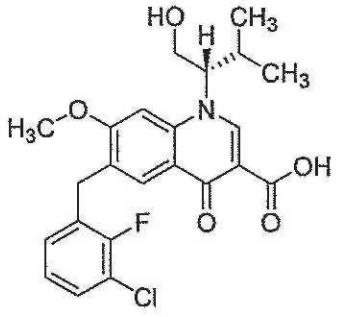
TABLETS FOR COMBINATION THERAPY

Priority of Invention

This application claims priority to United States Provisional Patent Application Number 61/150,655 filed 06 February 2009; and to United States Provisional Patent Application Number 61/150,652 filed 06 February 2009. The entire content of each of these applications is hereby incorporated herein by reference.

Background of the Invention

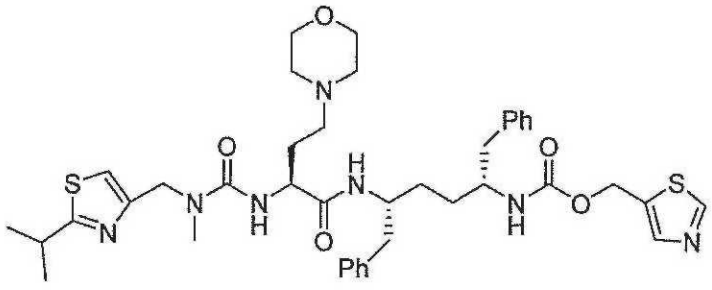
US Patent 7,176,220 discloses compounds that are reported to be useful as integrase inhibitors. One particular compound is the compound of Formula I (i.e. elvitegravir).



I

International Patent Application Publication Number WO 2005/113508 provides certain specific crystalline forms of the compound of Formula I. The specific crystalline forms are reported to have superior physical and chemical stability compared to other physical forms of the compound.

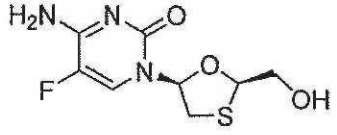
International Patent Application Publication Number WO 2008/010921 describes compounds that improve the pharmacokinetics of a co-administered drug by inhibiting cytochrome P450 monooxygenase. One such inhibitor is the compound of Formula II.



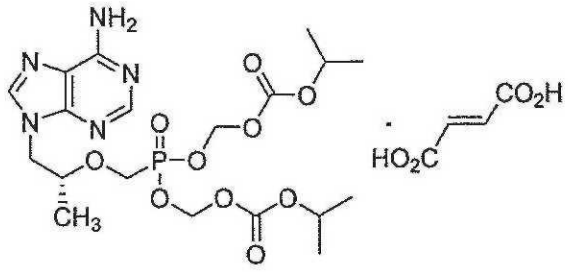
II

WO 2008/010921 also describes pharmaceutical compositions comprising the compounds and provides for compositions that include at least one additional therapeutic agent. On pages 170 and 174 WO 2008/010921 further describes combinations of the disclosed compounds (e.g. a

compound of Formula II). Included among the listing of different agents is the compound of Formula I (i.e. elvitegravir), the compound of Formula III (i.e. emtricitabine) and the salt of Formula IV (i.e. tenofovir disoproxil fumarate). A discussion regarding formulations of the compounds can also be found at pages 159-168.



III



IV

International Patent Application Publication Number WO 2008/103949 also discloses compounds that are reported to be useful to modify the pharmacokinetics of a co-administered drug by inhibiting cytochrome P450 monooxygenase. Page 199 of WO 2008/103949 describes the specific combination of the compounds of Formula I, Formula II, Formula III and Formula IV. Formulations are discussed on pages 181-190.

Despite these reports there currently is a need for improved solid dose forms (e.g. tablets) for the delivery of multi-agent compositions. There is also a need for improved solid dose forms comprising the compounds of Formulas I, II, III and IV that have enhanced or acceptable chemical stability, physical stability, release characteristics, or dosage uniformity. There is also a need for improved processes to prepare solid dose forms of compositions comprising the compounds of Formulas I, II, III and IV.

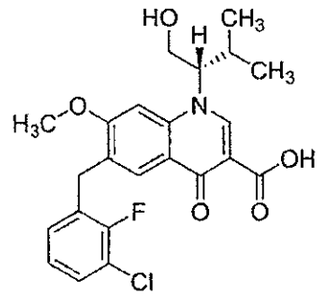
Summary of the Invention

The present invention provides solid dose forms (e.g. tablets) comprising the compounds of Formula I, Formula II, Formula III and the salt of Formula IV and methods to prepare the solid dose forms. Applicant has discovered that tablets prepared in a bilayer manner have superior properties to alternatively structured tablets. This discovery represents an advance in the development of combination therapy for the treatment of viral infections such as HIV.

