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DESCRIPTION

BACKGROUND

[0001] Decitabine is currently being developed as a pharmaceutical for the treatment of chronic myelogenous leukemia (CML), myelodysplastic syndrome (MDS), non-small cell lung (NSCL) cancer, sickle-cell anaemia, and acute myelogenous leukemia (AML). Decitabine possesses multiple pharmacological characteristics. Decitabine can be incorporated into DNA during the S phase of cell cycle, or can induce cell differentiation and exert haematological toxicity. Despite having a short physiological half-life, decitabine has an excellent tissue distribution.

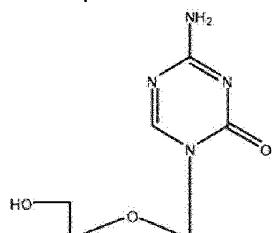
[0002] Despite its proven antileukemic effects in CML, MDS, and AML, the potential application of decitabine has been hampered by delayed and prolonged myelosuppression. Lower doses of decitabine, given over a longer period of time, have minimized myelosuppression to manageable levels without compromising its ability to suppress cancer via its hypomethylation effect. At higher doses, the associated toxicity was prohibitive. However, treatment of haematologic and solid tumours at maximally tolerated doses of decitabine has been ineffective. The cause of myelosuppression is not clear. It is plausible that since decitabine is randomly and extensively incorporated into the DNA of S phase cells, including bone marrow cells that are involved in normal haematopoiesis, the severe DNA damage due to the instability of decitabine leads to necrosis. Since incorporation of decitabine is not restricted to only the CpG-rich sequences, the DNA can break, due to the instability of decitabine, and require repair at numerous sites outside of the CpG islands.

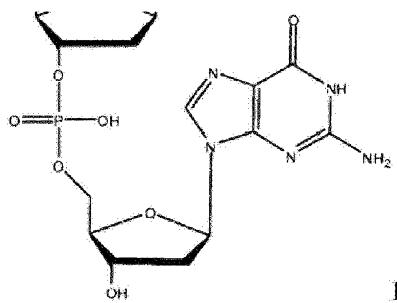
[0003] Decitabine and azacitidine are unstable in aqueous media and undergo hydrolytic degradation in aqueous media. The degradation is slowest at neutral pH. Pharmaceutically acceptable formulations of decitabine have been described previously in US 2003/0229047.

[0004] Dinucleotide compounds derived from decitabine for the development of therapies for similar indications had been described in U.S. Patent No. 7,700,567.

SUMMARY OF THE INVENTION

[0005] In a first aspect of the present invention there is provided a formulation comprising: (a) a compound of the formula:





or a pharmaceutically-acceptable salt thereof; dissolved in (b) a substantially anhydrous solvent comprising 60% to 70% propylene glycol; 20% to 30% glycerin; and 5% to 15% ethanol (w/w/w).

[0006] It has been found that the use of a substantially anhydrous solvent in the formulations of the invention produces a dramatic increase in the solubility (about 130 to about 150 mg/mL for the compound of formula 1-1). This improves subcutaneous administration, since such high concentrations lower the volumes of injection and increase the safety of the compound as less amounts of excipients are needed compared to lower concentrations of the same compound.

[0007] It has also been found that the use of substantially anhydrous solvents in the formulations of the invention exhibit increased shelf life stability (see Example 2 herein). For example, reconstituted dosage forms having a water content of 0.1% remain stable at 2-8°C for at least 12 months.

[0008] Ethanol is incorporated as a thinning agent.

[0009] In some embodiments, said solvent comprises about 65% propylene glycol; about 25% glycerin; and about 10% ethanol, for example being 65% propylene glycol; 25% glycerin; and 10% ethanol.

[0010] In some embodiments, said solvent comprises 65% to 70% propylene glycol and 25% to 30% glycerin, any balance being ethanol.

[0011] Embodiments of a pharmaceutically-acceptable salt include any salt described herein. In some embodiments, said salt is a sodium salt. The compound can be present at a concentration of 80 mg/mL to 110 mg/mL, for example about 100 mg/mL.

[0012] In some embodiments, the formulation further comprises dimethyl sulfoxide (DMSO), optionally at a DMSO: compound ratio of about 2: about 1; about 1: about 1; about 0.5: about 1; about 0.3: about 1; or 0.2 - 0.3: about 1.

[0013] In some embodiments, a formulation disclosed herein is suitable for administration by subcutaneous injection.

[0014] In another aspect, the invention provides a kit comprising: (a) a first vessel containing a compound or pharmaceutically-acceptable salt thereof as described herein; and (b) a second

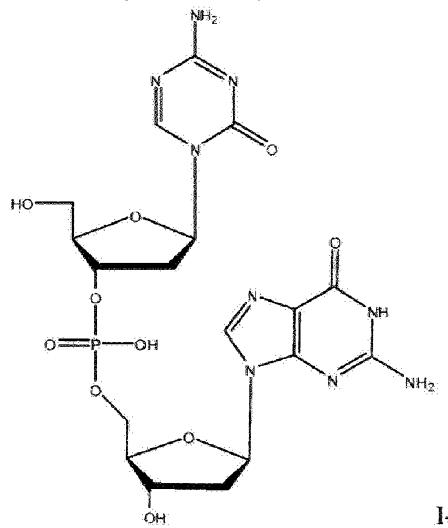
vessel containing a substantially anhydrous solvent comprising 60% to 70% propylene glycol; 20% to 30% glycerin; and 5% to 15% ethanol.

[0015] In some embodiments, the compound in the kit of the invention is present in the form of a substantially anhydrous powder, for example a lyophilized powder. The compound can be present in the first vessel in an amount of 80 mg to 110 mg, for example about 100 mg. In some embodiments, the kit further comprises instructions for administration by subcutaneous injection.

[0016] In another aspect, the invention provides a process for preparing a pharmaceutical composition, the process comprising dissolving a compound as described herein in a substantially anhydrous solvent. Examples of such a substantially anhydrous solvent include any substantially anhydrous solvent described herein.

[0017] In some embodiments, the process further comprises the steps of: dissolving said compound in DMSO to produce a solution of said compound in DMSO; and lyophilizing said solution to provide said compound as a substantially anhydrous powder.

[0018] In another aspect, the invention provides a substantially anhydrous powder consisting essentially of a compound of the formula:



or a pharmaceutically-acceptable salt thereof, and DMSO, the DMSO being present in an amount of up to about 200% w/w DMSO/compound of formula I-1. In one embodiment, the DMSO is present in an amount of up to about 100%, up to about 60%, up to about 50%, up to about 40%, or up to about 30% w/w DMSO/compound of formula I-1. In some embodiments, the DMSO is present in an amount of about 20 - about 30% w/w DMSO/compound of formula I-1. In some embodiments, the salt of the powder is a sodium salt.

[0019] In another aspect, the invention provides the formulation or kit for use in treating a cancer, myelodysplastic syndrome, leukemia or solid tumour.

[0020] The formulations, kits or uses of the invention find application in the treatment of a wide variety of diseases that are sensitive to the treatment with decitabine, including those

described herein as examples.

[0021] In some embodiments, the administration is subcutaneous administration.

[0022] Any compound described herein is suitable for use in any formulation or kit described herein.

BRIEF DESCRIPTION OF THE DRAWINGS

[0023]

FIGURE 1 illustrates the mean plasma concentrations of the compound I-1 in male and female cynomolgus monkeys given weekly subcutaneous doses of compound I-1 in a pharmacokinetic study.

FIGURE 2 illustrates the mean plasma concentrations of decitabine in male and female cynomolgus monkeys given weekly subcutaneous doses of decitabine in a pharmacokinetic study.

FIGURE 3 illustrates the decrease in LINE 1 methylation levels observed in blood samples drawn from cynomolgus monkeys on various days (D) after pretest.

FIGURE 4 illustrates the change in total related substances of the sodium salt of a compound of Formula I-1 in various DMSO and DMSO/water compositions.

DETAILED DESCRIPTION OF THE INVENTION

[0024] In current clinical treatment with decitabine, to minimize decomposition, decitabine is supplied as a lyophilized powder and reconstituted pre-administration in a cold solution containing at least 40% water (v/v), such as water for injection (WFI). This method requires refrigeration of decitabine in solution, but such storage is inconvenient and economically less desirable than storage at ambient temperatures. Due to rapid decomposition of decitabine in aqueous solution, the reconstituted decitabine solution can be infused only within hours of reconstitution. Refrigeration after reconstitution is undesirable because infusion of cold fluid can cause discomfort, pain, and subsequently, non-compliance in the subject. The inventions described herein solve these problems by providing formulations of decitabine derivatives in formulations that resist chemical decomposition and provide greater convenience and versatility in a therapeutic regimen.

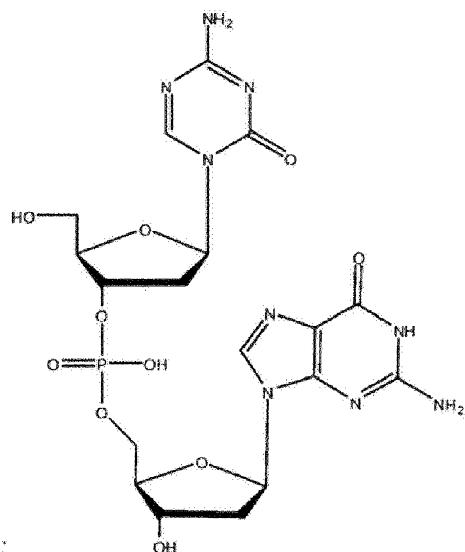
[0025] The inventions describe formulations of compounds derived from decitabine with improved chemical stability and greater ability to deliver pharmaceutically-active agent to a

subject in need or want thereof. The compounds incorporate a 5-aza-cytosine group in the form of a 5-aza-2'-deoxycytidine group (decitabine). The compounds also incorporate a guanine group in the form of a 2'- deoxyguanidine group. The 5-aza-cytosine group and the guanine group are linked by a phosphorus-containing linker.

[0026] The compounds are provided in formulations that preserve the efficacy of the compounds by providing media wherein the compounds exhibit good chemical stability.

Compounds

[0027] In some embodiments, the invention provides a formulation comprising: a) a compound of the formula:



or a pharmaceutically-acceptable salt thereof; b) a solvent comprising about 65% propylene glycol; about 25% glycerin; and about 10% ethanol, wherein the solvent is substantially anhydrous; and c) optionally, a pharmaceutically-acceptable excipient.

