine, proguanil, atovaquone, azithromycin, suramin, pentamidine, melarsoprol, nifurtimox, benznidazole, amphotericin B, pentavalent antimony compounds (e.g., sodium stibogluconate), interfereon gamma, itraconazole, a combination of dead promastigotes and BCG, leucovorin, corticosteroids, sulfonamide, spiramycin, IgG (serology), trimethoprim, and sulfamethoxazole.

[0101] Examples of second active agents that may be used for the treatment, prevention and/or management of immunodeficiency disorders include, but are not limited to: antibiotics (therapeutic or prophylactic) such as, but not limited to, ampicillin, tetracycline, penicillin, cephalosporins, streptomycin, kanamycin, and erythromycin; antivirals such as, but not limited to, amantadine, rimantadine, acyclovir, and ribavirin; immunoglobulin; plasma; immunologic

enhancing drugs such as, but not limited to, levamisole and isoprinosine; biologics such as, but not limited to, gammaglobulin, transfer factor, interleukins, and interferons; hormones such as, but not limited to, thymic; and other immunologic agents such as, but not limited to, B cell stimulators (e.g., BAFF/BlyS), cytokines (e.g., IL-2, IL-4, and IL-5), growth factors (e.g., TGF- α), antibodies (e.g., anti-CD40 and IgM), oligonucleotides containing unmethylated CpG motifs, and vaccines (e.g., viral and tumor peptide vaccines).

[0102] Examples of second active agents that may be used for the treatment, prevention and/or management of CNS disorders include, but are not limited to: opioids; a dopamine agonist or antagonist, such as, but not limited to, Levodopa, L-DOPA, cocaine, α -methyl-tyrosine, reserpine, tetrabenazine, benzotropine, pargyline, fenoldopam mesylate, cabergoline, pramipexole dihydrochloride, ropinirole, amantadine hydrochloride, selegiline hydrochloride, carbidopa, pergolide mesylate, Sinemet CR, and Symmetrel; a MAO inhibitor, such as, but not limited to, iproniazid, clorgyline, phenelzine and isocarboxazid; a COMT inhibitor, such as, but not limited to, tolcapone and entacapone; a cholinesterase inhibitor, such as, but not limited to, physostigmine salicylate, physostigmine sulfate, physostigmine bromide, meostigmine bromide, neostigmine methylsulfate, ambenonim chloride, edrophonium chloride, tacrine, pralidoxime chloride, obidoxime chloride, trimedoxime bromide, diacetyl monoxim, endrophonium, pyridostigmine, and demecarium; an anti-inflammatory agent, such as, but not limited to, naproxen sodium, diclofenac sodium, diclofenac potassium, celecoxib, sulindac, oxaprozin, diflunisal, etodolac, meloxicam, ibuprofen, ketoprofen, nabumetone, refecoxib, methotrexate, leflunomide, sulfasalazine, gold salts, Rho-D Immune Globulin, mycophenylate mofetil, cyclosporine, azathioprine, tacrolimus, basiliximab, daclizumab, salicylic acid, acetylsalicylic acid, methyl salicylate, diflunisal, salsalate, olsalazine, sulfasalazine, acetaminophen, indomethacin, sulindac, mefenamic acid, meclofenamate sodium, tolmetin, ketorolac, dichlofenac, flurbiprofen, oxaprozin, piroxicam, meloxicam, ampiroxicam, droxicam, piroxicam, tenoxicam, phenylbutazone, oxyphenbutazone, antipyrine, aminopyrine, apazone, zileuton, aurothioglucose, gold sodium thiomalate, auranofin, methotrexate, colchicine, allopurinol, probenecid, sulfapyrazone and benzbromarone or betamethasone and other glucocorticoids; and an antiemetic agent, such as, but not limited to, metoclopramide, domperidone, prochlorperazine, promethazine, chlorpromazine, trimethobenzamide, ondansetron, granisetron, hydroxyzine, acetylleucine monoethanolamine, alizapride, azasetron, benzquinamide, bethanidine, bromopride, buclizine, clobopride, cyclizine, dimenhydrinate, diphenidol, dolasetron, meclizine, methallatal, metopimazine, nabilone, oxypendyl, pipamazine, scopolamine, sulpiride, tetrahydrocannabinol, thiethylperazine, thioproperazine, tropisetron, and a mixture thereof.

[0103] Examples of second active agents that may be used for the treatment, prevention and/or management of CNS injuries and related syndromes include, but are not limited to, immunomodulatory agents, immunosuppressive agents, antihypertensives, anticonvulsants, fibrinolytic agents, antiplatelet agents, antipsychotics, antidepressants, benzodiazepines, buspirone, amantadine, and other known or conventional agents used in patients with CNS injury/damage and related syndromes. Specific examples include, but are not limited to: steroids (e.g., glucocorticoids, such as, but not limited to, methylprednisolone, dexamethasone

and betamethasone); an anti-inflammatory agent, including, but not limited to, naproxen sodium, diclofenac sodium, diclofenac potassium, celecoxib, sulindac, oxaprozin, diflunisal, etodolac, meloxicam, ibuprofen, ketoprofen, nabumetone, refecoxib, methotrexate, leflunomide, sulfasalazine, gold salts, RHo-D Immune Globulin, mycophenylate mofetil, cyclosporine, azathioprine, tacrolimus, basiliximab, daclizumab, salicylic acid, acetylsalicylic acid, methyl salicylate, diflunisal, salsalate, olsalazine, sulfasalazine, acetaminophen, indomethacin, sulindac, mefenamic acid, meclofenamate sodium, tolmetin, ketorolac, dichlofenac, flurbiprofen, oxaprozin, piroxicam, meloxicam, ampiroxicam, droxicam, pivoxicam, tenoxicam, phenylbutazone, oxyphenbutazone, antipyrine, aminopyrine, apazone, zileuton, aurothioglucose, gold sodium thiomalate, auranofin, methotrexate, colchicine, allopurinol, probenecid, sulfinpyrazone and benzbromarone; a cAMP analog including, but not limited to, db-cAMP; an agent comprising a methylphenidate drug, which comprises 1-threo-methylphenidate, d-threo-methylphenidate, dl-threo-methylphenidate, 1-erythro-methylphenidate, d-erythro-methylphenidate, dl-erythro-methylphenidate, and a mixture thereof; and a diuretic agent such as, but not limited to, mannitol, furosemide, glycerol, and urea.

[0104] Examples of second active agent that may be used for the treatment, prevention and/or management of dysfunctional sleep and related syndromes include, but are not limited to, a tricyclic antidepressant agent, a selective serotonin reuptake inhibitor, an antiepileptic agent (gabapentin, pregabalin, carbamazepine, oxcarbazepine, levitiracetam, topiramate), an antiarrhythmic agent, a sodium channel blocking agent, a selective inflammatory mediator inhibitor, an opioid agent, a second immunomodulatory compound, a combination agent, and other known or conventional agents used in sleep therapy. Specific examples include, but are not limited to, Neurontin, oxycontin, morphine, topiramate, amitryptiline, nortryptiline, carbamazepine, Levodopa, L-DOPA, cocaine, α -methyl-tyrosine, reserpine, tetrabenazine, benzotropine, pargyline, fenodolpam mesylate, cabergoline, pramipexole dihydrochloride, ropinirole, amantadine hydrochloride, selegiline hydrochloride, carbidopa, pramipexole mesylate, Sinemet CR, Symmetrel, iproniazid, clorgyline, phenelzine, isocarboxazid, tolcapone, entacapone, physostigmine salicylate, physostigmine sulfate, physostigmine bromide, meostigmine bromide, neostigmine methylsulfate, ambenonim chloride, edrophonium chloride, tacrine, pralidoxime chloride, obidoxime chloride, trimedoxime bromide, diacetyl monoxime, edrophonium, pyridostigmine, demecarium, naproxen sodium, diclofenac sodium, diclofenac potassium, celecoxib, sulindac, oxaprozin, diflunisal, etodolac, meloxicam, ibuprofen, ketoprofen, nabumetone, refecoxib, methotrexate, leflunomide, sulfasalazine, gold salts, RHo-D Immune Globulin, mycophenylate mofetil, cyclosporine, azathioprine, tacrolimus, basiliximab, daclizumab, salicylic acid, acetylsalicylic acid, methyl salicylate, diflunisal, salsalate, olsalazine, sulfasalazine, acetaminophen, indomethacin, sulindac, mefenamic acid, meclofenamate sodium, tolmetin, ketorolac, diclofenac, flurbiprofen, oxaprozin, piroxicam, meloxicam, ampiroxicam, droxicam, pivoxicam, tenoxicam, phenylbutazone, oxyphenbutazone, antipyrine, aminopyrine, apazone, zileuton, aurothioglucose, gold sodium thiomalate, auranofin, methotrexate, colchicine, allopurinol, probenecid, sulfinpyrazone, benzbromarone, betamethasone and other glucocorticoids, metoclopramide, domperidone, prochlorperazine, promethazine, chlorpromazine, trimethobenzamide, ondansetron, granisetron, hydroxyzine,