Accordingly, in one embodiment the invention provides a tablet comprising a first layer and a second layer wherein;

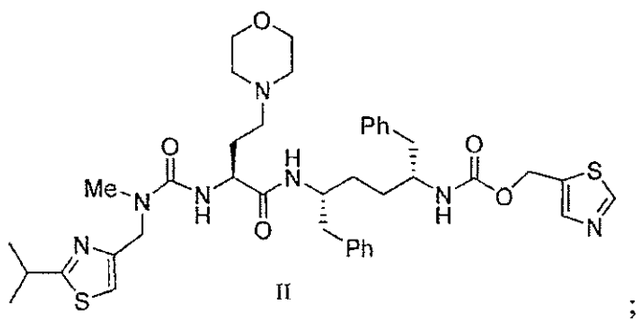
a) the first layer comprises:

a compound of Formula I:



I ; and

a compound of Formula II:

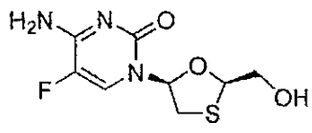


II ;

and optionally a pharmaceutically acceptable carrier; and

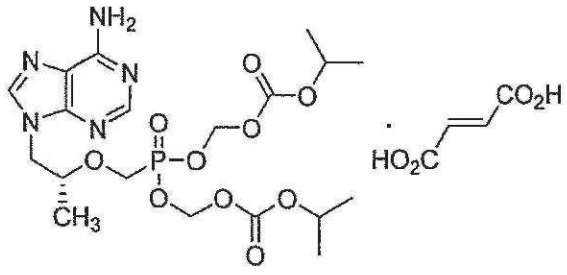
b) the second layer comprises:

a compound of Formula III:



III ; and

a salt of Formula IV:



IV ;

and optionally a pharmaceutically acceptable carrier.

In one embodiment the invention provides a pharmaceutical composition that comprises 150 mg ± 10% of the compound of Formula I; 150 mg ± 10% of the compound of Formula II; 200 mg ± 10% of the compound of Formula III; and 300 mg ± 10% of the compound of Formula IV.

In one embodiment the invention provides a pharmaceutical composition that comprises 150 mg ± 5% of the compound of Formula I; 150 mg ± 5% of the compound of Formula II; 200 mg ± 5% of the compound of Formula III; and 300 mg ± 5% of the compound of Formula IV.

In one embodiment the invention provides a pharmaceutical composition that comprises 150 mg ± 2% of the compound of Formula I; 150 mg ± 2% of the compound of Formula II; 200 mg ± 2% of the compound of Formula III; and 300 mg ± 2% of the compound of Formula IV.

In one embodiment the invention provides a pharmaceutical composition that consists of 150 mg ± 5% of the compound of Formula I; 150 mg ± 5% of the compound of Formula II; 200 mg ± 5% of the compound of Formula III; and 300 mg ± 5% of the compound of Formula IV as pharmaceutically active ingredients; and one or more pharmaceutically acceptable carriers.

In one embodiment the invention provides a pharmaceutical composition that consists of 150 mg ± 2% of the compound of Formula I; 150 mg ± 2% of the compound of Formula II; 200 mg ± 2% of the compound of Formula III; and 300 mg ± 2% of the compound of Formula IV as pharmaceutically active ingredients; and one or more pharmaceutically acceptable carriers.

In one embodiment the invention provides a tablet that consists of the following:

Components	Compound of Formula II 150 mg Formulation	
	% w/w	mg/tablet
Compound of Formula III	13.9	200.0
Salt of Formula IV	20.9	300.0
Compound of Formula I	10.4	150.0
Compound of Formula II	10.4	150.0
Colloidal Silicon Dioxide	12.0	172.5
Lactose Monohydrate	0.8	10.9
Microcrystalline Cellulose	20.8	299.5
Hydroxypropyl Cellulose	0.5	7.5
Hydroxypropyl Cellulose	0.6	9.0
Sodium Lauryl Sulfate	0.8	11.3
Croscarmellose Sodium	7.3	104.3
Magnesium Stearate	1.6	22.4
Total	100	1437.

The invention also provides methods for preparing tablets of the invention that are described herein, as well as intermediate compositions and articles that are useful for preparing compositions of the invention. Accordingly, in one embodiment the invention provides a method for preparing a tablet of the invention comprising: compressing a composition that comprises the compound of formula I and the compound of formula II to provide a first pressed layer; adding the compound of formula III and the salt of formula IV to the first pressed layer; and compressing to provide the tablet of the invention. In one embodiment of the invention, the composition that comprises the compound of formula I and the compound of formula II is compressed to provide a soft layer; a mixture that comprises the compound of formula III and the salt of formula IV is added to the soft layer to provide a final combination; and the final

combination is compressed to provide the tablet of the invention. In one embodiment, the methods of the invention can further comprise coating the tablet.

The invention also provides the methods illustrated in Figures 2 and 3 and in the Examples herein, which are useful for preparing intermediate compositions and tablets of the invention.

Brief Description of the Figures

FIG. 1. Illustrates a tablet of the invention.

FIG. 2. Illustrates the preparation of compositions of the invention.

FIG. 3. Illustrates the preparation of compositions of the invention.

Detailed Description

Specific values listed below for ranges and terms are for illustration only; they do not exclude other values.