[0028] The invention provides kits and formulations comprising pharmaceutically-acceptable salts of the compound described herein. Pharmaceutically-acceptable salts include, for example, acid-addition salts and base-addition salts. The acid that is added to a compound to form an acid-addition salt can be an organic acid or an inorganic acid. A base that is added to a compound to form a base-addition salt can be an organic base or an inorganic base. In some embodiments, a pharmaceutically-acceptable salt is a metal salt. In some embodiments, a pharmaceutically- acceptable salt is an ammonium salt.

[0029] Acid addition salts can arise from the addition of an acid to a compound described herein. In some embodiments, the acid is organic. In some embodiments, the acid is inorganic. Examples of suitable acids include hydrochloric acid, hydrobromic acid, hydroiodic acid, nitric acid, nitrous acid, sulfuric acid, sulfurous acid, a phosphoric acid, nicotinic acid, isonicotinic acid, lactic acid, salicylic acid, 4-aminosalicylic acid, tartaric acid, ascorbic acid, gentisic acid, gluconic acid, glucaronic acid, saccaric acid, formic acid, benzoic acid, glutamic acid,

pantothenic acid, acetic acid, propionic acid, butyric acid, fumaric acid, succinic acid, citric acid, oxalic acid, maleic acid, hydroxymaleic acid, methylmaleic acid, glycolic acid, malic acid, cinnamic acid, mandelic acid, 2-phenoxybenzoic acid, 2-acetoxybenzoic acid, embonic acid, phenylacetic acid, N-cyclohexylsulfamic acid, methanesulfonic acid, ethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, 2-hydroxyethanesulfonic acid, ethane-1,2-disulfonic acid, 4-methylbenzenesulfonic acid, naphthalene-2-sulfonic acid, naphthalene-1,5-disulfonic acid, 2-phosphoglyceric acid, 3-phosphoglyceric acid, glucose-6-phosphoric acid, and an amino acid.

[0030] Examples of suitable acid addition salts include a hydrochloride salt, a hydrobromide salt, a hydroiodide salt, a nitrate salt, a nitrite salt, a sulfate salt, a sulfite salt, a phosphate salt, a hydrogen phosphate salt, a dihydrogen phosphate salt, a carbonate salt, a bicarbonate salt, a nicotinate salt, an isonicotinate salt, a lactate salt, a salicylate salt, a 4- aminosalicylate salt, a tartrate salt, an ascorbate salt, a gentisinate salt, a gluconate salt, a glucaronate salt, a saccarate salt, a formate salt, a benzoate salt, a glutamate salt, a pantothenate salt, an acetate salt, a propionate salt, a butyrate salt, a fumarate salt, a succinate salt, a citrate salt, an oxalate salt, a maleate salt, a hydroxymaleate salt, a methylmaleate salt, a glycolate salt, a malate salt, a cinnamate salt, a mandelate salt, a 2-phenoxybenzoate salt, a 2-acetoxybenzoate salt, an embonate salt, a phenylacetate salt, an N-cyclohexylsulfamate salt, a methanesulfonate salt, an ethanesulfonate salt, a benzenesulfonate salt, a p-toluenesulfonate salt, a 2-hydroxyethanesulfonate salt, an ethane-1,2-disulfonate salt, a 4-methylbenzenesulfonate salt, a naphthalene-2-sulfonate salt, a naphthalene-1,5-disulfonate salt, a 2-phosphoglycerate salt, a 3-phosphoglycerate salt, a glucose-6-phosphate salt, and an amino acid salt.

[0031] Metal salts can arise from the addition of an inorganic base to a compound described herein. The inorganic base consists of a metal cation paired with a basic counterion, such as, for example, hydroxide, carbonate, bicarbonate, or phosphate. The metal can be an alkali metal, alkaline earth metal, transition metal, or main group metal. Examples of suitable metals include lithium, sodium, potassium, cesium, cerium, magnesium, manganese, iron, calcium, strontium, cobalt, titanium, aluminum, copper, cadmium, and zinc.

[0032] Examples of suitable metal salts include a lithium salt, a sodium salt, a potassium salt, a cesium salt, a cerium salt, a magnesium salt, a manganese salt, an iron salt, a calcium salt, a strontium salt, a cobalt salt, a titanium salt, an aluminum salt, a copper salt, a cadmium salt, and a zinc salt.

[0033] Ammonium salts can arise from the addition of ammonia or an organic amine to a compound described herein. Examples of suitable organic amines include triethyl amine, diisopropyl amine, ethanol amine, diethanol amine, triethanol amine, morpholine, N-methylmorpholine, piperidine, N-methylpiperidine, N-ethylpiperidine, dibenzyl amine, piperazine, pyridine, pyrazole, pipyrazole, imidazole, pyrazine, pipyrazine, ethylenediamine, N,N'-dibenzylethylene diamine, procaine, chloroprocaine, choline, dicyclohexyl amine, and N-methylglucamine.

[0034] Examples of suitable ammonium salts include a triethyl amine salt, a diisopropyl amine salt, an ethanol amine salt, a diethanol amine salt, a triethanol amine salt, a morpholine salt, an N-methylmorpholine salt, a piperidine salt, an N-methylpiperidine salt, an N-ethylpiperidine salt, a dibenzyl amine salt, a piperazine salt, a pyridine salt, a pyrazole salt, a pipyrazole salt, an imidazole salt, a pyrazine salt, a pipyrazine salt, an ethylene diamine salt, an N,N'-dibenzylethylene diamine salt, a procaine salt, a chloroprocaine salt, a choline salt, a dicyclohexyl amine salt, and a N-methylglucamine salt.

[0035] The compounds described herein can be synthesized by methods known in the art, for example, solution phase or solid phase synthesis. For descriptions of the synthesis of compounds of the invention, and for a description of the mechanism of action of compounds of the invention, see U.S. Patent No. 7,700,567.

Formulations of the invention

[0036] Formulations described herein provide pharmaceutically-useful compositions comprising the compound described herein in a form with high solubility, low injection volumes, and good chemical stability and shelf-life. These properties provide formulations that retain a high percentage of the initial efficacy and deliver a therapeutically-effective amount of the compound even after storage at or below room temperature for extended times.

[0037] Formulations can be solutions or suspensions of a compound in a solvent or a mixture of solvents. The suitable solvents include propylene glycol, glycerin, and ethanol. The formulations are substantially anhydrous.

[0038] The solvent contains a percentage of propylene glycol on a mass basis. The percentage of propylene glycol is 60% to 70%. In some embodiments, the percentage of propylene glycol can be 60%, 65% or 70%, or about 60%, about 65% or about 70%.

[0039] The solvent contains a percentage of glycerin on a mass basis. The percentage of glycerin is 20% to 30%. In some embodiments, the percentage of glycerin can be 20%, 25%, or 30%, or about 20%, about 25%, or about 30%.

[0040] The solvent contains a percentage of ethanol on a mass basis. The percentage of ethanol is 5% to 15%. In some embodiments, the percentage of ethanol can be 5%, 6%, 7%, 8%, 9%, 10%, 11%, 12%, 13%, 14%, or 15%, about 5%, about 6%, about 7%, about 8%, about 9%, about 10%, about 11%, about 12%, about 13%, about 14%, or about 15%.

[0041] The solvent or a mixture of solvents comprises 60% to 70% propylene glycol; 20% to 30% glycerin; and 5% to 15% ethanol. In some embodiments, a solvent or a mixture of solvents comprises about 60% to about 70% propylene glycol; about

20% to about 30% glycerin; and about 5% to about 15% ethanol. In some embodiments, a solvent or a mixture of solvents consists essentially of 60% to 70% propylene glycol; 20% to 30% glycerin; and 5% to 15% ethanol. In some embodiments, a solvent or a mixture of solvents consists essentially of about 60% to about 70% propylene glycol; about 20% to about 30% glycerin; and about 5% to about 15% ethanol. In some embodiments, a solvent or a mixture of solvents is 60% to 70% propylene glycol; 20% to 30% glycerin; and 5% to 15% ethanol. In some embodiments, a solvent or a mixture of solvents is about 60% to about 70% propylene glycol; about 20% to about 30% glycerin; and about 5% to about 15% ethanol.

[0042] In some embodiments, a solvent or a mixture of solvents comprises 65% propylene glycol; 25% glycerin; and 10% ethanol. In some embodiments, a solvent or a mixture of solvents comprises about 65% propylene glycol; about 25% glycerin; and about 10% ethanol. In some embodiments, a solvent or a mixture of solvents consists essentially of 65% propylene glycol; 25% glycerin; and 10% ethanol. In some embodiments, a solvent or a mixture of solvents consists essentially of about 65% propylene glycol; about 25% glycerin; and about 10% ethanol. In some embodiments, a solvent or a mixture of solvents is 65% propylene glycol; 25% glycerin; and 10% ethanol. In some embodiments, a solvent or a mixture of solvents is about 65% propylene glycol; about 25% glycerin; and about 10% ethanol.

[0043] A formulation can be prepared, stored, transported, and handled in anhydrous or substantially-anhydrous form. A solvent can be dried prior to preparing a formulation, and a compound can be dried, for example, by lyophilization. A drying agent, or dessicant, can be used during preparation, storage, transportation, or handling to regulate water content. Examples of drying agents include silica gel, calcium sulfate, calcium chloride, calcium phosphate, sodium chloride, sodium bicarbonate, sodium sulfate, sodium phosphate, montmorillonite, molecular sieves (beads or powdered), alumina, titania, zirconia, and sodium pyrophosphate. A drying agent can contact a formulation directly, be inserted into the formulation in the form of a packet with a permeable membrane, or be stored with the formulation in a sealed environment, such as a dessicator, such that the drying agent and the formulation are simultaneously exposed to the same controlled atmosphere. A drying agent can be removed from a formulation, for example, by filtration or cannulation. Additionally, a formulation can be stored in a sealed container within a controlled atmosphere consisting essentially of, or enriched in, nitrogen or argon.

[0044] Anhydrous or substantially-anhydrous conditions benefit the shelf-life of a formulation disclosed herein at both ambient and reduced temperatures. This benefit reduces the costs associated with the storage, transportation, and spoilage of a formulation, increases the convenience of storage and handling, and avoids the need to administer cold formulations, thereby improving subject tolerance and compliance to a regimen of a formulation of the invention.