acetylleucine monoethanolamine, alizapride, azasetron, benzquinamide, bietanautine, bromopride, buclizine, clebopride, cyclizine, dimenhydrinate, diphenidol, dolasetron, meclizine, methallatal, metopimazine, nabilone, oxyperndyl, pipamazine, scopolamine, sulpiride, tetrahydrocannabinol, thiethylperazine, thioproperazine, tropisetron, and a mixture thereof.

[0105] Examples of second active agents that may be used for the treatment, prevention and/or management of hemoglobinopathy and related disorders include, but are not limited to: interleukins, such as IL-2 (including recombinant IL-II ("rIL2") and canarypox IL-2), IL-10, IL-12, and IL-18; interferons, such as interferon alfa-2a, interferon alfa-2b, interferon alfa-n1, interferon alfa-n3, interferon beta-1 a, and interferon gamma-1 b; and G-CSF; hydroxyurea; butyrates or butyrate derivatives; nitrous oxide; hydroxy urea; HEMOXIN™ (NIPRISAN™; see United States Patent No. 5,800,819); Gardos channel antagonists such as clotrimazole and triaryl methane derivatives; Deferoxamine; protein C; and transfusions of blood, or of a blood substitute such as Hemospan™ or Hemospan™ PS (Sangart).

4.2. Process for Making Dosage Forms

[0106] Dosage forms provided herein can be prepared by any of the methods of pharmacy, but all methods include the step of bringing the active ingredient into association with the excipient, which constitutes one or more necessary ingredients. In general, the compositions are prepared by uniformly admixing (e.g., direct blend) the active ingredient with liquid excipients or finely divided solid excipients or both, and then, if necessary, shaping the product into the desired presentation (e.g., compaction such as roller-compaction). If desired, tablets can be coated by standard aqueous or non-aqueous techniques.

[0107] A dosage form provided herein can be prepared by compression or molding, optionally with one or more accessory ingredients. Compressed tablets can be prepared by compressing in a suitable machine the active ingredient in a free-flowing form such as powder or granules, optionally mixed with an excipient as above and/or a surface active or dispersing agent. Molded tablets can be made by molding in a suitable machine a mixture of the powdered compound moistened with an inert liquid diluent. Encapsulation of the dosage forms provided herein can be done using capsules of methylcellulose, calcium alginate, or gelatin.

[0108] As described herein, the active ingredients and excipients are directly blended and loaded into, for example, a capsule, or compressed directly into tablets. A direct-blended dosage form may be more advantageous than a compacted (e.g., roller-compacted) dosage form in certain instances, since direct-blending can reduce or eliminate the harmful health effects that may be caused by airborne particles of ingredients during the manufacture using compaction process.

[0109] Direct blend formulations may be advantageous in certain instances because they require only one blending step, that of the active and excipients, before being processed into the final dosage form, e.g., tablet or capsule. This can reduce the production of airborne

particle or dust to a minimum, while roller-compaction processes may be prone to produce dust. In roller-compaction process, the compacted material is often milled into smaller particles for further processing. The milling operation can produce significant amounts of airborne particles, since the purpose for this step in manufacturing is to reduce the materials particle size. The milled material is then blended with other ingredients prior to manufacturing the final dosage form.

[0110] For certain active ingredients, in particular for a compound with a low solubility, the active ingredient's particle size is reduced to a fine powder in order to help increase the active ingredient's rate of solubilization. The increase in the rate of solubilization is often necessary for the active ingredient to be effectively absorbed in the gastrointestinal tract. However for fine powders to be directly-blended and loaded onto capsules, the excipients should preferably provide certain characteristics which render the ingredients suitable for the direct-blend process. Examples of such characteristics include, but are not limited to, acceptable flow characteristics. In one embodiment, therefore, provided herein is the use of, and compositions comprising, excipients which may provide characteristics, which render the resulting mixture suitable for direct-blend process, e.g., good flow characteristics. As described herein, a dry blend tablet formulation is the preferred way of making the tablets provided herein.

4.2.1. Screening

[0111] The process for making the pharmaceutical compositions of the invention preferably includes the screening of the active ingredient and the excipient(s). For example, the active ingredient is passed through a screen having openings of about 200 microns to about 750 microns. For example, the active ingredient is passed through a screen with openings of about 200 microns to about 400 microns. For example, the active ingredient is passed through a screen having openings of about 300 to about 400 microns. Depending on the excipient(s) used, the screen openings vary. For example, disintegrants and binders are passed through openings of about 430 microns to about 750 microns, from about 600 microns to about 720 microns, or about 710 microns. Lubricants are typically passed through smaller openings, e.g., about 150 microns to about 250 microns screen. For example, the lubricant is passed through a screen opening of about 210 microns.

4.2.2. Pre-blending

[0112] After the ingredients are screened, the excipient and active ingredient are mixed in a diffusion mixer. For example, the mixing time is from about 1 minute to about 50 minutes, from about 5 minutes to about 45 minutes, from about 10 minutes to about 40 minutes, or from about 10 minutes to about 25 minutes. For example, the mixing time is about 15 minutes.

[0113] When more than one excipient is used, the excipients may be admixed in a tumble

blender for about 1 minute to about 20 minutes, or for about 5 minutes to about 10 minutes, prior to mixing with the active ingredient.

4.2.3. Roller Compaction

[0114] As described herein, the pre-blend may optionally be passed through a roller compactor with a hammer mill attached at the discharge of the compactor.

4.2.4. Final Blend

[0115] When a lubricant, *e.g.*, sodium stearyl fumarate and magnesium stearate, is used, the lubricant is mixed with the pre-blend at the end of the process to complete the pharmaceutical composition. This additional mixing is from about 1 minute to about 10 minutes, or from about 3 minutes to about 5 minutes.

4.2.5. Tableting

[0116] The formulation mixture can be tableted (*e.g.*, via compaction, compression, or molding) into the desired size and shape tablet using, for example, a tablet press or other conventional tableting equipment and standard techniques.

4.2.6. Encapsulation

[0117] The formulation mixture can also be optionally encapsulated into the desired size capsule shell using, for example, a capsule filling machine or a rotary tablet press.

4.3. Kits

[0118] Pharmaceutical packs or kits which comprise pharmaceutical compositions or dosage forms provided herein are also described. An example of a kit comprises notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration.

4.4. Compositions for use in Methods of Treatment, Prevention, and Management

[0119] Provided herein are compositions provide herein for use in methods of treating, preventing, and/or managing certain diseases or disorders.