Compound Amounts and Compound Weight Ratios

The tablets of the invention comprise compounds of Formula I, Formula II, Formula III and the salt of Formula IV. The weights and ratios described herein relate to the combination therapy as described. It is to be understood that the amounts of the various compounds can vary while remaining within the scope of the invention. In one embodiment the amount of the compound of Formula I in a tablet of the invention is 150 mg ± 10%. In another embodiment the amount of the compound of Formula I in a tablet of the invention is 150 mg ± 5%. In another embodiment the amount of the compound of Formula I in a tablet of the invention is 150 mg ± 2%. In one embodiment the amount of the compound of Formula II in a tablet of the invention is 150 mg ± 10%. In another embodiment the amount of the compound of Formula II in a tablet of the invention is 150 mg ± 5%. In another embodiment the amount of the compound of Formula II in a tablet of the invention is 150 mg ± 2%. In one embodiment of the amount of the compound of Formula III in a tablet of the invention is 200 mg ± 10%. In another embodiment the amount of the compound of Formula III in a tablet of the invention is 200 mg ± 5%. In another embodiment the amount of the compound of Formula III in a tablet of the invention is 200 mg ± 2%. In one embodiment the amount of the salt of Formula IV in a tablet of the invention is 300 mg ±10%. In another embodiment the amount of the salt of Formula IV in a tablet of the invention is 300 mg ± 5%. In another embodiment the amount of the salt of Formula IV in a tablet of the invention is 300 mg ± 2%.

It is also to be understood that various weight ratios of the compounds to one another

within the combination therapy as described can vary while remaining within the scope of the invention. In one embodiment the weight ratio of the compound of Formula I to the compound of Formula II in a tablet of the invention is 1 ± 0.8. In another embodiment the weight ratio of the compound of Formula I to the compound of Formula II in a tablet of the invention is 1 ± 0.5. In another embodiment the weight ratio of the compound of Formula I to the compound of Formula II in a tablet of the invention is 1 ± 0.3. In another embodiment the weight ratio of the compound of Formula I to the compound of Formula II in a tablet of the invention is 1 ± 0.1.

Pharmaceutical Formulations

The pharmaceutical compositions of the invention may be formulated with conventional carriers. While it is possible for the active ingredients of the composition to be administered alone it may be preferable to present them as pharmaceutical formulations. As described herein the pharmaceutical formulations of the invention may comprise compounds of Formulas I, II, III and IV together with one or more acceptable carriers. The carrier(s) should be "acceptable" in the sense of being compatible with the other ingredients of the formulation and physiologically innocuous to the recipient thereof. Accordingly, in one embodiment, the application provides for pharmaceutical compositions comprising the compounds of Formulas I, II, III and IV and a pharmaceutically acceptable carrier

As used herein the term carrier includes excipients, glidants, fillers, binders, lubricant, diluents, preservatives, surface active agents, dispersing agents and the like. The term carrier also includes agents such sweetening agents, flavoring agents, coloring agents and preserving agents. Furthermore, these terms include the values mentioned herein as well as values in accord with ordinary practice. Tablets will generally contain excipients, glidants, fillers, binders and the like. All formulations can optionally contain excipients such as those set forth in the Handbook of Pharmaceutical Excipients (APhA Publications, Washington, DC), herein incorporated by reference in its entirety. Excipients include ascorbic acid and other antioxidants, chelating agents such as EDTA, carbohydrates such as dextrin, hydroxyalkylcellulose, hydroxyalkylmethylcellulose, stearic acid and the like. The pH of the formulations ranges from about 3 to about 11, but is ordinarily about 7 to 10.

The pharmaceutical formulations include those suitable for the administration routes discussed herein. The pharmaceutical formulations may conveniently be presented in unit dosage form and may be prepared by any of the methods well known in the art of pharmacy. Techniques and formulations generally are found in Remington's Pharmaceutical Sciences

(Mack Publishing Co., Easton, PA), herein incorporated by reference in its entirety. Such methods include the step of bringing into association the active ingredient(s) with the carrier which constitutes one or more accessory ingredients. In general the formulations can be prepared by uniformly and intimately bringing into association the active ingredient(s) with liquid carriers or finely divided solid carriers or both, and then, if necessary, shaping the product (e.g. as a unit dosage form).

A particular carrier that can be used in combination with the compound of Formula II is a silica particle. Typically, silica particles comprise a granular hydrophilic fumed silica that has a mean particle diameter of 10 to 120 micron and a BET surface area of 40 to 400 m^2/g (determined according to DIN 66 131 with nitrogen). The silica particles also typically have a pore volume of about 0.5 to 2.5 mL/g, wherein less than about 5% of the overall pore volume has a pore diameter of less than about 5 nm, the remainder being mesopores and macropores. Additionally, the silica particles typically have a pH in the range of about 3.6 to about 8.5 and a tapped density of about 220 to about 700 g/L.

A specific silica material (particle) that is particularly useful in the compositions of the invention is AEROPERL® 300 (fumed silica), which is available from Evonik Degussa AG, Dusseldorf, Germany. However, other materials having physical and chemical properties similar to the silica materials described herein can also be used, e.g. calcium silicate (such as Zeopharm), or magnesium aluminometasilicate (such as Neusilin). Silica particles that have a mean grain diameter of 10 to about 120 micron are useful. Silica particles that have a mean grain diameter of 20-40 micron are also useful. Silica particles that have a BET surface area of about 40 to 400 m^2/g are useful. Silica particles that have a BET surface area of at least 150 m^2/g, or at least 200 m^2/g, or at least 250 m^2/g or at least 275 m^2/g are also useful.