[0045] A formulation can further include a pharmaceutically-acceptable excipient. Examples of excipients include mannitol, sorbitol, lactose, dextrose, and cyclodextrins. Excipients can be added to modulate the density, rheology, uniformity, and viscosity of the formulation.

[0046] A formulation can include acidic or basic excipients to modulate the acidity or basicity of the formulation. Examples of acids suitable to increase the acidity of a formulation include hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, phosphoric acid, nitric acid, ascorbic acid, citric acid, tartaric acid, lactic acid, oxalic acid, formic acid, benzenesulphonic acid, benzoic acid, maleic acid, glutamic acid, succinic acid, aspartic acid, diatrizoic acid, and acetic acid. Examples of bases suitable to increase the basicity of a formulation include lithium hydroxide, sodium hydroxide, potassium hydroxide, sodium carbonate, sodium bicarbonate, sodium phosphate, potassium phosphate, sodium acetate, sodium benzoate, tetrabutylammonium acetate, tetrabutylammonium benzoate, and trialkyl amines. Polyfunctional excipients, such as ethylene diamine tetraacetic acid (EDTA), or a salt thereof, can also be used to modulate acidity or basicity.

[0047] The compound disclosed herein can be present in a formulation in any amount. In some embodiments, the compound is present in a concentration of 1 mg/mL to 130 mg/mL, 10 mg/mL to 130 mg/mL, 40 mg/mL to 120 mg/mL, 80 mg/mL to 110 mg/mL, about 1 mg/mL to about 130 mg/mL, about 10 mg/mL to about 130 mg/mL, about 40 mg/mL to about 120 mg/mL, or about 80 mg/mL to about 110 mg/mL. In some embodiments, the compound is present in a concentration of 10 mg/mL, 20 mg/mL, 30 mg/mL, 40 mg/mL, 50 mg/mL, 60 mg/mL, 70 mg/mL, 80 mg/mL, 90 mg/mL, 100 mg/mL, 110 mg/mL, 120 mg/mL, 130 mg/mL, 140 mg/mL, 150 mg/mL, 160 mg/mL, 170 mg/mL, 180 mg/mL, 190 mg/mL, 200 mg/mL, about 10 mg/mL, about 20 mg/mL, about 30 mg/mL, about 40 mg/mL, about 50 mg/mL, about 60 mg/mL, about 70 mg/mL, about 80 mg/mL, about 90 mg/mL, about 100 mg/mL, about 110 mg/mL, about 120 mg/mL, about 130 mg/mL, about 140 mg/mL, about 150 mg/mL, about 160 mg/mL, about 170 mg/mL, about 180 mg/mL, about 190 mg/mL, or about 200 mg/mL. In some embodiments, the compound is present in a concentration of 100 mg/mL. In some embodiments, the compound is present in a concentration of about 100 mg/mL.

[0048] A formulation can be prepared by contacting a compound described herein with a solvent or a mixture of solvents. Alternatively, the compound can be contacted with a single solvent, and other solvents can be added subsequently, as a mixture, or sequentially. When the final formulation is a solution, complete solvation can be achieved at whatever step of the process is practical for manufacturing. Optional excipients can be added to the formulation at whatever step is practical for manufacturing.

[0049] Preparation of the formulation can be optionally promoted by agitation, heating, or extension of the dissolution period. Examples of agitation include shaking, sonication, mixing, stirring, vortex, and combinations thereof.

[0050] In some embodiments, a formulation is optionally sterilized. Examples of sterilization techniques include filtration, chemical disinfection, irradiation, and heating.

[0051] Formulations of the invention are effective for maintaining the therapeutic compound and retarding decomposition during storage and handling, thereby sustaining the efficacy of

the compound and the formulation thereof.

[0052] One example of storage conditions is to store a formulation of the invention at 2-8 °C for a period of time, for example, a day, a week, a month, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12 months, about a year, or longer than a year. In some embodiments, the formulation retains about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95%, or about 100% efficacy after storage for 3 months at 2-8 °C.

[0053] One example of storage conditions is to store a formulation of the invention at 25 °C and 60% relative humidity for a period of time, for example, a day, a week, a month, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12 months, about a year, or longer than a year. In some embodiments, the formulation retains about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95%, or about 100% efficacy after storage for 3 months at 25°C and 60% relative humidity.

Dimethyl sulfoxide (DMSO) for use according to the invention

[0054] The use of DMSO as a solvent according to the invention can reduce bulk solution and fill volumes (both bulk and fill volumes can be reduced to 1/5th of those used with aqueous systems) and to remove time and temperature restrictions on scale-up.

Moreover, the use of substantially anhydrous DMSO greatly increases stability: increasing water concentration is correlated with a decrease in stability (as shown in Figure 4, which shows the % change in total related substances of the sodium salt of a compound of Formula I-1 when stored in DMSO or DMSO/water (water for injection, "WFI") at 25°C/60% RH for 24 hours).

[0055] Any source of DMSO can be used according to the invention. In some embodiments, the DMSO source is suitable for healthcare and drug delivery applications, for example, conforming to USP or Ph. Eur monographs, or manufactured under cGMP and API guidelines. Grades such as anhydrous, analytical grade, HPLC grade, or Pharma Solvent can be used according to the invention.

[0056] In some embodiments, the DMSO for use according to the invention has impurities in low levels, for example <0.2% water by KF, <0.01% non-volatile residue, and/or <0.1% of related compounds.

[0057] In some embodiments, the isosteres of DMSO can be used in place of DMSO. In some embodiments, an isostere of DMSO is one in which one or more atom(s) is(are) replaced by a cognate isotope, for example hydrogen by deuterium.

FURTHER EMBODIMENTS

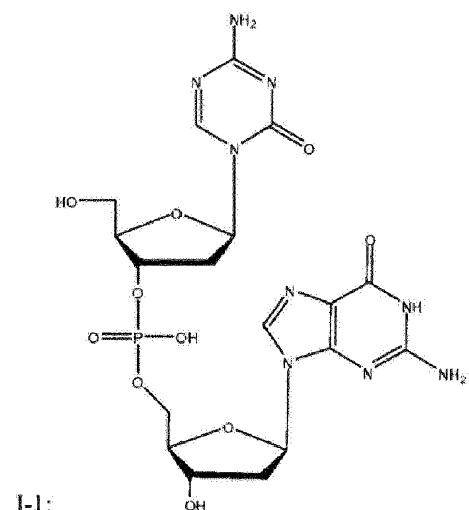
[0058] In one preferred embodiment, the formulation is substantially anhydrous.

[0059] In one preferred embodiment, the compound is present in a concentration of about 80 mg/mL to about 110 mg/mL.

[0060] In one preferred embodiment, the formulation is a solution.

[0061] In one preferred embodiment, the formulation retains about 95% efficacy after storage for 3 months at 2-8 °C, or about 68% efficacy after storage for 3 months at 25 °C and 60% relative humidity.

[0062] In one preferred embodiment, the formulation comprises: a) a compound of the formula:

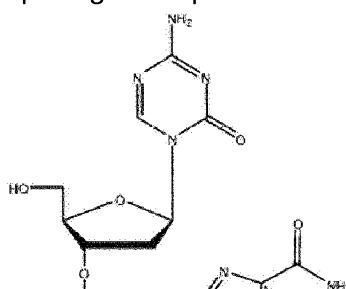


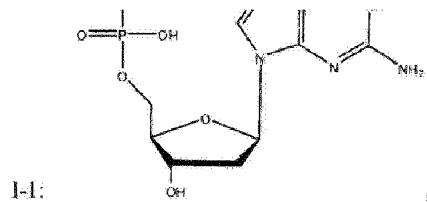
or a pharmaceutically-acceptable salt thereof; b) a solvent comprising about 65% propylene glycol; about 25% glycerin; and about 10% ethanol, wherein the solvent is substantially anhydrous; and c) optionally, a pharmaceutically-acceptable excipient.

[0063] Preferably, for this embodiment the compound exists as a sodium salt.

[0064] More preferably, the solvent is 65% propylene glycol; 25% glycerin; and 10% ethanol. More preferably, the compound is present in a concentration of about 100 mg/mL.

[0065] One embodiment of the invention relates to a formulation for use in the treatment of one or more myelodysplastic syndromes, leukemia, or solid tumours, the formulation comprising a compound having the formula:





or a pharmaceutically-acceptable salt thereof; wherein the compound is provided in a solvent comprising about 65% propylene glycol; about 25% glycerin; and about 10% ethanol, wherein the solvent is substantially anhydrous, and optionally with a pharmaceutically-acceptable excipient.

[0066] Preferably, for this embodiment, the myelodysplastic syndrome is acute myeloid leukemia (AML), acute promyelocytic leukemia (APL), acute lymphoblastic leukemia (ALL), or chronic myelogenous leukemia (CML).

[0067] Preferably, for this embodiment, the compound exists as a sodium salt.

[0068] Preferably, for this embodiment, the solvent is 65% propylene glycol; 25% glycerin; and 10% ethanol.

[0069] Preferably, for this embodiment, the compound is present in a concentration of about 100 mg/mL. Preferably, for this embodiment, the compound provided in the solvent is suitable for subcutaneous administration.

Dosing and Administration.

[0070] Doses of formulations of the invention can be administered to a subject by a method known in the art. Examples of methods of administration include subcutaneous injection, intravenous injection, and infusion. In some embodiments, a subject is in need or want of the formulation.

[0071] A dose of a formulation contains an amount that is therapeutically-effective for an indication. In some embodiments, a subject is in need or want of therapy for the indication.

[0072] A therapeutically-effective amount of a compound of the invention can be expressed as mg of the compound per kg of subject body mass. In some embodiments, a therapeutically-effective amount is 1-1,000 mg/kg, 1-500 mg/kg, 1-250 mg/kg, 1-100 mg/kg, 1-50 mg/kg, 1-25 mg/kg, or 1-10 mg/kg. In some embodiments, a therapeutically-effective amount is 5 mg/kg, 10 mg/kg, 25 mg/kg, 50 mg/kg, 75 mg/kg, 100 mg/kg, 150 mg/kg, 200 mg/kg, 250 mg/kg, 300 mg/kg, 400 mg/kg, 500 mg/kg, 600 mg/kg, 700 mg/kg, 800 mg/kg, 900 mg/kg, 1,000 mg/kg, about 5 mg/kg, about 10 mg/kg, about 25 mg/kg, about 50 mg/kg, about 75 mg/kg, about 100 mg/kg, about 150 mg/kg, about 200 mg/kg, about 250 mg/kg, about 300 mg/kg, about 400 mg/kg, about 500 mg/kg, about 600 mg/kg, about 700 mg/kg, about 800 mg/kg, about 900 mg/kg, or about 1,000 mg/kg.