[0120] Examples of diseases or disorders include, but are not limited to, those disorders related to PDE4, TNF α , cAMP and/or angiogenesis and include diseases or disorders such as various inflammatory diseases, pulmonary diseases, autoimmune diseases and immunological diseases. Specific examples include, but are not limited to, inflammation and various forms thereof, cancer, disorders associated with angiogenesis, pain including, but not limited to, Complex Regional Pain Syndrome ("CRPS"), Macular Degeneration ("MD") and related syndromes, skin diseases, pulmonary disorders, asbestos-related disorders, parasitic diseases, immunodeficiency disorders, CNS disorders, CNS injury, atherosclerosis and related disorders, dysfunctional sleep and related disorders, hemoglobinopathy and related disorders (e.g., anemia), tuberculosis and related disorders, PDE4/TNF α related disorders, infectious diseases, and other various diseases and disorders.

[0121] Exemplary diseases or disorders include, but are not limited to, inflammatory, viral, genetic, allergic, skin, and autoimmune diseases. Specific examples include, but are not limited to, arthritis, HIV, hepatitis, acne, adult respiratory distress syndrome, bone resorption diseases, chronic pulmonary inflammatory diseases, dermatitis, dermatomyositis, cystic fibrosis, Lichen Planus, septic shock, sepsis, endotoxic shock, hemodynamic shock, sepsis syndrome, post ischemic reperfusion injury, meningitis, psoriasis, fibrotic disease, cachexia, graft versus host disease, graft rejection, auto-immune disease, rheumatoid spondylitis, Behcet disease, dermatitis, Crohn's disease, ulcerative colitis, inflammatory-bowel disease, rosacea, multiple sclerosis, systemic lupus erythematosus, ENL in leprosy, sarcoidosis, radiation damage, cancer, asthma, uveitis, or hyperoxic alveolar injury.

[0122] In one embodiment, the disease is psoriasis. In another embodiment, psoriasis is plaque psoriasis.

[0123] In another embodiment, the disease is arthritis. In another embodiment, arthritis is psoriatic arthritis, rheumatoid arthritis, osteoarthritis or acute gouty arthritis.

[0124] In another embodiment, the disease is a skin disease. In another embodiment, skin disease is acne, dermatitis or dermatomyositis. In another embodiment, dermatitis is atopic dermatitis or contact dermatitis.

[0125] In another embodiment, the disease is ulcerative colitis.

[0126] In another embodiment, the disease is Behcet's disease.

[0127] In another embodiment, the disease is Crohn's disease.

[0128] In another embodiment, the disease is sarcoidosis. In another embodiment, sarcoidosis is chronic cutaneous sarcoidosis.

[0129] In another embodiment, the disease is uveitis.

[0130] In another embodiment, the disease is rosacea.

[0131] In another embodiment, the disease is Lichen Planus.

[0132] Also described herein are methods of treating, preventing and/or managing various other diseases or disorders using the compositions and formulations provided herein. Examples of other diseases or disorders are described in the following.

[0133] Examples of cancer and precancerous conditions include, but are not limited to, those described in U.S. patent nos. 6,281,230 and 5,635,517 to Muller et al., in various U.S. patent publications to Zeldis, including publication nos. 2004/0220144A1, published November 4, 2004 (Treatment of Myelodysplastic Syndrome); 2004/0029832A1, published February 12, 2004 (Treatment of Various Types of Cancer); and 2004/0087546, published May 6, 2004 (Treatment of Myeloproliferative Diseases). Examples also include those described in WO 2004/103274, published December 2, 2004.

[0134] Certain examples of cancer include, but are not limited to, cancers of the skin, such as melanoma; lymph node; breast; cervix; uterus; gastrointestinal tract; lung; ovary; prostate; colon; rectum; mouth; brain; head and neck; throat; testes; kidney; pancreas; bone; spleen; liver; bladder; larynx; nasal passages; and AIDS-related cancers. The compounds are also useful for treating cancers of the blood and bone marrow, such as multiple myeloma and acute and chronic leukemias, for example, lymphoblastic, myelogenous, lymphocytic, and myelocytic leukemias. The compounds provided herein can be used for treating, preventing or managing either primary or metastatic tumors.

[0135] Other cancers include, but are not limited to, advanced malignancy, amyloidosis, neuroblastoma, meningioma, hemangiopericytoma, multiple brain metastases, glioblastoma multiforme, glioblastoma, brain stem glioma, poor prognosis malignant brain tumor, malignant glioma, recurrent malignant glioma, anaplastic astrocytoma, anaplastic oligodendroglioma, neuroendocrine tumor, rectal adenocarcinoma, Dukes C & D colorectal cancer, unresectable colorectal carcinoma, metastatic hepatocellular carcinoma, Kaposi's sarcoma, karyotype acute myeloblastic leukemia, chronic lymphocytic leukemia (CLL), Hodgkin's lymphoma, non-Hodgkin's lymphoma, cutaneous T-Cell lymphoma, cutaneous B-Cell lymphoma, diffuse large B-Cell lymphoma, low grade follicular lymphoma, metastatic melanoma (localized melanoma, including, but not limited to, ocular melanoma), malignant mesothelioma, malignant pleural effusion mesothelioma syndrome, peritoneal carcinoma, papillary serous carcinoma, gynecologic sarcoma, soft tissue sarcoma, scleroderma, cutaneous vasculitis, Langerhans cell histiocytosis, leiomyosarcoma, fibrodysplasia ossificans progressiva, hormone refractory prostate cancer, resected high-risk soft tissue sarcoma, unresectable hepatocellular carcinoma, Waldenstrom's macroglobulinemia, smoldering myeloma, indolent myeloma, fallopian tube cancer, androgen independent prostate cancer, androgen dependent stage IV

non-metastatic prostate cancer, hormone-insensitive prostate cancer, chemotherapy-insensitive prostate cancer, papillary thyroid carcinoma, follicular thyroid carcinoma, medullary thyroid carcinoma, and leiomyoma. For example, the cancer is metastatic. For example, the cancer is refractory or resistance to chemotherapy or radiation.

[0136] For example, the diseases or disorders are various forms of leukemias such as chronic lymphocytic leukemia, chronic myelocytic leukemia, acute lymphoblastic leukemia, acute myelogenous leukemia and acute myeloblastic leukemia, including leukemias that are relapsed, refractory or resistant, as disclosed in U.S. publication no. 2006/0030594, published February 9, 2006.

[0137] The term "leukemia" refers malignant neoplasms of the blood-forming tissues. The leukemia includes, but is not limited to, chronic lymphocytic leukemia, chronic myelocytic leukemia, acute lymphoblastic leukemia, acute myelogenous leukemia and acute myeloblastic leukemia. The leukemia can be relapsed, refractory or resistant to conventional therapy. The term "relapsed" refers to a situation where patients who have had a remission of leukemia after therapy have a return of leukemia cells in the marrow and a decrease in normal blood cells. The term "refractory or resistant" refers to a circumstance where patients, even after intensive treatment, have residual leukemia cells in their marrow.

[0138] As described herein, the diseases or disorders are various types of lymphomas, including Non-Hodgkin's lymphoma (NHL). The term "lymphoma" refers a heterogenous group of neoplasms arising in the reticuloendothelial and lymphatic systems. "NHL" refers to malignant monoclonal proliferation of lymphoid cells in sites of the immune system, including lymph nodes, bone marrow, spleen, liver and gastrointestinal tract. Examples of NHL include, but are not limited to, mantle cell lymphoma (MCL), lymphocytic lymphoma of intermediate differentiation, intermediate lymphocytic lymphoma (ILL), diffuse poorly differentiated lymphocytic lymphoma (PDL), centrocytic lymphoma, diffuse small-cleaved cell lymphoma (DSCCL), follicular lymphoma, and any type of the mantle cell lymphomas that can be seen under the microscope (nodular, diffuse, blastic and mentle zone lymphoma).