In one embodiment, the compound of Formula II is associated (i.e. coated in the pores and on the surface) with the silica particles prior to combining with the other components of the compositions of the invention. In one embodiment of the invention the weight percentage of the compound of Formula II to the silica particles is 20% ± 15%. In one embodiment of the invention the weight percentage of the compound of Formula II to the silica particles is 50% ± 10%. In one embodiment of the invention the weight percentage of the compound of Formula II to the silica particles is 45% ± 15%. In one embodiment of the invention the weight percentage of the compound of Formula II to the silica particles is 100% ± 20%. In one embodiment of the

invention the weight percentage of the compound of Formula II to the silica particles is 85% ± 15%.

Loading

The compound of Formula II can be loaded on the silica particles using any suitable method. For example the compound of Formula II can be loaded on the silica particles by:

a) spraying a solution of the compound (e.g. a solution of the compound in an alcohol solvent such as ethanol) onto the silica particles;

b) combining the compound of Formula II, a suitable solvent (e.g. a volatile solvent such as dichloromethane), and the silica particles; evaporating the solvent; and isolating the resulting solid material; or

c) combining the compound of Formula I and a suitable volatile solvent (e.g. a halogenated hydrocarbon such as dichloromethane), and the silica particles; adding an antisolvent (e.g. a highly non-polar solvent such as hexanes or heptane) and isolating the resulting solid material (as illustrated in Example 4).

The compound of Formula II can be combined with a suitable solvent and a plurality of silica particles to provide a mixture. Optionally, the compound of Formula II can be combined with the suitable solvent with concurrent mixing. Typically, the weight percentage of the compound of Formula II to the silica particles prior to combining is about is 50% ± 10%. In one embodiment of the invention the weight percentage of the compound of Formula II to the silica particles prior to combining is about 20% ± 10%. In another embodiment of the invention the weight percentage of the compound of Formula II to the silica particles prior to combining is about 30% ± 10%. Any solvent in which the compound of Formula II is soluble can be used. Typically, the solvent comprises a volatile organic solvent, such as, for example, a (C_1-C_6) alcohol (e.g. ethanol).

A compound of Formula II can also be loaded into a silica material by dissolving the compound in a suitable solvent to provide a solution comprising the compound of Formula II; adding silica particles to the solution to provide a mixture; optionally agitating or stirring the mixture; adding an antisolvent to the mixture; and isolating the solid mixture that comprises the compound of Formula II on the silica particles. Suitable solvents include organic solvents such as ketones (e.g. acetone), alcohols (e.g. ethanol) and halogenated hydrocarbons (e.g. dichloromethane). Suitable antisolvents include highly non-polar solvents (e.g. hexane or

heptane). The final solid mixture can be isolated by any suitable separation technique (e.g. filtration).

One or more pharmaceutically acceptable excipients can be combined with the mixture to provide a second mixture. These pharmaceutically acceptable excipients can include fillers, binders, and disintegrants. In order to improve the processability of the mixture in the subsequent aqueous granulation process, it can be beneficial to select fillers and disintegrants that are compatible with this aqueous process. For example microcrystalline cellulose (filler) and croscarmellose sodium (disintegrant) were found to be particularly compatible with the subsequent aqueous granulation process. Hydroxypropyl cellulose (binder) was also found to be particularly compatible with the subsequent granulation process. In one embodiment of the invention the weight percentage of microcrystalline cellulose to the total weight of the second mixture is about 50% ± 20%. In one embodiment of the invention the weight percentage of hydroxypropyl cellulose to the total weight of the second mixture is 2% ± 1%. In one embodiment of the invention the weight percentage of croscarmellose sodium is 5% ± 2%. Following addition of the pharmaceutically acceptable excipients, the second mixture can be mixed, for example, using a mechanical mixer, such as a high shear granulator (Niro-Fielder, model PMA-25).

Water can be added to the second mixture to provide a wet granulate, which can subsequently be de-agglomerated, e.g. with a 20 mesh sieve. Drying, for example using a fluid bed dryer (Fluid Air, model 20), provides a dried material that comprises solid particles. In one embodiment the dried material has less than about 10.0% moisture content as determined by loss on drying (LOD). In another embodiment the dried material has less than about 5.0% moisture content as determined by loss on drying (LOD). In another embodiment the dried material has less than about 1.0% moisture content as determined by loss on drying (LOD). The size of these particles can be reduced, e.g. using a 40 mesh sieve or a suitable mill (Quadro CoMil, model 197/S) to provide a third mixture.

A suitable pharmaceutically acceptable lubricant/glidant (e.g. magnesium stearate, stearic acid, calcium stearate, zinc stearate, or pregelatinized starch) can be combined with the third mixture to provide a fourth mixture. In one embodiment the weight percentage of magnesium stearate to the total weight of the fourth mixture is 1% ± 0.5%.

In one embodiment, the invention provides a composition prepared by the methods described herein. The invention also provides a product prepared by any of the process steps described herein.

Formulations suitable for oral administration may be presented as discrete units such as tablets each containing a predetermined amount of the active ingredients; as a powder or granules.