[0073] In some embodiments, a therapeutically-effective amount can be administered 1-35 times per week, 1-14 times per week, or 1-7 times per week. In some embodiments, a therapeutically-effective amount can be administered 1-10 times per day, 1-5 times per day, 1 time, 2 times, or 3 times per day.

Therapeutic Uses

[0074] The pharmaceutical formulations according to the present invention can be used to treat a wide variety of diseases that are sensitive to treatment with decitabine, including those described herein.

[0075] Examples of indications that can be treated using the pharmaceutical formulations of the present invention include those involving undesirable or uncontrolled cell proliferation. Such indications include benign tumors, various types of cancers such as primary tumors and tumor metastasis, restenosis (e.g. coronary, carotid, and cerebral lesions), hematological disorders, abnormal stimulation of endothelial cells (atherosclerosis), insults to body tissue due to surgery, abnormal wound healing, abnormal angiogenesis, diseases that produce fibrosis of tissue, repetitive motion disorders, disorders of tissues that are not highly vascularized, and proliferative responses associated with organ transplants.

[0076] Generally, cells in a benign tumor retain their differentiated features and do not divide in a completely uncontrolled manner. A benign tumor is usually localized and nonmetastatic. Specific types of benign tumors that can be treated using the present invention include hemangiomas, hepatocellular adenoma, cavernous haemangioma, focal nodular hyperplasia, acoustic neuromas, neurofibroma, bile duct adenoma, bile duct cystanoma, fibroma, lipomas, leiomyomas, mesotheliomas, teratomas, myxomas, nodular regenerative hyperplasia, trachomas and pyogenic granulomas.

[0077] In a malignant tumor cells become undifferentiated, do not respond to the body's growth control signals, and multiply in an uncontrolled manner. The malignant tumor is invasive and capable of spreading to distant sites (metastasizing). Malignant tumors are generally divided into two categories: primary and secondary. Primary tumors arise directly from the tissue in which they are found. A secondary tumor, or metastasis, is a tumor which is originated elsewhere in the body but has now spread to a distant organ. The common routes for metastasis are direct growth into adjacent structures, spread through the vascular or lymphatic systems, and tracking along tissue planes and body spaces (peritoneal fluid, cerebrospinal fluid, etc.)

[0078] Specific types of cancers or malignant tumors, either primary or secondary, that can be treated using this invention include breast cancer, skin cancer, bone cancer, prostate cancer, liver cancer, lung cancer, brain cancer, cancer of the larynx, gall bladder, pancreas, rectum, parathyroid, thyroid, adrenal, neural tissue, head and neck, colon, stomach, bronchi, kidneys,

basal cell carcinoma, squamous cell carcinoma of both ulcerating and papillary type, metastatic skin carcinoma, osteo sarcoma, Ewing's sarcoma, veticulum cell sarcoma, myeloma, giant cell tumor, small-cell lung tumor, gallstones, islet cell tumor, primary brain tumor, acute and chronic lymphocytic and granulocytic tumors, hairy-cell tumor, adenoma, hyperplasia, medullary carcinoma, pheochromocytoma, mucosal neuromas, intestinal ganglioneuromas, hyperplastic corneal nerve tumor, marfanoid habitus tumor, Wilm's tumor, seminoma, ovarian tumor, leiomyomater tumor, cervical dysplasia and in situ carcinoma, neuroblastoma, retinoblastoma, soft tissue sarcoma, malignant carcinoid, topical skin lesion, mycosis fungoide, rhabdomyosarcoma, Kaposi's sarcoma, osteogenic and other sarcoma, malignant hypercalcemia, renal cell tumor, polycythermia vera, adenocarcinoma, glioblastoma multiforma, leukemias, lymphomas, malignant melanomas, epidermoid carcinomas, and other carcinomas and sarcomas.

[0079] Hematologic disorders include abnormal growth of blood cells which can lead to dysplastic changes in blood cells and hematologic malignancies such as various leukemias. Examples of hematologic disorders include but are not limited to acute myeloid leukemia, acute promyelocytic leukemia, acute lymphoblastic leukemia, chronic myelogenous leukemia, the myelodysplastic syndromes, and sickle cell anemia.

[0080] Treatment of abnormal cell proliferation due to insults to body tissue during surgery can be possible for a variety of surgical procedures, including joint surgery, bowel surgery, and cheloid scarring. Diseases that produce fibrotic tissue include emphysema.

[0081] Repetitive motion disorders that can be treated using the present invention include carpal tunnel syndrome. An example of cell proliferative disorders that can be treated using the invention is a bone tumor.

[0082] The proliferative responses associated with organ transplantation that can be treated using this invention include those proliferative responses contributing to potential organ rejections or associated complications. Specifically, these proliferative responses can occur during transplantation of the heart, lung, liver, kidney, and other body organs or organ systems.

[0083] Abnormal angiogenesis that can be treated using this invention include those abnormal angiogenesis accompanying rheumatoid arthritis, ischemic-reperfusion related brain edema and injury, cortical ischemia, ovarian hyperplasia and hypervascularity, (polycystic ovary syndrome), endometriosis, psoriasis, diabetic retinopathy, and other ocular angiogenic diseases such as retinopathy of prematurity (retrolental fibroplastic), muscular degeneration, corneal graft rejection, neurosular glaucoma and Oster Webber syndrome.

[0084] Diseases associated with abnormal angiogenesis require or induce vascular growth. For example, corneal angiogenesis involves three phases: a pre-vascular latent period, active neovascularization, and vascular maturation and regression. The identity and mechanism of various angiogenic factors, including elements of the inflammatory response, such as

leukocytes, platelets, cytokines, and eicosanoids, or unidentified plasma constituents have yet to be revealed.

[0085] In some embodiments, the pharmaceutical formulations of the present invention can be used for treating diseases associated with undesired or abnormal angiogenesis, either alone, or in combination with an anti-neoplastic agent whose activity as an anti-neoplastic agent in vivo is adversely affected by high levels of DNA methylation. The particular dosage of these agents required to inhibit angiogenesis and/or angiogenic diseases can depend on the severity of the condition, the route of administration, and related factors that can be decided by the attending physician. Generally, accepted and effective daily doses are the amount sufficient to effectively inhibit angiogenesis and/or angiogenic diseases.

[0086] Pharmaceutical formulations of the present invention can be used to treat a variety of diseases associated with undesirable angiogenesis such as retinal/choroidal neovascularization and corneal neovascularization. Examples of retinal/choroidal neovascularization include, but are not limited to, Bests diseases, myopia, optic pits, Stargarts diseases, Pagets disease, vein occlusion, artery occlusion, sickle cell anemia, sarcoid, syphilis, pseudoxanthoma elasticum carotid abostructive diseases, chronic uveitis/vitritis, mycobacterial infections, Lyme's disease, systemic lupus erythematosis, retinopathy of prematurity, Eales disease, diabetic retinopathy, macular degeneration, Bechets diseases, infections causing a retinitis or chroiditis, presumed ocular histoplasmosis, pars planitis, chronic retinal detachment, hyperviscosity syndromes, toxoplasmosis, trauma and post-laser complications, diseases associated with rubesis (neovascularization of the angle) and diseases caused by the abnormal proliferation of fibrovascular or fibrous tissue including all forms of proliferative vitreoretinopathy. Examples of corneal neovascularization include, but are not limited to, epidemic keratoconjunctivitis, Vitamin A deficiency, contact lens overwear, atopic keratitis, superior limbic keratitis, pterygium keratitis sicca, sjogrens, acne rosacea, phylectenulosis, diabetic retinopathy, retinopathy of prematurity, corneal graft rejection, Mooren ulcer, Terrien's marginal degeneration, marginal keratolysis, polyarteritis, Wegener sarcoidosis, Scleritis, periphigoid radial keratotomy, neovascular glaucoma and retrobulbar fibroplasia, syphilis, Mycobacteria infections, lipid degeneration, chemical burns, bacterial ulcers, fungal ulcers, Herpes simplex infections, Herpes zoster infections, protozoan infections and Kaposi sarcoma.

[0087] In some embodiments, the pharmaceutical formulations of the present invention can be used for treating chronic inflammatory diseases associated with abnormal angiogenesis, either alone, or in combination with an anti-neoplastic agent whose activity as an anti-neoplastic agent in vivo is adversely affected by high levels of DNA methylation. The chronic inflammation depends on continuous formation of capillary sprouts to maintain an influx of inflammatory cells. The influx and presence of the inflammatory cells produce granulomas and thus, maintains the chronic inflammatory state. Inhibition of angiogenesis using the pharmaceutical formulations of the present invention can prevent the formation of the granulomas, thereby alleviating the disease. Examples of chronic inflammatory disease include, but are not limited to, inflammatory bowel diseases such as Crohn's disease and ulcerative colitis, psoriasis,

sarcoidosis, and rheumatoid arthritis.

[0088] Inflammatory bowel diseases such as Crohn's disease and ulcerative colitis are characterized by chronic inflammation and angiogenesis at various sites in the gastrointestinal tract. For example, Crohn's disease occurs as a chronic transmural inflammatory disease that most commonly affects the distal ileum and colon but can also occur in any part of the gastrointestinal tract from the mouth to the anus and perianal area. Patients with Crohn's disease generally have chronic diarrhea associated with abdominal pain, fever, anorexia, weight loss and abdominal swelling. Ulcerative colitis is also a chronic, nonspecific, inflammatory and ulcerative disease arising in the colonic mucosa and is characterized by the presence of bloody diarrhea. These inflammatory bowel diseases are generally caused by chronic granulomatous inflammation throughout the gastrointestinal tract, involving new capillary sprouts surrounded by a cylinder of inflammatory cells. Inhibition of angiogenesis by the pharmaceutical formulations of the present invention should inhibit the formation of the sprouts and prevent the formation of granulomas. The inflammatory bowel diseases also exhibit extra intestinal manifestations, such as skin lesions. Such lesions are characterized by inflammation and angiogenesis and can occur at many sites other the gastrointestinal tract. Inhibition of angiogenesis by the pharmaceutical formulations of the present invention should reduce the influx of inflammatory cells and prevent the lesion formation.