[0139] Examples of diseases and disorders associated with, or characterized by, undesired angiogenesis include, but are not limited to, inflammatory diseases, autoimmune diseases, viral diseases, genetic diseases, allergic diseases, bacterial diseases, ocular neovascular diseases, choroidal neovascular diseases, retina neovascular diseases, and rubeosis (neovascularization of the angle). Specific examples of the diseases and disorders associated with, or characterized by, undesired angiogenesis include, but arc not limited to, arthritis, endometriosis, Crohn's disease, heart failure, advanced heart failure, renal impairment, endotoxemia, toxic shock syndrome, osteoarthritis, retrovirus replication, wasting, meningitis, silica-induced fibrosis, asbestos-induced fibrosis, veterinary disorder, malignancy-associated hypercalcemia, stroke, circulatory shock, periodontitis, gingivitis, macrocytic anemia, refractory anemia, and 5q-deletion syndrome.

[0140] Examples of pain include, but are not limited to those described in U.S. patent

publication no. 2005/0203142, published September 15, 2005. Specific types of pain include, but are not limited to, nociceptive pain, neuropathic pain, mixed pain of nociceptive and neuropathic pain, visceral pain, migraine, headache and post-operative pain.

[0141] Examples of nociceptive pain include, but are not limited to, pain associated with chemical or thermal burns, cuts of the skin, contusions of the skin, osteoarthritis, rheumatoid arthritis, tendonitis, and myofascial pain.

[0142] Examples of neuropathic pain include, but are not limited to, CRPS type I, CRPS type II, reflex sympathetic dystrophy (RSD), reflex neurovascular dystrophy, reflex dystrophy, sympathetically maintained pain syndrome, causalgia, Sudeck atrophy of bone, algoneurodystrophy, shoulder hand syndrome, post-traumatic dystrophy, trigeminal neuralgia, post herpetic neuralgia, cancer related pain, phantom limb pain, fibromyalgia, chronic fatigue syndrome, spinal cord injury pain, central post-stroke pain, radiculopathy, diabetic neuropathy, post-stroke pain, luetic neuropathy, and other painful neuropathic conditions such as those induced by drugs such as vincristine and velcade.

[0143] As used herein, the terms "complex regional pain syndrome," "CRPS" and "CRPS and related syndromes" mean a chronic pain disorder characterized by one or more of the following: pain, whether spontaneous or evoked, including allodynia (painful response to a stimulus that is not usually painful) and hyperalgesia (exaggerated response to a stimulus that is usually only mildly painful); pain that is disproportionate to the inciting event (e.g., years of severe pain after an ankle sprain); regional pain that is not limited to a single peripheral nerve distribution; and autonomic dysregulation (e.g., edema, alteration in blood flow and hyperhidrosis) associated with trophic skin changes (hair and nail growth abnormalities and cutaneous ulceration).

[0144] Examples of MD and related syndromes include, but are not limited to, those described in U.S. patent publication no. 2004/0091455, published May 13, 2004. Specific examples include, but are not limited to, atrophic (dry) MD, exudative (wet) MD, age-related maculopathy (ARM), choroidal neovascularisation (CNVM), retinal pigment epithelium detachment (PED), and atrophy of retinal pigment epithelium (RPE).

[0145] Examples of skin diseases include, but are not limited to, those described in U.S. publication no. 2005/0214328A1, published September 29, 2005. Specific examples include, but are not limited to, keratoses and related symptoms, skin diseases or disorders characterized with overgrowths of the epidermis, acne, and wrinkles.

[0146] As used herein, the term "keratosis" refers to any lesion on the epidermis marked by the presence of circumscribed overgrowths of the horny layer, including but not limited to actinic keratosis, seborrheic keratosis, keratoacanthoma, keratosis follicularis (Darier disease), inverted follicular keratosis, palmoplantar keratoderma (PPK, keratosis palmaris et plantaris), keratosis pilaris, and stucco keratosis. The term "actinic keratosis" also refers to senile keratosis, keratosis senilis, verruca senilis, plana senilis, solar keratosis, keratoderma or

keratoma. The term "seborrheic keratosis" also refers to seborrheic wart, senile wart, or basal cell papilloma. Keratosis is characterized by one or more of the following symptoms: rough appearing, scaly, erythematous papules, plaques, spicules or nodules on exposed surfaces (e.g., face, hands, ears, neck, legs and thorax), excrescences of keratin referred to as cutaneous horns, hyperkeratosis, telangiectasias, elastosis, pigmented lentigines, acanthosis, parakeratosis, dyskeratoses, papillomatosis, hyperpigmentation of the basal cells, cellular atypia, mitotic figures, abnormal cell-cell adhesion, dense inflammatory infiltrates and small prevalence of squamous cell carcinomas.

[0147] Examples of skin diseases or disorders characterized with overgrowths of the epidermis include, but are not limited to, any conditions, diseases or disorders marked by the presence of overgrowths of the epidermis, including but not limited to, infections associated with papilloma virus, arsenical keratoses, sign of Leser-Trélat, warty dyskeratoma (WD), trichostasis spinulosa (TS), erythrokeratoderma variabilis (EKV), ichthyosis fetalis (harlequin ichthyosis), knuckle pads, cutaneous melanoacanthoma, porokeratosis, psoriasis, squamous cell carcinoma, confluent and reticulated papillomatosis (CRP), acrochordons, cutaneous horn, cowden disease (multiple hamartoma syndrome), dermatosis papulosa nigra (DPN), epidermal nevus syndrome (ENS), ichthyosis vulgaris, molluscum contagiosum, prurigo nodularis, and acanthosis nigricans (AN).

[0148] Examples of pulmonary disorders include, but are not limited to, those described in U.S. publication no. 2005/0239842A1, published October 27, 2005. Specific examples include pulmonary hypertension and related disorders. Examples of pulmonary hypertension and related disorders include, but are not limited to: primary pulmonary hypertension (PPH); secondary pulmonary hypertension (SPH); familial PPH; sporadic PPH; precapillary pulmonary hypertension; pulmonary arterial hypertension (PAH); pulmonary artery hypertension; idiopathic pulmonary hypertension; thrombotic pulmonary arteriopathy (TPA); plexogenic pulmonary arteriopathy; functional classes I to IV pulmonary hypertension; and pulmonary hypertension associated with, related to, or secondary to, left ventricular dysfunction, mitral valvular disease, constrictive pericarditis, aortic stenosis, cardiomyopathy, mediastinal fibrosis, anomalous pulmonary venous drainage, pulmonary venoocclusive disease, collagen vascular disease, congenital heart disease, HIV virus infection, drugs and toxins such as fenfluramines, congenital heart disease, pulmonary venous hypertension, chronic obstructive pulmonary disease, interstitial lung disease, sleep-disordered breathing, alveolar hypoventilation disorder, chronic exposure to high altitude, neonatal lung disease, alveolar-capillary dysplasia, sickle cell disease, other coagulation disorder, chronic thromboemboli, connective tissue disease, lupus including systemic and cutaneous lupus, schistosomiasis, sarcoidosis or pulmonary capillary hemangiomatosis.

[0149] Examples of asbestos-related disorders include, but are not limited to, those described in U.S. publication no. 2005/0100529, published May 12, 2005. Specific examples include, but are not limited to, mesothelioma, asbestosis, malignant pleural effusion, benign exudative effusion, pleural plaques, pleural calcification, diffuse pleural thickening, rounded atelectasis, fibrotic masses, and lung cancer.