A tablet can be made by compression or molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine the active ingredient in a free-flowing form such as a powder or granules, optionally mixed with a binder, lubricant, inert diluent, preservative, surface active agent or dispersing agent. Molded tablets may be made by molding in a suitable machine a mixture of the powdered active ingredient moistened with an inert liquid diluent. The tablets may optionally be coated or scored and optionally are formulated so as to provide slow or controlled release of the active ingredient.

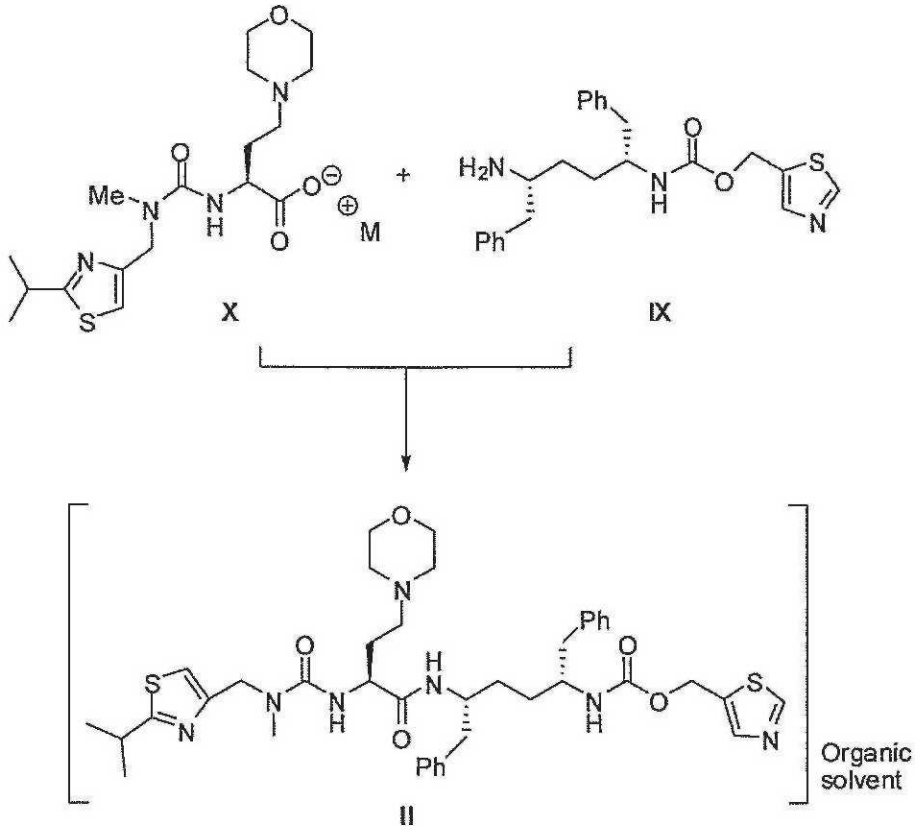
Tablets of the invention may contain one or more agents including sweetening agents, flavoring agents, coloring agents and preserving agents, in order to provide a palatable preparation. Tablets containing the active ingredients in admixture with non-toxic pharmaceutically acceptable excipients which are suitable for manufacture of tablets are acceptable. These excipients may be, for example, inert diluents, such as calcium or sodium carbonate, lactose, lactose monohydrate, croscarmellose sodium, povidone, calcium or sodium phosphate; granulating and disintegrating agents, such as maize starch, or alginic acid; binding agents, such as cellulose, microcrystalline cellulose, starch, gelatin or acacia; and lubricating agents, such as magnesium stearate, stearic acid or talc. Tablets may be uncoated or may be coated by known techniques including microencapsulation to delay disintegration and adsorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate alone or with a wax may be employed.

International Patent Application Publication Number WO 2005/113508 provides certain specific crystalline forms of the compound of Formula I. The entire content of International Patent Application Publication Number WO 2005/113508 is incorporated herein by reference (in particular, see pages 12-62 therein). The specific crystalline forms are identified therein as Crystal Form II and Crystal Form III. Crystal form II has an X-ray powder diffraction pattern having characteristic diffraction peaks at diffraction angles 2θ(°) of 6.56, 13.20, 19.86, 20.84,

21.22, and 25.22 as measured by an X-ray powder diffractometer. Crystal form III has an X-ray powder diffraction pattern having characteristic diffraction peaks at diffraction angles 2θ(°) of 8.54, 14.02, 15.68, 17.06, 17.24, 24.16, and 25.74 as measured by an X-ray powder diffractometer. In one embodiment the compositions of the invention include a compound of formula I that is in Crystal Form II or Crystal Form III.

Preparation of a Compound of Formula II and Preloading on Silica

A compound of Formula II or a salt thereof can be prepared by coupling an acid salt of formula **X** wherein M is a counterion with an amine of formula **IX** to form the corresponding amide of Formula II as described in International Patent Application Publication Number WO 2008/103949 (for example, see page 254).