[0089] Sarcoidosis, another chronic inflammatory disease, is characterized as a multi-system granulomatous disorder. The granulomas of this disease can form anywhere in the body and, thus, the symptoms depend on the site of the granulomas and whether the disease is active. The granulomas are created by the angiogenic capillary sprouts providing a constant supply of inflammatory cells. By using the pharmaceutical formulations of the present invention to inhibit angiogenesis, such granulomas formation can be inhibited. Psoriasis, also a chronic and recurrent inflammatory disease, is characterized by papules and plaques of various sizes. Treatment using the pharmaceutical formulations of the present invention should prevent the formation of new blood vessels necessary to maintain the characteristic lesions and provide the patient relief from the symptoms.

[0090] Rheumatoid arthritis (RA) is also a chronic inflammatory disease characterized by non-specific inflammation of the peripheral joints. It is believed that the blood vessels in the synovial lining of the joints undergo angiogenesis. In addition to forming new vascular networks, the endothelial cells release factors and reactive oxygen species that lead to pannus growth and cartilage destruction. The factors involved in angiogenesis can actively contribute to, and help maintain, the chronically inflamed state of rheumatoid arthritis. Treatment using the pharmaceutical formulations of the present invention alone or in conjunction with other anti-RA agents can prevent the formation of new blood vessels necessary to maintain the chronic inflammation and provide the RA patient relief from the symptoms.

[0091] In some embodiments, the pharmaceutical formulations of the present invention can be used for treating diseases associated with abnormal hemoglobin synthesis. Decitabine

containing formulations stimulate fetal hemoglobin synthesis because the mechanism of incorporation into DNA is associated with DNA hypomethylation. Examples of diseases associated with abnormal hemoglobin synthesis include, but are not limited to, sickle cell anemia and β -thalassemia.

[0092] In some embodiments, the pharmaceutical formulations of the present invention can be used to control intracellular gene expression. DNA methylation is associated with the control of gene expression. Specifically, methylation in or near promoters inhibit transcription while demethylation restores expression. Examples of the possible applications of the described mechanisms include therapeutically modulated growth inhibition, induction of apoptosis, and cell differentiation.

[0093] Gene activation facilitated by the pharmaceutical formulations of the present invention can induce differentiation of cells for therapeutic purposes. Cellular differentiation is induced through the mechanism of hypomethylation. Examples of morphological and functional differentiation include, but are not limited to differentiation towards formation of muscle cells, myotubes, cells of erythroid and lymphoid lineages.

[0094] Myelodysplastic syndromes (MDS) are heterogeneous clonal hematopoietic stem cell disorders associated with the presence of dysplastic changes in one or more of the hematopoietic lineages, including dysplastic changes in the myeloid, erythroid, and megakaryocytic series. These changes result in cytopenias in one or more of the three lineages. Subjects afflicted with MDS typically develop complications related to anemia, neutropenia (infections), or thrombocytopenia (bleeding). Generally, from about 10% to about 70% of subjects with MDS develop acute leukemia. Representative myelodysplastic syndromes include acute myeloid leukemia, acute promyelocytic leukemia, acute lymphoblastic leukemia, and chronic myelogenous leukemia.

[0095] Acute myeloid leukemia (AML) is the most common type of acute leukemia in adults. Several inherited genetic disorders and immunodeficiency states are associated with an increased risk of AML. These include disorders with defects in DNA stability leading to random chromosomal breakage, such as Bloom's syndrome, Fanconi's anemia, Li-Fraumeni kindreds, ataxia-telangiectasia, and X-linked agammaglobulinemia.

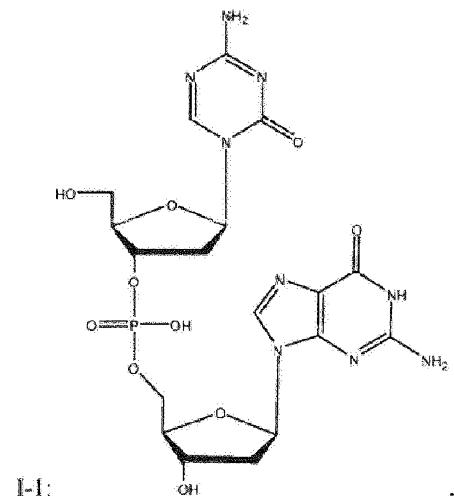
[0096] Acute promyelocytic leukemia (APML) represents a distinct subgroup of AML. This subtype is characterized by promyelocytic blasts containing the 15; 17 chromosomal translocation. This translocation leads to the generation of a fusion transcript comprising a retinoic acid receptor sequence and a promyelocytic leukemia sequence.

[0097] Acute lymphoblastic leukemia (ALL) is a heterogeneous disease with distinct clinical features displayed by various subtypes. Reoccurring cytogenetic abnormalities have been demonstrated in ALL. The most common associated cytogenetic abnormality is the 9; 22 translocation leading to development of the Philadelphia chromosome.

[0098] Chronic myelogenous leukemia (CML) is a clonal myeloproliferative disorder of a pluripotent stem cell, generally caused by ionizing radiation. CML is characterized by a specific chromosomal abnormality involving the translocation of chromosomes 9 and 22, creating the Philadelphia chromosome.

[0099] Formulations described herein can be used to provide therapy for a MDS. In some embodiments, a formulation can provide therapy for more than one MDS in a single administration.

[0100] In some embodiments, the invention provides a formulation comprising: a) a compound of the formula:



or a pharmaceutically-acceptable salt thereof; b) a solvent comprising about 65% propylene glycol; about 25% glycerin; and about 10% ethanol, wherein the solvent is substantially anhydrous; and c) optionally, a pharmaceutically-acceptable excipient.

[0101] In some embodiments, the compound exists as a sodium salt.

[0102] In some embodiments, the solvent is 65% propylene glycol; 25% glycerin; and 10% ethanol.

[0103] In some embodiments, the compound is present in a concentration of about 100 mg/mL.

EXAMPLES

EXAMPLE 1: Inhibition of DNA Methylation by Compounds of the Invention.

[0104] The demethylating activity of compounds of the invention was tested in a cell-based green fluorescent protein (GFP) assay. In the assay, a decrease in methylation resulting from exposure to a methylation inhibitor led to GFP expression, and was readily scored.

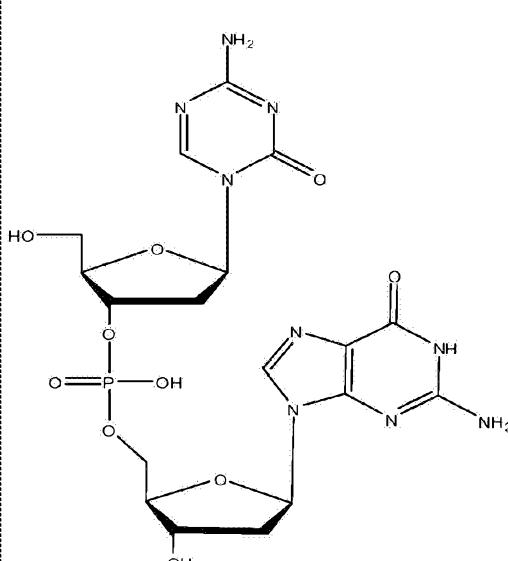
[0105] The CMV-EE210 cell line containing the epigenetically silenced GFP transgene was used to assay for reactivation of GFP expression by flow cytometry. CMV-EE210 was made by transfecting NIH 3T3 cells with the pTR-UF/UF1/UF2 plasmid, which contained pBS(+) (Stratagene, Inc.) with a cytomegalovirus (CMV) promoter driving a humanized GFP gene adapted for expression in mammalian cells. After transfection, high-level GFP expressing cells were initially selected by FACS analysis and sorting using a MoFlo cytometer (Cytomation, Inc.).

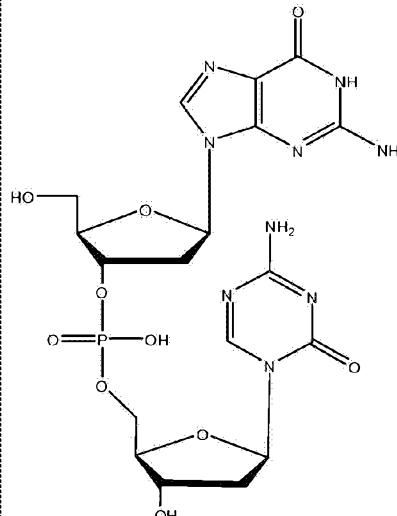
[0106] Decitabine, a potent inhibitor of mammalian DNMT1, was used as a positive control. To screen for reactivation of CMV-EE210, decitabine (1 μ M) or a test compound (30-50 μ M) was added to complete medium (phenol red free DMEM (Gibco, Life Technologies) supplemented with 10% fetal bovine serum (Hyclone)). Cells were then seeded to 30% confluence (~5000 cell/well) in a 96-well plate containing the test compounds, and grown for three days at 37 °C in 5% CO_2 .

[0107] The plates were examined under a fluorescent microscope using a 450-490 excitation filter (13 filter cube, Leica, Deerfield 111.). Wells were scored g1 positive, g2 positive, or g3 if GFP was expressed in 10%, 30%, >75% of viable cells, respectively.

[0108] Table 1 provides the results of the test for decitabine and the test compounds as DNA methylation inhibitors. GFP₅₀ is the concentration of an inhibitor at which the Green Fluorescent Protein (GFP) expression level is reduced from g3 to g1/2. Table 1 demonstrates that the tested compounds inhibited DNA methylation effectively at low concentrations, resulting in reactivation of GFP gene transcription.

TABLE 1.

Compound	GFP Expression Level	GFP ₅₀ (nM)
Decitabine	g3	500
 I-1:	g3	400

Compound	GFP Expression Level	GFP ₅₀ (nM)
I-2: 	g3	700

EXAMPLE 2: Stability of a Representative Compound in Solvent Formulations.