[0150] Examples of parasitic diseases include, but are not limited to, those described in U.S. publication no. 2006/0154880, published July 13, 2006. Parasitic diseases include diseases and disorders caused by human intracellular parasites such as, but not limited to, *P. falcifarum*, *P. ovale*, *P. vivax*, *P. malariae*, *L. donovari*, *L. infantum*, *L. aethiopica*, *L. major*, *L. tropica*, *L. mexicana*, *L. braziliensis*, *T. Gondii*, *B. microti*, *B. divergens*, *B. coli*, *C. parvum*, *C. cayetanensis*, *E. histolytica*, *I. belli*, *S. mansonii*, *S. haematobium*, *Trypanosoma ssp.*, *Toxoplasma ssp.*, and *O. volvulus*. Other diseases and disorders caused by non-human intracellular parasites such as, but not limited to, *Babesia bovis*, *Babesia canis*, *Babesia Gibsoni*, *Besnoitia darlingi*, *Cytauxzoon felis*, *Eimeria ssp.*, *Hammondia ssp.*, and *Theileria ssp.*, are also encompassed. Specific examples include, but are not limited to, malaria, babesiosis, trypanosomiasis, leishmaniasis, toxoplasmosis, meningoencephalitis, keratitis, amebiasis, giardiasis, cryptosporidiosis, isosporiasis, cyclosporiasis, microsporidiosis, ascariasis, trichuriasis, ancylostomiasis, strongyloidiasis, toxocariasis, trichinosis, lymphatic filariasis, onchocerciasis, filariasis, schistosomiasis, and dermatitis caused by animal schistosomes.

[0151] Examples of immunodeficiency disorders include, but are not limited to, those described in U.S. publication no. 2006/0188475, published August 24, 2006. Specific examples include, but not limited to, adenosine deaminase deficiency, antibody deficiency with normal or elevated Igs, ataxia-telangiectasia, bare lymphocyte syndrome, common variable immunodeficiency, Ig deficiency with hyper-IgM, Ig heavy chain deletions, IgA deficiency, immunodeficiency with thymoma, reticular dysgenesis, Nezelof syndrome, selective IgG subclass deficiency, transient hypogammaglobulinemia of infancy, Wiscott-Aldrich syndrome, X-linked agammaglobulinemia, X-linked severe combined immunodeficiency.

[0152] Examples of CNS disorders include, but are not limited to, those described in U.S. publication no. 2005/0143344, published June 30, 2005. Specific examples include, but are not limited to, include, but are not limited to, Amyotrophic Lateral Sclerosis, Alzheimer Disease, Parkinson Disease, Huntington's Disease, Multiple Sclerosis other neuroimmunological disorders such as Tourette Syndrome, delirium, or disturbances in consciousness that occur over a short period of time, and amnesic disorder, or discrete memory impairments that occur in the absence of other central nervous system impairments.

[0153] Examples of CNS injuries and related syndromes include, but are not limited to, those described in U.S. publication no. 2006/0122228, published June 8, 2006. Specific examples include, but are not limited to, CNS injury/damage and related syndromes, include, but are not limited to, primary brain injury, secondary brain injury, traumatic brain injury, focal brain injury, diffuse axonal injury, head injury, concussion, post-concussion syndrome, cerebral contusion and laceration, subdural hematoma, epidural hematoma, post-traumatic epilepsy, chronic vegetative state, complete SCI, incomplete SCI, acute SCI, subacute SCI, chronic SCI, central cord syndrome, Brown-Sequard syndrome, anterior cord syndrome, conus medullaris syndrome, cauda equina syndrome, neurogenic shock, spinal shock, altered level of consciousness, headache, nausea, emesis, memory loss, dizziness, diplopia, blurred vision,

emotional lability, sleep disturbances, irritability, inability to concentrate, nervousness, behavioral impairment, cognitive deficit, and seizure.

[0154] Examples of atherosclerosis and related conditions include, but are not limited to, those disclosed in U.S. publication no. 2002/0054899, published May 9, 2002. Specific examples include, but are not limited to, all forms of conditions involving atherosclerosis, including restenosis after vascular intervention such as angioplasty, stenting, atherectomy and grafting. Forms of vascular intervention include diseases of the cardiovascular and renal system, such as, but not limited to, renal angioplasty, percutaneous coronary intervention (PCI), percutaneous transluminal coronary angioplasty (PTCA), carotid percutaneous transluminal angioplasty (PTA), coronary by-pass grafting, angioplasty with stent implantation, peripheral percutaneous transluminal intervention of the iliac, femoral or popliteal arteries, and surgical intervention using impregnated artificial grafts. The following chart provides a listing of the major systemic arteries that may be in need of treatment:

Artery	Body Areas Supplied
Axillary	Shoulder and axilla
Brachial	Upper arm
Brachiocephalic	Head, neck, and arm
Celiac	Divides into left gastric, splenic, and hepatic arteries
Common carotid	Neck
Common iliac	Divides into external and internal iliac arteries
Coronary	Heart
Deep femoral	Thigh
Digital	Fingers
Dorsalis pedis	Foot
External carotid	Neck and external head regions
External iliac	Femoral artery
Femoral	Thigh
Gastric	Stomach
Hepatic	Liver, gallbladder, pancreas, and duodenum
Inferior mesenteric	Descending colon, rectum, and pelvic wall
Internal carotid	Neck and internal head regions
Internal iliac	Rectum, urinary bladder, external genitalia, buttocks muscles, uterus and vagina
Left gastric	Esophagus and stomach
Middle sacral	Sacrum
Ovarian	Ovaries
Palmar arch	Hand

Peroneal	Calf
Popliteal	Knee
Posterior tibial	Calf
Pulmonary	Lungs
Radial	Forearm
Renal	Kidney
Splenic	Stomach, pancreas, and spleen
Subclavian	Shoulder
Superior mesenteric	Pancreas, small intestine, ascending and transverse colon
Testicular	Testes
Ulnar	Forearm

[0155] Examples of dysfunctional sleep and related syndromes include, but are not limited to, those disclosed in U.S. publication no. 2005/0222209A1, published October 6, 2005. Specific examples include, but are not limited to, snoring, sleep apnea, insomnia, narcolepsy, restless leg syndrome, sleep terrors, sleep walking sleep eating, and dysfunctional sleep associated with chronic neurological or inflammatory conditions. Chronic neurological or inflammatory conditions, include, but are not limited to, Complex Regional Pain Syndrome, chronic low back pain, musculoskeletal pain, arthritis, radiculopathy, pain associated with cancer, fibromyalgia, chronic fatigue syndrome, visceral pain, bladder pain, chronic pancreatitis, neuropathies (diabetic, post-herpetic, traumatic or inflammatory), and neurodegenerative disorders such as Parkinson's Disease, Alzheimer's Disease, amyotrophic lateral sclerosis, multiple sclerosis, Huntington's Disease, bradykinesia; muscle rigidity; parkinsonian tremor; parkinsonian gait; motion freezing; depression; defective long-term memory, Rubinstein-Taybi syndrome (RTS); dementia; postural instability; hypokinetic disorders; synuclein disorders; multiple system atrophies; striatonigral degeneration; olivopontocerebellar atrophy; Shy-Drager syndrome; motor neuron disease with parkinsonian features; Lewy body dementia; Tau pathology disorders; progressive supranuclear palsy; corticobasal degeneration; frontotemporal dementia; amyloid pathology disorders; mild cognitive impairment; Alzheimer disease with parkinsonism; Wilson disease; Hallervorden-Spatz disease; Chediak-Hagashi disease; SCA-3 spinocerebellar ataxia; X-linked dystonia parkinsonism; prion disease; hyperkinetic disorders; chorea; ballismus; dystonia tremors; Amyotrophic Lateral Sclerosis (ALS); CNS trauma and myoclonus.

[0156] Examples of hemoglobinopathy and related disorders include, but are not limited to, those described in U.S. publication no. 2005/0143420A1, published June 30, 2005. Specific examples include, but are not limited to, hemoglobinopathy, sickle cell anemia, and any other disorders related to the differentiation of CD34+ cells.