This amide forming reaction can be carried out under standard conditions. For example, it can be carried out in a suitable organic solvent (e.g. tetrahydrofuran or dichloromethane) in the presence of a suitable coupling agent (e.g. EDC•HCl and HOBt). Other suitable amide coupling reagents and conditions are known in the field. The reaction can typically be carried out at a temperature from about -30 °C to about 20 °C. The final reaction solution containing the compound of Formula II in dichloromethane (DCM) can be directly utilized in the processes illustrated in Figure 2 to provide representative compositions of the invention, or the dichloromethane solution of the compound can be combined with ethanol and the resulting mixture can be distilled to remove the dichloromethane, leaving a solution of the compound of Formula II in ethanol. This ethanol solution can be combined with the silicon dioxide particles and evaporated (as illustrated in the left column of Figure 2) to provide a composition

comprising the compound of Formula II loaded on silicon dioxide particles. Alternatively, the dichloromethane solution of the compound can be combined with silicon dioxide particles, an antisolvent can be added, and the resulting mixture can be filtered and dried (as illustrated in the right column of Figure 2) to provide a composition comprising the compound of Formula II loaded

Figure 1 shows a cross-section of a tablet (101) of the invention. The tablet includes a first layer (103) that comprises a compound of Formula I and a compound of Formula II. The tablet also includes a second layer (105) that comprises a compound of Formula III and a salt of Formula IV.

Figure 2 illustrates processes that can be used to prepare a tablet comprising a compound of formula II.

Figure 3 illustrates a process for preparing a bilayer tablet of the invention. This process is further detailed in Example 1.

The invention will now be illustrated by the following non-limiting Examples.

Examples

Example 1: Tablet Formation.

The manufacturing procedure for a fixed dose combination tablet containing the compounds of Formulas I, II, III and IV include the following steps: 1) fluid-bed granulation and drying of the compound of Formula I, 2) high-shear granulation and fluid-bed drying of the compound of Formula II or the compound of Formula II loaded on silica particles, 3) dry granulation of the compound of Formula III and dry granulation of the salt of Formula IV, 4) milling of the dry granulation of the compound of Formula III and milling of the dry granulation of the salt of Formula IV, 5) blending of the compound of Formula III and the salt of Formula IV, 6) blending of the compound of Formula I and the compound of Formula II, 7) bilayer compression with one layer consisting of the blend of the compounds of Formula I and Formula II and the other layer consisting of the blend of the compounds of Formula III and Formula IV to form a tablet, 8) coating of the tablet and 9) packaging of the coated tablet.

The in-process weight control for a bilayer tablet was superior compared to a trilayer tablet configuration. Bilayer weight control for the layer containing the compounds of Formula I and Formula II was between 100.2% and 100.8% of the mean target layer weight. Mean weights for the total tablet was between 99.5% and 100.7% of the mean target tablet weight. The relative standard deviation (RSD) value for the layer containing the compounds of Formula I and Formula II was between 1.4% and 2.2%, while the RSD for the total tablet was between 0.7% and 1.2%. These low RSD values indicate very low weight variability during the bilayer tablet compression process. The friability at the start and end of the compression process was 0.0%. No chipped, capped, or broken tablets were observed during bilayer compression.

Significant material property and compressibility differences between layer 1 and layer 2 presented the risk of layer separation during the tablet compression process using the process of preloading Formula II on a solid carrier. However, the resulting tablets were intact with no defects and withstood the aggressive film-coating process.

As described above, an intermediate mixture that is useful for preparing a layer of a tablet of the invention is a composition that comprises the compound of formula I and the compound of formula II. Accordingly, in one embodiment, the invention provides a composition comprising the compound of formula I and the compound of formula II. The compounds of formula I and II may be present in any of the amounts or ratios described herein

that are useful for preparing tablets of the invention. For example, the following tables illustrate representative components and weight ratios for intermediate compositions that can be used to prepare tablets of the invention.

Components	% w/w	mg/tablet
Compound of Formula I	26.77	150.0
Compound of Formula II	13.39	75.0
Colloidal Silicon Dioxide	15.40	86.3
Lactose Monohydrate	1.94	10.9
Microcrystalline Cellulose	29.32	164.3
Hydroxypropyl Cellulose	1.34	7.5
Hydroxypropyl Cellulose	0.80	4.5
Sodium Lauryl Sulfate	2.01	11.3
Croscarmellose Sodium	8.03	45.1
Magnesium Stearate	1.00	5.6
Total	100.0	560

Components	% w/w	mg/tablet
Compound of Formula I	23.61	150.0
Compound of Formula II	15.74	100.0
Colloidal Silicon Dioxide	18.10	115.0
Lactose Monohydrate	1.71	10.9
Microcrystalline Cellulose	28.26	179.5

Hydroxypropyl Cellulose	1.18	7.5
Hydroxypropyl Cellulose	0.94	6.0
Sodium Lauryl Sulfate	1.77	11.3
Croscarmellose Sodium	7.67	48.8
Magnesium Stearate	1.00	6.4
Total	100.0	635

Components	% w/w	mg/tablet
Compound of Formula I	19.10	150.0
Compound of Formula II	19.10	150.0
Colloidal Silicon Dioxide	21.97	172.5
Lactose Monohydrate	1.38	10.9
Microcrystalline Cellulose	26.74	210.0
Hydroxypropyl Cellulose	0.96	7.5
Hydroxypropyl Cellulose	1.15	9.0
Sodium Lauryl Sulfate	1.43	11.3
Croscarmellose Sodium	7.16	56.3
Magnesium Stearate	1.00	7.9
Total	100.0	785