[0109] The stability of a compound of the invention in various formulations under various storage conditions was investigated. Stability was determined by HPLC at the designated time intervals. The results are summarized in Table 2 for formulations comprising a sodium salt of compound I-1:

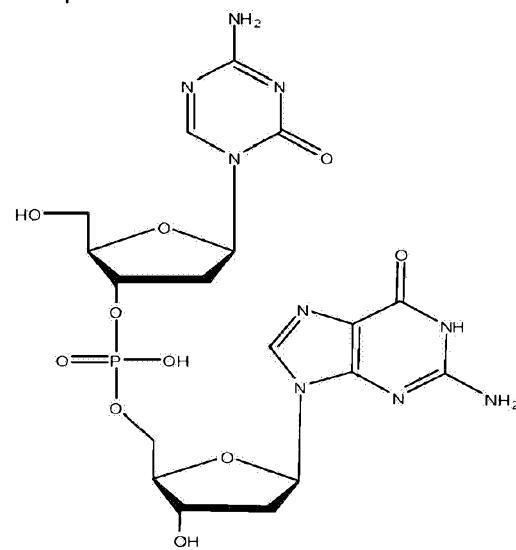


TABLE 2.

Formulation	Storage Conditions	Time Point	Percent compound detected	% decomposition per hour
		0	95.8%	

Formulation	Storage Conditions	Time Point	Percent compound detected	% decomposition per hour
water, pH 7.0	2-8 °C	5 hours	95.1%	0.14
water, pH 7.0	Room temperature	0	95.8%	
		5 hours	90.4%	1.1
DMSO / water (1:1, w/w)	25 °C / 60% relative humidity	0	93.7%	
		5 hours	90.1%	0.72
DMSO / water (3:1, w/w)	25 °C / 60% relative humidity	0	96.6%	
		24 hours	94.2%	0.10
Propylene glycol / Glycerin (70:30, v/v)	Room temperature	0	96.8%	0.021
Propylene Glycol / Glycerin / Ethanol (65:25:10, w/w/w)	2-8 °C	24 hours	96.3%	
		0	95.8%	
	25°C/60% relative humidity	3 months	95.1%	0.00032
		0	95.8%	
		3 months	67.6%	0.013

[0110] Solution of compound 1-1 in water at pH 7, the pH at which compounds of this class are most stable, led to rapid decomposition in a few hours, even at lower temperatures. Use of DMSO / water (1:1) gave slightly better results at higher temperatures. An improvement was noted in using 3:1 DMSO / water formulation. The compound was stable in anhydrous DMSO. This stability can facilitate a manufacturing process.

[0111] In regard to selection of pharmaceutically acceptable solvents for final formulation ready for administration, the anhydrous propylene glycol / glycerin system provided better stability. The final formulation was prepared by substituting small amounts of propylene glycol and glycerin with ethanol, to provide propylene glycol / glycerin / ethanol (65:25:10). This formulation provided a great improvement in the solubility and stability of the compound at both higher and lower temperatures.

[0112] Based on the experiments conducted in water, a 10-fold improvement in stability could have been expected upon changing from room temperature to colder (2-8 °C) storage conditions. However, in the propylene glycol / glycerin / ethanol (65:25:10) system, changing

from warmer to colder storage conditions provided a 40-fold improvement in stability. The combined effects of cooling plus the addition of ethanol to the propylene glycol / glycerin system provided a 66-fold improvement in stability. Such great improvements in the stability of compound 1-1 during storage could not have been expected.

[0113] The propylene glycol / glycerin / ethanol (65:25:10) system provided compound 1-1 as a solution, which was smooth, free-flowing, and suitable for passage through a 23-gauge needle without complications or clogging. The maximum solubility of the compound in this medium was determined to be about 130-150 mg/mL, which compares favorably to the aqueous solubility of 20 mg/mL. The good chemical stability taken together with the excellent solubility identified the glycol / glycerin / ethanol (65:25:10) system as a formulation for use in animal experiments.

EXAMPLE 3: Animal Studies with the Formulation of EXAMPLE 2.

[0114] The glycol / glycerin / ethanol (65:25:10) formulation of EXAMPLE 2, containing 100 mg/mL free base equivalent of the sodium salt of compound I-1 was administered to live animals. An analogous decitabine formulation was used for comparison (50 mg lyophilized decitabine powder vial reconstituted to 10 mg/mL with water for injection and administered as infusions by diluting in infusion bags).

[0115] Administration of a single dose of the formulations to monkeys (10 mg/kg) produced higher physiological concentrations of compound I-1 (C_{max} 1,130 ng/mL; AUC of 1,469 ng•hr/mL) than of decitabine (C_{max} 160 ng/mL; AUC of 340 ng•hr/mL).

[0116] In a repeat dose study, monkeys were dosed 3x weekly subcutaneously (3 mg/kg). At day 15, the systemic exposure to compound I-1 (C_{max} 181 ng/mL; AUC of 592 ng•hr/mL) was greater than that of decitabine (C_{max} 28 ng/mL; AUC of 99 ng•hr/mL). The pharmacokinetic parameters of the compounds did not vary significantly over the 22-day observation period, and minimal accumulation was detected. (FIGURES 1 and 2.) Pharmacodynamic properties (not shown) were monitored and were acceptable. Blood samples were drawn periodically to assay LINE-1 DNA methylation.

[0117] Decreases in LINE-1 DNA methylation, the indicator of biological activity, were observed, and the decrease continued until termination of the study on day 22. The observed LINE-1 methylation was significantly different ($p < 0.05$) from the methylation level observed prior to initial dosing. (FIGURE 3.)

[0118] The formulation was well-tolerated in the species tested. Three regimens were evaluated: a) once daily subcutaneous dose in rats and rabbits for 5 days; b) once weekly subcutaneous dose in rabbits and cynomolgus monkeys for 28 days as tolerated; and c) twice weekly subcutaneous dose in rats for 28 days as tolerated. Rabbits tolerated the 5-day

regimen well, up to a dose of 1.5 mg/kg/day, which is equivalent to 18 mg/kg/day in humans, and the weekly regimen up to a dose of 1.5 mg/kg/week for 3 weeks.

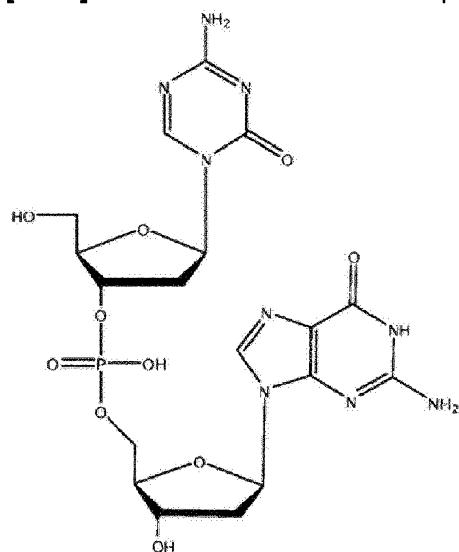
[0119] Cynomolgus monkeys tolerated the weekly regimen well, up to a dose of 3.0 mg/kg/week for 3 weeks, which is equivalent to 36 mg/kg/week. Rats tolerated much higher doses: 30 mg/kg/day over 5 days; and 20 mg/kg twice weekly for 4 weeks.

[0120] The main toxicity in all experiments was myelosuppression. However, the subcutaneous formulation tested exhibited less myelosuppression and faster recovery.

EXAMPLE 4: Preparation of a kit according to the invention

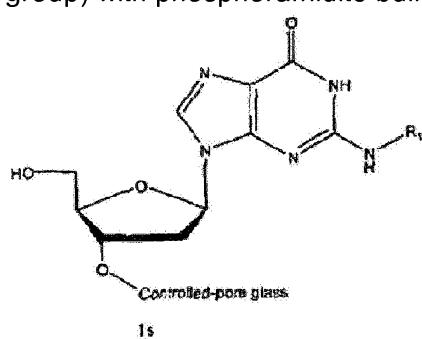
First vessel: Compound of formula 1-1 for Injection. 100 mg

[0121] The sodium salt of the compound of the formula:

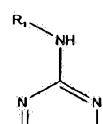


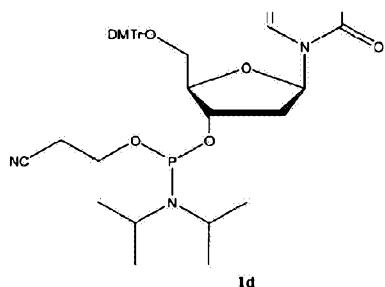
I-1

was prepared as described in US 7,700,567 by coupling **1s** (where R_1 = carbamate protective group) with phosphoramidite building block **1d**:



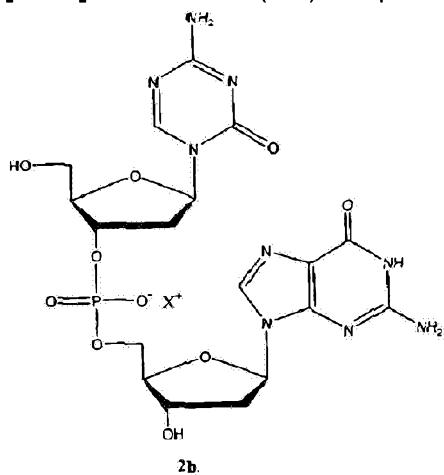
1s





[0122] A protected 2'-deoxyguanosine-linked CPG solid support **1s** (where R_1 = *tert*-butyl phenoxyacetyl) was coupled with 2-2.5 equivalents of phenoxyacetyl decitabine phosphoramidite (**1d**, where R_1 = phenoxyacetyl) in the presence of 60% of 0.3 M benzylthiotetrazole activator (in acetonitrile) for 10 minutes. The CPG solid support containing protected DpG dinucleotide was treated with 20 mL of 50 mM K_2CO_3 in methanol for 1 hour and 20 minutes. The coupled product was oxidized, the protective group was removed, and the resultant compound was washed, filtered, and purified by the ÄKTA Explorer 100 HPLC with a Gemini C18 preparative column (Phenomenex), 250x21.2 mm, 10 μ m with guard column (Phenomenex), 50x21.2mm, 10 μ m, with 50 mM triethylammonium acetate (pH 7) in MilliQ water (Mobile Phase A) and 80% acetonitrile in MilliQ water (Mobile Phase B), with 2% to 20/25% Mobile Phase **B** in column volumes.