[0157] Examples of tuberculosis (TB) and related disorders include, but are not limited to,

those described in PCT publication no. WO 2010/093588, published February 9, 2010. Specific examples include, but are not limited to, pulmonary TB and extrapulmonary TB (remote TB lesions) such as, but not limited to, genitourinary TB (e.g., kidney TB), tuberculous meningitis, military TB, tuberculous peritonitis, tuberculous pericarditis, tuberculous lymphadenitis, TB of bones and joints, gastrointestinal TB, and TB of the liver. Also described herein are methods of treating, preventing, and/or managing the symptoms associated with TB. Examples include, but are not limited to, cough, dyspnea, hilar lymphadenopathy, segmental atelectasis, swelling of the nodes, lobar atelectasis, pulmonary cavitation, fever, unremitting headache, nausea, drowsiness, stupor, coma, stiff neck, weakness, and malaise.

[0158] Disorders related to TB often include other mycobacterial infections, symptom of which resemble those of TB. Examples of such disorders include, but are not limited to, disorders caused by *M. avium* complex (MAC; *M. avium* and *M. intracellulare*), *M. kansasii*, *M. xenopus*, *M. marinum*, *M. ulcerans*, *M. leprae*, and *M. fortuitum* complex (*M. fortuitum* and *M. chelonae*). Examples of disorders caused by these mycobacteria include, but are not limited to, pulmonary diseases, lymphadenitis, cutaneous diseases, wounds, and foreign body infections. Also described herein is the treatment, prevention and/or management of other granulomatous diseases. Examples of such diseases include, but are not limited to: infectious agents caused diseases such as histoplasmosis, cryptococcus, schistosomiasis, and leishmaniasis; allergic reactions caused diseases such as berylliosis; non-infectious agents caused diseases such as aspiration pneumonia and foreign body reaction; genetically caused diseases such as chronic granulomatous disease; and diseases of unknown causes such as sarcoidosis, Crohn's disease and cat-scratch fever.

[0159] Examples of TNF α related disorders include, but are not limited to, those described in WO 98/03502 and WO 98/54170. Specific examples include, but are not limited to: endotoxemia or toxic shock syndrome; cachexia; adult respiratory distress syndrome; bone resorption diseases such as arthritis; hypercalcemia; Graft versus Host Reaction; cerebral malaria; inflammation; tumor growth; chronic pulmonary inflammatory diseases; reperfusion injury; myocardial infarction; stroke; circulatory shock; rheumatoid arthritis; Crohn's disease; HIV infection and AIDS; other disorders such as rheumatoid arthritis, rheumatoid spondylitis, osteoarthritis, psoriatic arthritis and other arthritic conditions, septic shock, sepsis, endotoxic shock, graft versus host disease, wasting, Crohn's disease, ulcerative colitis, multiple sclerosis, systemic lupus erythematosus, ENL in leprosy, HIV, AIDS, and opportunistic infections in AIDS; disorders such as septic shock, sepsis, endotoxic shock, hemodynamic shock and sepsis syndrome, post ischemic reperfusion injury, malaria, mycobacterial infection, meningitis, psoriasis, congestive heart failure, fibrotic disease, cachexia, graft rejection, oncogenic or cancerous conditions, asthma, autoimmune disease, radiation damages, and hyperoxic alveolar injury; viral infections, such as those caused by the herpes viruses; viral conjunctivitis; or atopic dermatitis.

[0160] In other embodiments, the use of formulations, compositions or dosage forms provided herein in various immunological applications, in particular, as vaccine adjuvants, particularly anticancer vaccine adjuvants, as disclosed in U.S. Publication No. 2007/0048327, published

March 1, 2007, is also encompassed. These embodiments also relate to the uses of the compositions, formulations, or dosage forms provided herein in combination with vaccines to treat or prevent cancer or infectious diseases, and other various uses such as reduction or desensitization of allergic reactions.

5. EXAMPLES

[0161] Embodiments provided herein may be more fully understood by reference to the following examples. These examples are meant to be illustrative of pharmaceutical compositions and dosage forms provided herein, but are not in any way limiting.

5.1. Core Formulation

[0162] Table 1 illustrates a batch formulation and oral dosage formulation for a core portion containing Compound A.

Table 1

Trade Name	Common Name	Weight Percent
Compound A		10.00%
316, Fast-Flo®	Lactose Monohydrate	60.00%
Avicel® PH-102	Microcrystalline Cellulose	26.25%
Ac-Di-Sol®	Croscarmellose Sodium	3.00%
	Magnesium stearate	<u>0.75%</u>
		100.0%

[0163] The core formulation was prepared according to the weight percentage provided above. In the table, the value of Compound A assumes the potency of 100% w/w. The actual input weight was adjusted depending on the assigned purity and water content. Lactose was adjusted to maintain the batch weight.

5.2. Coating Formulations

5.2.1. Coating Formulation 1

[0164] Three coating formulations, having pink, brown and beige color, respectively, were made using the ingredients listed in Table 2 below:

Table 2

Ingredient (%w/w)	Pink	Brown	Beige
Poly Vinyl Alcohol	40.0	40.0	40.0
Macrogol/PEG 3350	20.2	20.2	20.2
Talc	14.8	14.8	14.8
Titanium Dioxide	24.6	12.13	22.99
Red Iron Oxide	0.4	1.22	1.18
Yellow Iron Oxide	-	11.65	0.43
Black Iron Oxide	-	-	0.4

5.2.2. Coating Formulation 2 (Reference Example)

[0165] Three coating formulations, having pink, brown and beige color, respectively, were made using the ingredients listed in Table 3 below:

Table 3

Ingredient (%w/w)	Pink	Brown	Beige
Lactose Monohydrate	33.0	33.0	31.0
Hypromellose 6cP	31.0	31.0	31.0
Macrogol/PEG 3350	5.0	5.0	5.0
Triacetin	4.0	4.0	4.0
Titanium Dioxide	26.5	12.47	26.0
Red Iron Oxide	0.5	2.1	1.7
Yellow Iron Oxide	-	12.43	0.6
Black Iron Oxide	-	-	0.7

5.2.3. Coating Formulation 3 (Reference Example)

[0166] Three coating formulations, having pink, brown and beige color, respectively, were made using the ingredients listed in Table 4 below:

Table 4

Ingredient (%w/w)	Pink	Brown	Beige
Polydextrose FCC	26.0	26.0	26.0
Hypromellose 15cP	31.0	31.0	31.0
Talc	7.0	7.0	7.0
Maltodextrin	5.0	5.0	5.0

Ingredient (%w/w)	Pink	Brown	Beige
Medium Chain Triglycerides	2.0	2.0	2.0
Titanium Dioxide	28.5	14.47	26.0
Red Iron Oxide	0.5	2.1	1.7
Yellow Iron Oxide	-	12.43	0.6
Black Iron Oxide	-	-	0.7

5.2.4. Coating Formulation 4 (Reference Example)

[0167] Three coating formulations, having pink, brown and beige color, respectively, were made using the ingredients listed in Table 5 below:

Table 5

Ingredient (%w/w)	Pink	Brown	Beige
Polydextrose FCC	26.0	26.0	26.0
Hypromellose 15cP	31.0	31.0	31.0
Talc	7.0	7.0	7.0
Maltodextrin	5.0	5.0	5.0
Titanium Dioxide	30.5	16.47	28.0
Red Iron Oxide	0.5	2.1	1.7
Yellow Iron Oxide	-	12.43	0.6
Black Iron Oxide	-	-	0.7

5.2.5. Coating Formulation 5 (Reference Example)

[0168] Three coating formulations, having pink, brown and beige color, respectively, were made using the ingredients listed in Table 6 below:

Table 6

Ingredient (%w/w)	Pink	Brown	Beige
Polydextrose FCC	26.0	26.0	26.0
Hypromellose 15cP	31.0	31.0	31.0
Talc	7.0	7.0	7.0
Maltodextrin	5.0	5.0	5.0
Triacetin	4.0	4.0	4.0
Titanium Dioxide	26.5	12.47	24.0

Ingredient (%w/w)	Pink	Brown	Beige
Red Iron Oxide	0.5	2.1	1.7
Yellow Iron Oxide	-	12.43	0.6
Black Iron Oxide	-	-	0.7

5.2.6. Coating Formulation 6 (Reference Example)

[0169] Three coating formulations, having pink, brown and beige color, respectively, were made using the ingredients listed in Table 7 below:

Table 7

Ingredient (%w/w)	Pink	Brown	Beige
Polydextrose FCC	13.0	13.0	13.0
Hypromellose 15cP	44.0	44.0	44.0
Talc	7.0	7.0	7.0
Maltodextrin	5.0	5.0	5.0
Medium Chain Triglycerides	4.0	4.0	4.0
Titanium Dioxide	26.5	12.47	24.0
Red Iron Oxide	0.5	2.1	1.7
Yellow Iron Oxide	-	12.43	0.6
Black Iron Oxide	-	-	0.7

5.3. Tablet Formulation (Core + Coating)

[0170] A complete tablet formulation, including core and coating, was prepared using the ingredients listed in Table 8 below:

Ingredient (%w/w)	Pink	Brown	Beige
<u>Core</u>			
Compound A	10.00	10.00	10.00
Lactose Monohydrate (316, Fast-Flo®)	60.00	60.00	60.00
Microcrystalline Cellulose (Avicel® PH-102)	26.25	26.25	26.25
Croscarmellose Sodium (Ac-di-sol®)	3.00	3.00	3.00
Magnesium Stearate	0.75	0.75	0.75
Total	100	100	100

<u>Coating</u>			
Poly Vinyl Alcohol	40.00	40.00	40.00
Macrogol/PEG 3350	20.20	20.20	20.20
Talc	14.80	14.80	14.80
Titanium Dioxide	24.60	12.13	22.99
Red Iron Oxide	0.40	1.22	1.18
Yellow Iron Oxide	-	11.65	0.43
Black Iron Oxide	-	-	0.40
Total	100	100	100

5.4. Stability of Formulation

[0171] Tablets were coated with different color formulations based on the poly vinyl alcohol. (See Section 5.3, above). Color coating was applied to achieve 4% weight gain of core formulations. For some of the tablets, coating was continued with clear coating suspension to achieve a target of 1% weight gain (*i.e.*, a total of 5% weight gain). Tablets were placed in open dish stability chambers under accelerated temperature and humidity conditions (40°C/75%RH) to assess color changes.

[0172] There was no color fading observed for tablets after 24 hours, 48 hours and 72 hours. A slight dullness in the color was observed for color and clear coated tablets after 72 hours. The results indicate that the formulation described in Section 5.3 above, with or without clear coating, have adequate stability, particularly with regard to color changes.

REFERENCES CITED IN THE DESCRIPTION

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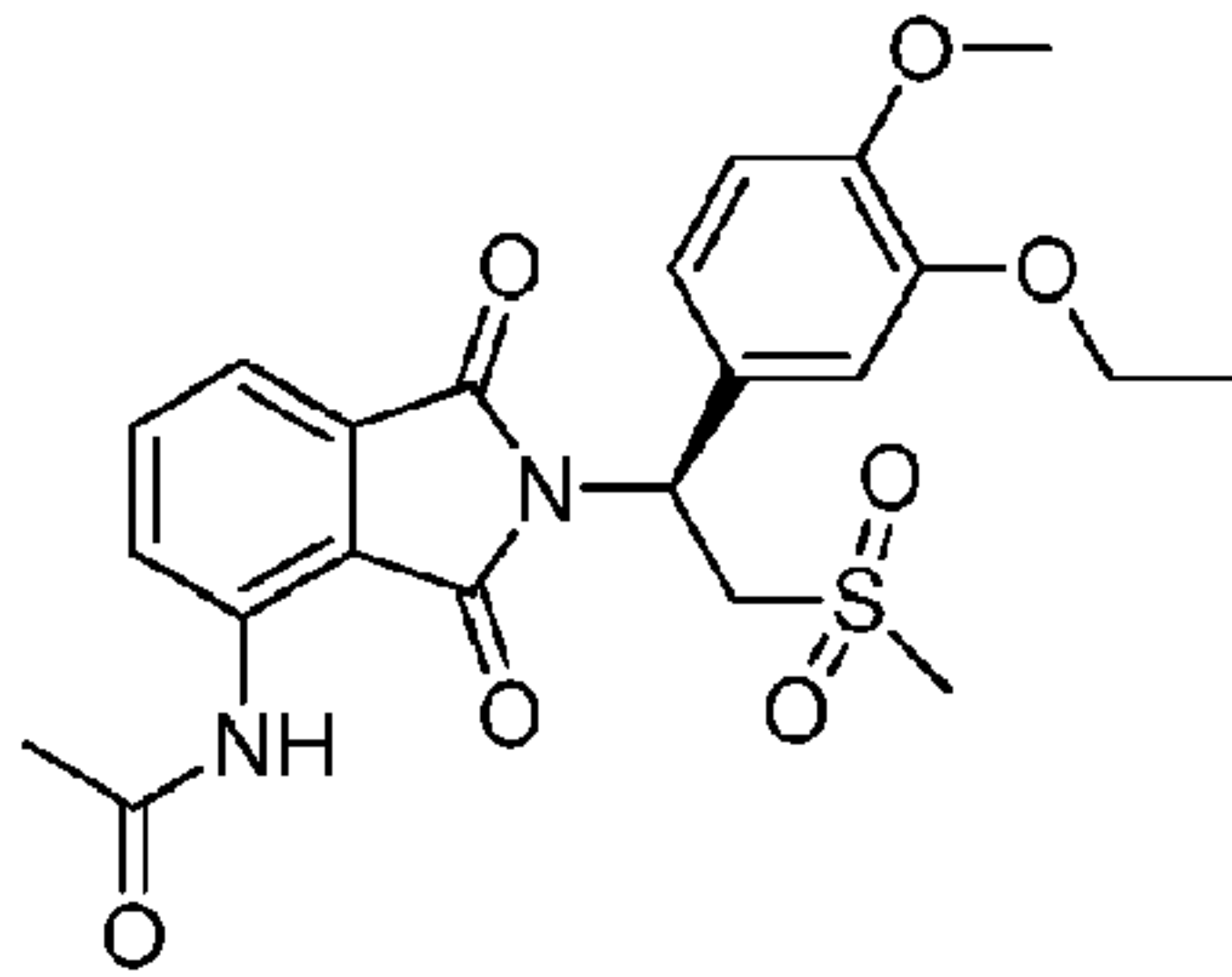
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Patentkrav

1. Farmaceutisk sammensætning, der omfatter:

(A) forbindelse A:



5 (A)

eller et farmaceutisk acceptabelt salt deraf i en mængde på 5 til 25 vægt-% af den samlede sammensætning; et fyldstof; et desintegreringsmiddel; og et smøremiddel,

hvor fyldstoffet omfatter lactosemonohydrat i en mængde på 50 til 65 vægt-% af den samlede sammensætning; og

(B) en overtræksformulering, der omfatter polyvinylalkohol.

2. Farmaceutisk sammensætning ifølge krav 1, hvor polyvinylalkohol er til stede i en mængde på 35 til 45 vægt-% af den samlede overtræksformulering.

3. Farmaceutisk sammensætning ifølge krav 1, hvor sammensætningen yderligere omfatter cellulose som et fyldstof, hvor eventuelt cellulose er til stede i en mængde på 10 til 50 vægt-% af den samlede sammensætning.