Attempts were also made to manufacture the tablet as a trilayer. During trilayer compression, the tablet was manufactured with a layer of the compound of Formula I, a second layer containing the compound of Formula II and a third layer containing a blend of the compound of Formula III and the salt of Formula IV. The initial attempt was to apply the compound of Formula II as the first layer, followed by the compound of Formula I on the second layer, and lastly the third layer of a blend of the compound of Formula III and the salt of Formula IV. The weight of the layer containing the compound of Formula II was too small, which resulted in a very thin tablet layer that caused a collision of the upper and lower punches during the compression process. The weight of the layer containing the compound of Formula I was similar to the weight of the layer containing the compound of Formula II so punch collision would prevent the reverse configuration. The weight of the blend containing the compounds of Formula III and IV was large enough to present a layer thickness that prevented punch collision

during the compression process. This layer was followed by a second layer containing the compound of Formula I and a third layer containing the compound of Formula II.

Severe tablet defects were observed during the tablet compression process involving the three layer process. There was also difficultly controlling the tablet weight with this three layer configuration as well. In addition, the second and third layers separated from the tablet core. An intact tablet, free of defects could not be produced for the trilayer tablet configuration.

Combining all of the compounds of Formula I, II, III, and IV in a mixture to produce a mono layer tablet compromises the chemical stability of Formula IV. Additionally, improper combinations of the compounds of Formula I, II, III, or IV in a bilayer tablet will also compromise the chemical stability of Formula IV. For example, Formula IV in the presence of silicon dioxide from the Formula II formulation will decompose Formula IV; and Formula IV in the in the presence of sodium lauryl sulphate from the Formula I formulation will decompose Formula IV.

Example 2

The following table illustrates representative compositions of the invention that are useful in preparing tablets of the invention.

	Compound of Formula II 75 mg Formulation		Compound of Formula II 100 mg Formulation	
Components	% w/w	mg/tablet	% w/w	mg/tablet
Compound of Formula III	16.5	200.0	15.5	200.0
Salt of Formula IV	24.7	300.0	23.3	300.0
Compound of Formula I	12.4	150.0	11.7	150.0
Compound of Formula II	6.2	75.0	7.8	100.0
Colloidal Silicon Dioxide	7.1	86.3	8.9	115.0
Lactose Monohydrate	0.9	10.9	0.8	10.9
Microcrystalline Cellulose	20.9	253.8	20.9	269.0
Hydroxypropyl Cellulose	0.6	7.5	0.6	7.5

Hydroxypropyl Cellulose	0.4	4.5	0.5	6.0
Sodium Lauryl Sulfate	0.9	11.3	0.9	11.3
Croscarmellose Sodium	7.7	93.1	7.5	96.8
Magnesium Stearate	1.7	20.1	1.6	20.9
Total	100	1212	100	1287

Example 3

The following table illustrates a representative composition of the invention that is useful in preparing tablets of the invention.

Components	Compound of Formula II 150 mg Formulation	
	% w/w	mg/tablet
Compound of Formula III	13.9	200.0
Salt of Formula IV	20.9	300.0
Compound of Formula I	10.4	150.0
Compound of Formula II	10.4	150.0
Colloidal Silicon Dioxide	12.0	172.5
Lactose Monohydrate	0.8	10.9
Microcrystalline Cellulose	20.8	299.5
Hydroxypropyl Cellulose	0.5	7.5
Hydroxypropyl Cellulose	0.6	9.0
Sodium Lauryl Sulfate	0.8	11.3
Croscarmellose Sodium	7.3	104.3
Magnesium Stearate	1.6	22.4
Total	100	1437

Example 4

The following table illustrates a representative composition of the invention that is useful in preparing tablets of the invention.

Components	Compound of Formula II 150 mg Formulation	
	% w/w	mg/tablet
Compound of Formula III	14.8	200.0
Salt of Formula IV	22.2	300.0
Compound of Formula I	11.1	150.0
Compound of Formula II	11.1	150.0
Lactose Monohydrate	0.8	10.9
Microcrystalline Cellulose	19.5	267.1
Colloidal Silicon Dioxide	11.1	150.0
Croscarmellose Sodium	6.1	81.75
Hydroxypropyl Cellulose	0.6	7.5
Sodium Lauryl Sulfate	0.8	11.25
Magnesium Stearate	1.6	21.5
Tablet Core Weight	100	1350

Example 5.

A pharmacokinetic dose-ranging study involving the compound of Formula II was conducted. Doses for the compound of Formula II for the dose-ranging study included 50 mg, 100 mg and 200 mg. Pharmacokinetic data for the compound of Formula II from the dose-ranging study for the 100 mg dose is provided in the table below.