[0123] The ESI-MS (-ve) of DpG dinucleotide **2b**:



where X^+ = triethylammonium (calculated exact mass for the neutral compound $C_{18}H_{24}N_9O_{10}P$ is 557.14), exhibited m/z 556.1 $[M-H]^-$ and 1113.1 for $[2M-H]^-$ (see mass spectrum in Figure 31 of US 7700567).

[0124] The sodium salt of the compound of formula I-1, i.e. DpG dinucleotide **2b**, where X^+ = sodium, was obtained by re-dissolving the triethylammonium salt in 4 mL water, 0.2 mL 2M $NaClO_4$ solution. When 36 mL acetone was added, the dinucleotide precipitated. The solution was kept at -20°C for several hours and centrifugated at 4000 rpm for 20 minutes. The supernatant was discarded and the solid was washed with 30 mL acetone followed by an additional centrifugation at 4000 rpm for 20 minutes. The precipitate, which was dissolved in

water and freeze dried, exhibited m/z 556.0 [M-H]⁻ (see mass spectrum in Figure 36 of US 7700567).

Compounding and filling of bulk formulation

[0125]

1. 1. Based on the assay value of the lot of the sodium salt of the compound of formula I-1, needed quantities the salt and DMSO were calculated and weighed appropriately for the intended batch scale.
2. 2. The sodium salt of the compound of formula I-1 was dissolved in DMSO utilizing an overhead mixer in an appropriately sized stainless steel (SS) vessel.
3. 3. Upon complete solubilization of the drug in DMSO, samples of the bulk solution were tested using a UV or HPLC in-process method to determine that the amount of the sodium salt of the compound of formula I-1 was within 95-105% of the target concentration.
4. 4. Bulk solution was filtered through a series of two pre-sterilized 0.2 micron sterilizing filters that were DMSO compatible, and collected into a 2L SS surge vessel.
5. 5. Filtration rate was continuously adjusted by visual monitoring of quantity available for filling in the surge vessel.
6. 6. One gram of the filtered bulk solution was filled into each of the 5 cc depyrogenated, clear glass vials and the operation was continued with until all of the filtered bulk solution was filled.
7. 7. Each vial was automatically and partially stoppered on the fill line with a fluoropolymer coated, chlorobutyl rubber lyo stopper that was pre-sterilized.
8. 8. Product vials were transferred to lyophilizer under aseptic transfer conditions for initiation of lyophilization cycle.

Lyophilization and capping of vials

[0126]

1. Vials were lyophilized using the cycle parameters as below.

Freezing	Primary/Secondary Drying					Final Set point (stoppering conditions)
Temperature	-40° C	-5° C	10° C	30° C	60° C	25° C
Ramp time (min)	133	117	50	67	100	-

Freezing	Primary/Secondary Drying					Final Set point (stoppering conditions)
Time (min.)	360	1440	1440	1440	1440	hold
Vacuum (mTorr)	- (note: 100 mT for evacuation at -50°C)	100	100	50	50	50 mT before back fill

- Upon completion of the lyophilization cycle, the lyophilizer was back filled with nitrogen, and the vials were completely and automatically stoppered.
- Vials were aseptically transferred to an isolator where each of the vials was automatically capped with a blue aluminum flip-off cap.
- Vials were visually inspected before proceeding with sampling for release testing, and the labeling and packaging operation. Vials were kept at 2-8°C until ready.

Labeling and Packaging

[0127] Each vial was labeled per approved content, and packaged individually into a heat-sealed aluminum foil pouch with a desiccant under vacuum. The foil pouch was labeled outside with the same label as was used for the product vial. Labeled and packaged vials were stored at 2-8°C until further distribution.

Residual DMSO

[0128] Four batches of the same scale of 3000 vials/batch were prepared using the same process as described above. DMSO was consistently removed to the following residual levels to yield a solid white powder, demonstrating that lyophilization of the sodium salt of the compound of formula I-1 out of DMSO as described above yielded a safe and chemically stable sodium salt of the compound of formula I-1 as a powder:

#	DMSO in mg/vial
Batch 1	25
Batch 2	28
Batch 3	27
Batch 4	29

Second vessel: Diluent for reconstitution of the sodium salt of the compound of formula I-1, 3 mL

Compounding and filling of bulk formulation**[0129]**

1. Calculated quantities (see table below) of propylene glycol, ethanol, and glycerin in the aforementioned order were added into an appropriately sized stainless steel vessel equipped with an overhead mixer.

	% of each ingredient	Grade	Function
Propylene glycol	65	NF, PhEur	Solvent
Glycerin	25	NF, PhEur	Solvent
Alcohol/Ethanol	10	USP, PhEur	Thinning agent

2. Intermittent mixing during addition of components was followed by at least 30 minutes of mixing to yield a well-mixed solution.
3. Bulk solution was filtered through a series of two pre-sterilized 0.2 micron compatible sterilizing filters, and collected into a 2L SS surge vessel.
4. Filtration rate was adjusted by visual monitoring of quantity available for filling in the surge vessel.
5. At least 3.15 g, equivalent to 3.0 mL, of the filtered bulk solution was filled into each of the 5 cc depyrogenated, clear glass vials followed by automatic stoppering using fluoropolymer coated chlorobutyl rubber closures.
6. Stoppered vials were capped with sterilized white aluminum flip-off caps.
7. Vials were visually inspected prior to sampling for the release testing and labeling operation and were stored at 2-30°C until ready.

Labeling and Packaging

[0130] Each diluent vial was labeled per approved content. Labeled vials were stored at 2-30°C until further distribution.

REFERENCES CITED IN THE DESCRIPTION

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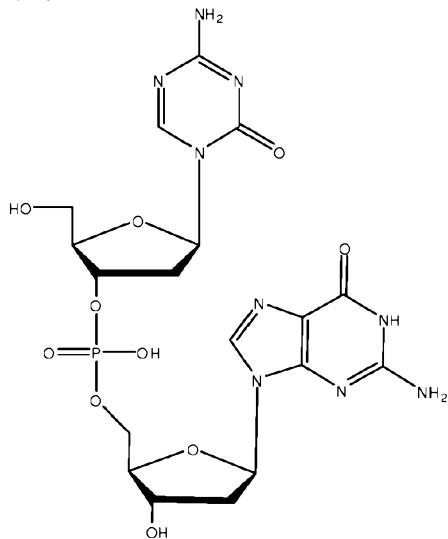
Patent documents cited in the description

- US20030229047A [0003]
- US7700567B [0004] [0035] [0121] [0123] [0124]

Patentkrav

1. Formulering, der omfatter:

(a) en forbindelse med formlen:



I-1,

5

eller et farmaceutisk acceptabelt salt deraf; opløst i

(b) et i det væsentlige vandfrit opløsningsmiddel, der omfatter 60 % til 70 % propylenglycol; 20 % til 30 % glycerin og 5 % til 15 % ethanol (w/w/w).

10

2. Formulering ifølge 1, hvor opløsningsmidlet er 65 % propylenglycol; 25 % glycerin og 10 % ethanol (w/w/w).

3. Formulering ifølge et hvilket som helst af de foregående 15 krav, hvor saltet er et natriumsalt.

4. Formulering ifølge et hvilket som helst af de foregående krav, hvor forbindelsen er til stede i en koncentration på fra 80 mg/ml til 110 mg/ml, eventuelt på omkring 100 mg/ml.

20

5. Formulering ifølge et hvilket som helst af de foregående krav, som yderligere omfatter DMSO, eventuelt i et DMSO:forbindelse-forhold på omkring 2:omkring 1; omkring 1:omkring 1; omkring 0,5:omkring 1; omkring 0,3:omkring 1 eller 0,2 - 0,3:omkring 1.

25

6. Formulering ifølge et hvilket som helst af de foregående krav, som er egnet til indgivelse ved subkutan injektion.

7. Kit, der omfatter:

(a) en første beholder, der indeholder en forbindelse som defineret i krav 1 eller krav 3; og

5 (b) en anden beholder, der indeholder et i det væsentlige vandfrit opløsningsmiddel som defineret i krav 1 eller 2.

8. Kit ifølge krav 7, hvor forbindelsen er lyofiliseret.

10 9. Kit ifølge et hvilket som helst af kravene 7-8, hvor den første beholder indeholder 80 mg til 110 mg af forbindelsen.

10. Kit ifølge et hvilket som helst af kravene 7-9, hvor den første beholder indeholder omkring 100 mg af forbindelsen.

15

11. Kit ifølge et hvilket som helst af kravene 7-10, der yderligere omfatter instruktioner i indgivelse ved subkutan injektion.

20 12. Kit ifølge et hvilket som helst af kravene 7-11, hvor forbindelsen er i form af et i det væsentlige vandfrit pulver.

13. Kit ifølge krav 12, hvor det i det væsentlige vandfrie pulver hovedsagelig består af forbindelsen og DMSO, idet DMSO 25 er til stede i en mængde på op til 200 % w/w.

14. Kit ifølge krav 13, hvor DMSO er til stede i en mængde på 20-30% w/w DMSO/forbindelse.

30 15. Fremgangsmåde til fremstilling af en farmaceutisk sammensætning, hvilken fremgangsmåde omfatter at op löse en forbindelse som defineret i krav 1 eller krav 3 i et i det væsentlige vandfrit opløsningsmiddel som defineret i krav 1 eller 2.

35

16. Fremgangsmåde ifølge krav 15, der yderligere omfatter trinnene:

(a) at op löse forbindelsen i DMSO for at opnå en opløsning af

forbindelsen i DMSO; og

(b) at lyofilisere opløsningen fra trin (a) for at tilvejebringe forbindelsen som et i det væsentlige vandfrit pulver.

5

17. Formulering ifølge et hvilket som helst af kravene 1-6 eller kit ifølge et hvilket som helst af kravene 7-14 til anvendelse ved behandling af cancer, myelodysplastisk syndrom, leukæmi eller en solid tumor.