4. Farmaceutisk sammensætning ifølge krav 1, hvor desintegreringsmidlet er croscarmellose, hvor eventuelt croscarmellose er til stede i en mængde på 2 til 8 vægt-% af den samlede sammensætning.

5. Farmaceutisk sammensætning ifølge krav 1, hvor smøremidlet er magnesiumstearat, hvor eventuelt magnesiumstearat er til stede i en mængde på 0,25 til 5 % af den samlede sammensætning.

6. Farmaceutisk sammensætning ifølge krav 1, hvor

overtræksformuleringen omfatter et eller flere yderligere excipienser, hvor excipienet er et overtræksmiddel, et bindemiddel, et smøremiddel, et stabiliseringsmiddel, et plastificeringsmiddel, et klæbemiddel, et glidemiddel, et
5 fortyndingsmiddel eller en kombination deraf.

7. Farmaceutisk sammensætning ifølge krav 6, hvor excipienet er polyethylenglycol, hvor eventuelt polyethylenglycol er til stede i en mængde på 20 til 25 vægt-%
10 af den samlede overtræksformulering.

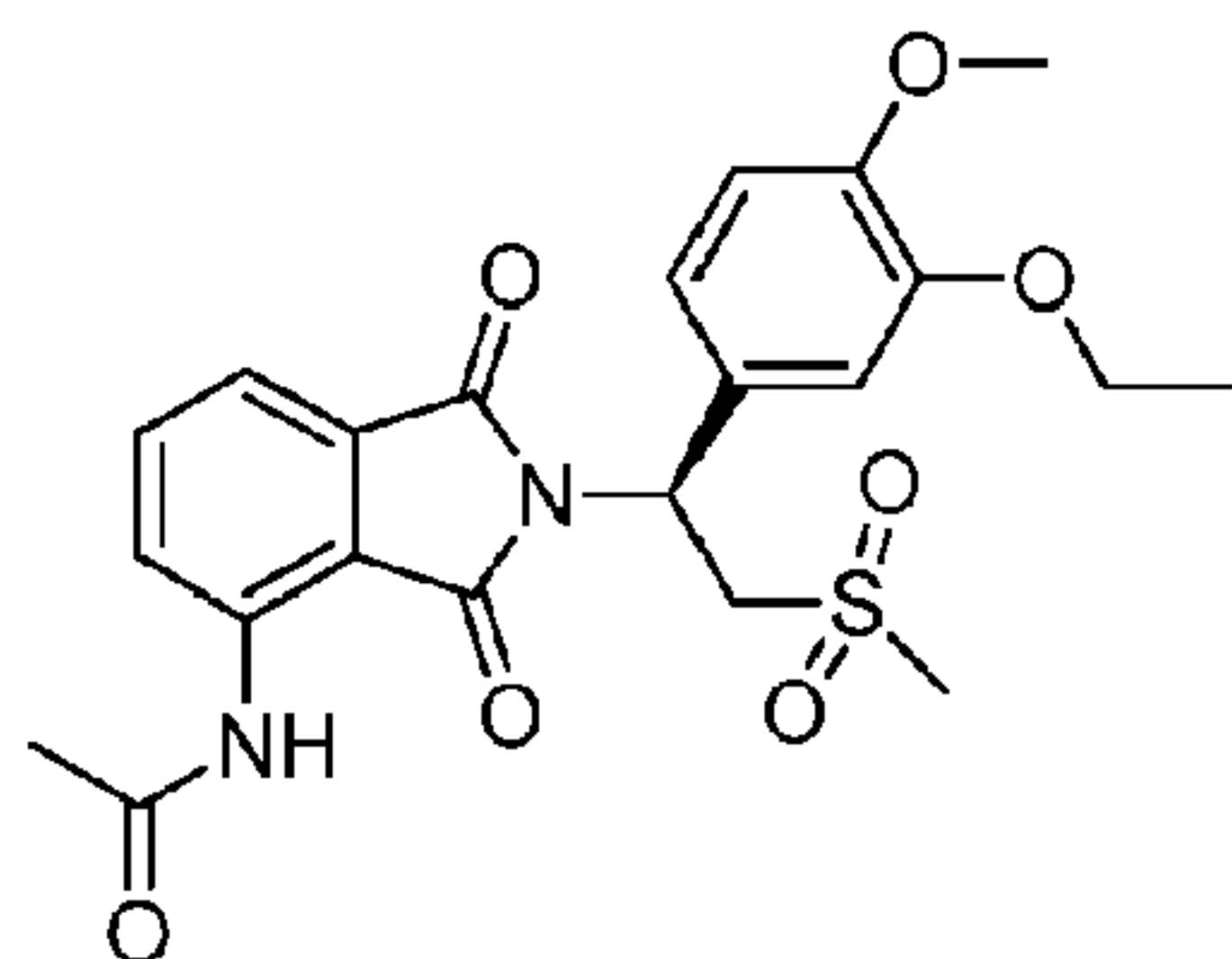
8. Farmaceutisk sammensætning ifølge krav 6, hvor excipienet er talkum, hvor eventuelt talkum er til stede i en mængde på 10 til 15 vægt-% af den samlede
15 overtræksformulering.

9. Farmaceutisk sammensætning ifølge krav 6, hvor excipienserne er et eller flere farvestoffer, hvor eventuelt farvestofferne er til stede i en mængde på 25 til 30 vægt-% af
20 den samlede overtræksformulering.

10. Farmaceutisk sammensætning ifølge krav 1, hvor sammensætningen er en tablet, der omfatter:

(A) en kernesammensætning, hvor kernesammensætningen omfatter:

25 (i) forbindelse A



(A)

eller et farmaceutisk acceptabelt salt deraf i en mængde på 10 vægt-% af den samlede kernesammensætning;

(ii) lactosemonohydrat i en mængde på 60 vægt-% af den samlede
30 kernesammensætning;

(iii) mikrokrySTALLinsk cellulose i en mængde på 26,25 vægt-% af den samlede kernesammensætning;

(iv) croscarmellose i en mængde på 3 vægt-% af den samlede

kernesammensætning; og

(v) magnesiumstearat i en mængde på 0,75 vægt-% af den samlede kernesammensætning; og

(B) en overtræksformulering, hvor overtræksformuleringen
5 omfatter:

(i) polyvinylalkohol i en mængde på 40 vægt-% af den samlede overtræksformulering;

(ii) polyethylenglycol i en mængde på 20 vægt-% af den samlede overtræksformulering;

10 (iii) talkum i en mængde på 15 vægt-% af den samlede overtræksformulering; og

(iv) en blanding af farvestoffer i en mængde på 25 vægt-% af den samlede overtræksformulering.

15 11. Farmaceutisk sammensætning til anvendelse i en fremgangsmåde til behandling, forebyggelse eller håndtering af en sygdom eller forstyrrelse hos en patient, hvilken fremgangsmåde omfatter administrering til patienten af den farmaceutiske sammensætning ifølge krav 1 eller krav 10, hvor
20 sygdommen eller forstyrrelsen er psoriasis, arthritis, dermatitis, akne, dermatomyositis, ulcerativ colitis, Behcets sygdom, Crohns sygdom, sarkoidose, uveitis, rosacea eller lichen planus.

25 12. Farmaceutisk sammensætning til anvendelse ifølge krav 11, hvor psoriasis er typen plaque psoriasis.

13. Farmaceutisk sammensætning til anvendelse ifølge krav 11, hvor arthritis er psoriasis arthritis, rheumatoid arthritis,
30 osteoarthritis eller akut arthritis urica.

14. Farmaceutisk sammensætning til anvendelse ifølge krav 11, hvor dermatitis er atopisk dermatitis eller kontaktdermatitis.

35 15. Farmaceutisk sammensætning til anvendelse ifølge krav 11, hvor sarkoidose er kronisk kutan sarkoidose.