10

18. Formulering ifølge et hvilket som helst af kravene 1-6 eller kit ifølge et hvilket som helst af kravene 7-14 til anvendelse ved behandling af brystcancer, hudcancer, knoglecancer, prostatacancer, levercancer, lungecancer,

15 hjerne cancer, cancer af strubehovedet, galdeblæren, pankreas, rectum, parathyreoidea, thyreoidea, binyren, nervevæv, hoved og hals, colon, mave, bronkier, nyrer, basalcellekarcinom, pladecellekarcinom af både den ulcererende og den papilære type, metastatisk hudkarcinom, osteosarkom, Ewings sarkom,

20 veticulumcellesarkom, myelom, kæmpecelletumor, småcellet lungetumor, galdesten, ø-celletumor, primær hjernetumor, akutte og kroniske lymfatiske og granulocytiske tumorer, hårcelletumor, adenom, hyperplasi, medullært karcinom, fæokromocytom, slimhindeneuromer, intestinale ganglioneuromer,

25 hyperplastisk hornhindenervetumor, marfanoid habitus-tumor, Wilms tumor, seminom, ovarietumor, leiomyomater-tumor, cervikal dysplasi og in situ-karcinom, neuroblastom, retinoblastom, blødvævssarkom, malignt carcinoid, topisk hudlæsion, mycosis fungoides, rhabdomyosarkom, Kaposis sarkom,

30 osteogene og andre sarkomer, malign hyperkalcæmi, renalcelletumor, polycythermia vera, adenocarkarcinom, glioblastoma multiforme, leukæmier, lymfomer, maligne melanomer, epidermoide karcinomer eller andre karcinomer og sarkomer.

35

19. Formulering eller kit til anvendelse ifølge krav 17, hvor leukæmien er valgt blandt akut myeloid leukæmi (AML), akut promyelocytisk leukæmi (APML), akut lymfoblastisk leukæmi

(ALL) og kronisk myelogen leukæmi (CML).

20. Formulering ifølge et hvilket som helst af kravene 1-6, eller kit ifølge et hvilket som helst af kravene 7-14, til 5 anvendelse ved behandling af en lidelse forbundet med unormal hæmoglobinsyntese.

21. Formulering eller kit til anvendelse ifølge et hvilket som helst af kravene 17 til 19, hvor formuleringen er til 10 subkutan indgivelse.

DRAWINGS

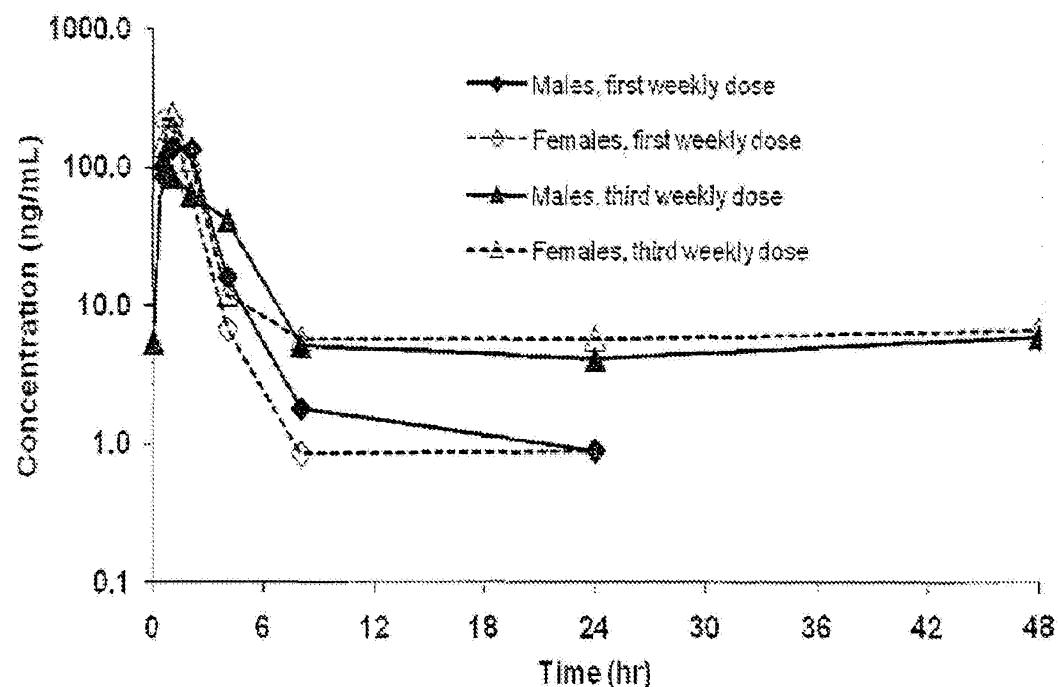


FIGURE 1

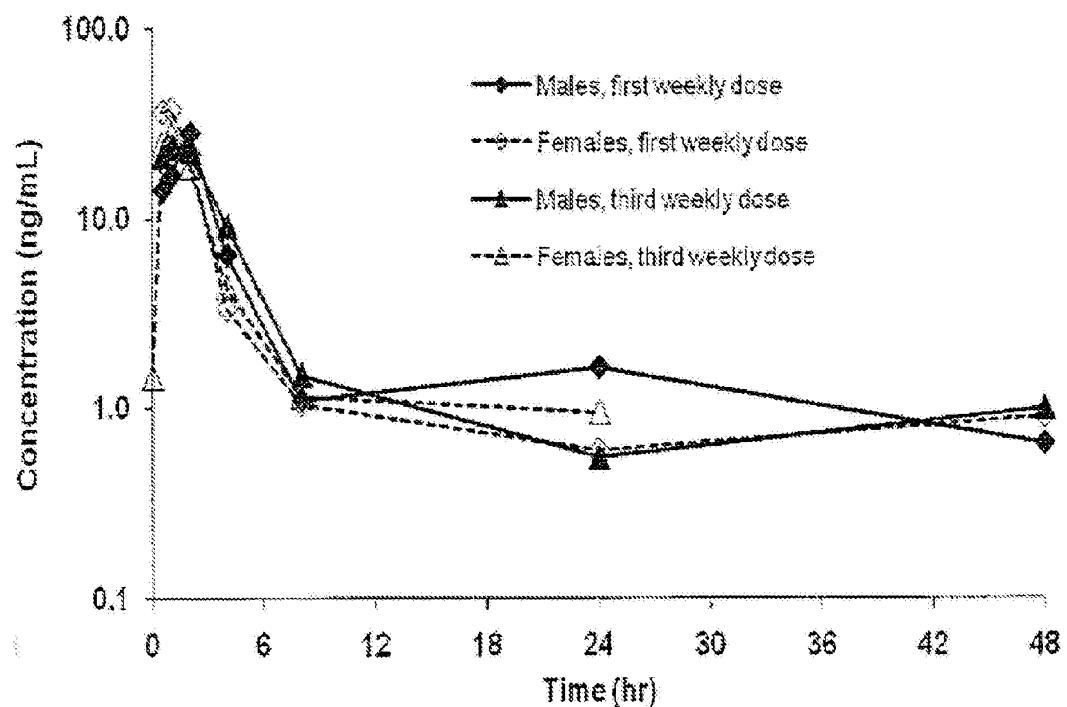


FIGURE 2

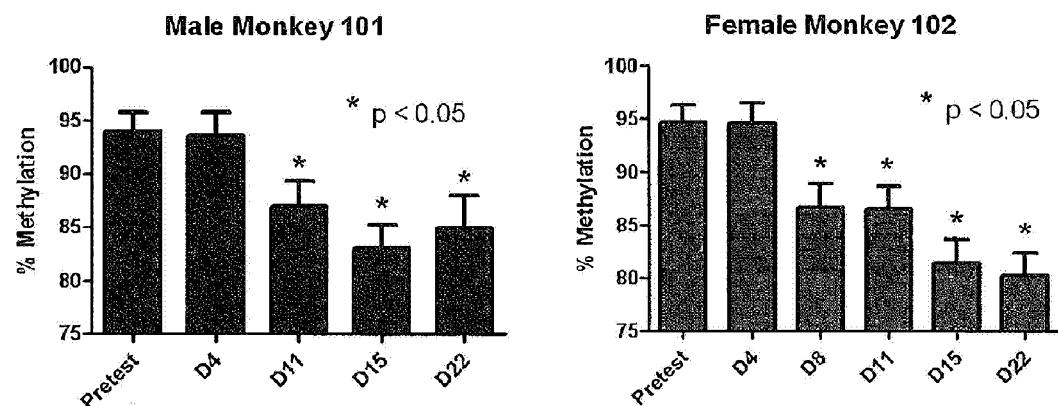


FIGURE 3

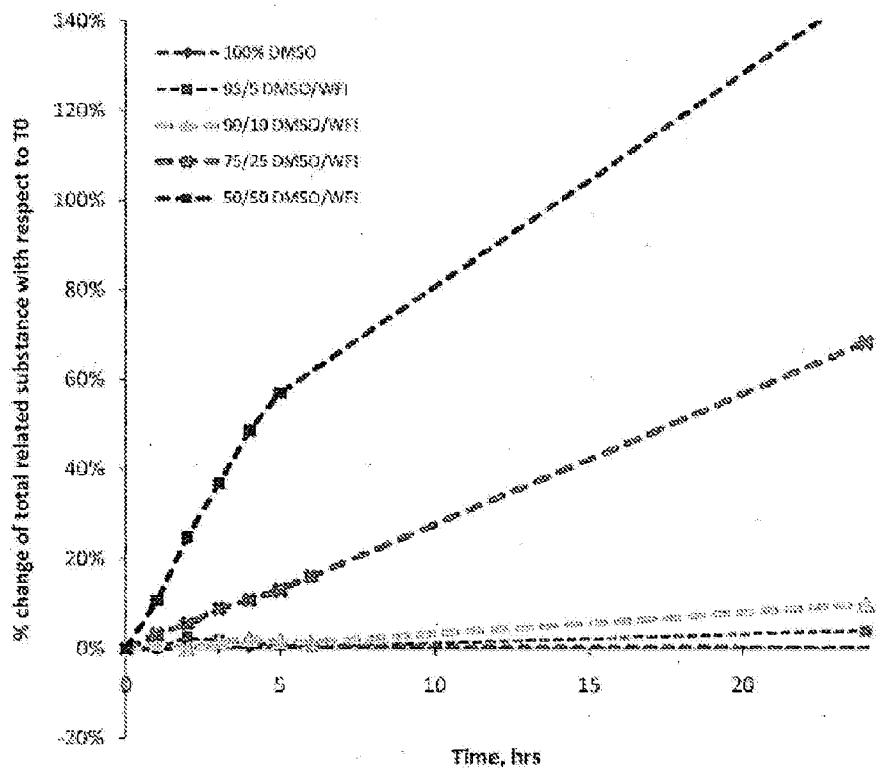


FIGURE 4