PK Data for the compound of Formula II (dose-ranging study; 100 mg dose; n=11)	
AUC_{tau} (ng.hr/mL)	3440 (34.3)
C_{max} (ng/mL)	563 (30.7)
$T_{1/2}$ (h)	3.12 (2.55, 3.36)
CL/F (L/h)	33.2 (43.6)

Q1 = first quartile; Q3 = third quartile

Data presented as arithmetic mean (%CV); $T_{½}$ presented as median (Q1, Q3); %CV, percent coefficient of variation

A pharmacokinetic study involving two fixed-dose combinations of the compounds of Formula I/ Formula II/ Formula III/ Formula IV was also conducted. Doses for the fixed-dose combination of the compounds of Formula I/ Formula II/ Formula III/ Formula IV were 150 mg/100 mg/ 200 mg/300 mg (Dose A) and 150 mg/150 mg/ 200 mg/300 mg (Dose B) respectively. Phamacokinetic data for the compound of Formula II from the study involving the two fixed dose combinations (Dose A and Dose B) is provided in the following table.

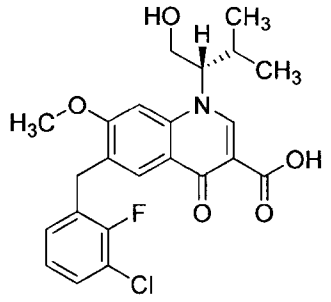
PK Data for the compound of Formula II (fixed dose combination study)		
Mean (CV%) PK	Dose A (n = 43)	Dose B (n = 42)
AUC_{tau} (ng.hr/mL)	5150 (31.7)	10400 (35.1)
C_{max} (ng/mL)	855 (27.6)	1570 (29.7)
C_{tau} (ng/mL)	10.8 (83.6)	23.3 (103)

The overall exposure, as measured by area under the curve (AUC), was 1.5 to 2-fold higher for the compound of Formula II in the fixed-dose combination study (Dose A) versus the exposure as measured by area under the curve (AUC) for the dose-ranging study involving the 100 mg dose of the compound of Formula II. Due to this increased exposure to the compound of Formula II, either less of the compound of Formula I is required to provide an equivalent antiviral effect, or the same amount of the compound of Formula I will provide an increased antiviral effect when the compound of Formula I is administered with the compound of Formula II. This unexpected increased exposure to the compound of Formula II, also suggests that less compound of Formula II is required to achieve effective exposure to the compound of Formula II.

All publications, patents, and patent documents are incorporated by reference herein, as though individually incorporated by reference. The invention has been described with reference to various specific and preferred embodiments and techniques. However, it should be understood that many variations and modifications may be made while remaining within the spirit and scope of the invention.

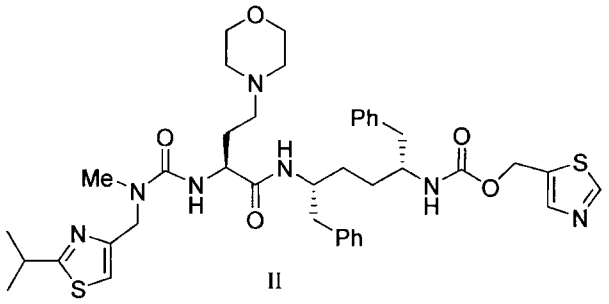
The claims defining the invention are as follows:

1. Use of a compound of Formula I:



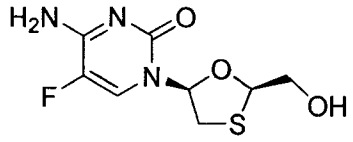
I ; and

a compound of Formula II:



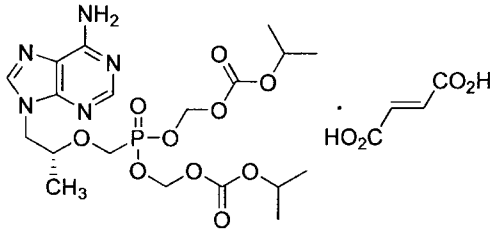
II ; and

a compound of Formula III:



III ; and

a salt of Formula IV:



IV ;

in the manufacture of a medicament for treating HIV;

wherein the medicament is in the form of a tablet comprising a first layer and a second layer, wherein the first layer comprises the compound of Formula I and the compound of formula II, and optionally a pharmaceutically acceptable carrier, and the second layer comprises the compound of formula III and the salt of formula IV, and optionally a pharmaceutically acceptable carrier.

2. The use of claim 1, wherein the first layer of the tablet is in contact with the second layer.

3. The use of claim 1 or claim 2, wherein the weight of the compound of Formula I divided by the weight of the compound of Formula II is 1.0 ± 0.5.

4. The use of any one of claims 1 to 3, wherein the first layer of the tablet further comprises a plurality of silica particles.

5. The use of claim 4, wherein the compound of Formula II is associated with silica particles.

6. The use of any one of claims 1 to 5, wherein the tablet comprises 150 mg ± 10% of the compound of Formula I; 150 mg± 10% of the compound of Formula II;
200 mg ± 10% of the compound of Formula III; and 300 mg ± 10% of the salt of Formula IV.

7. The use of claim 6, wherein the tablet comprises 150 mg ± 5% of the compound of Formula I.

8. The use of claim 7, wherein the tablet comprises 150 mg ± 2% of the compound of Formula I.

9. The use of any one of claims 6 to 8, wherein the tablet comprises 150 mg ± 5% of the compound of Formula II.

10. The use of claim 9, wherein the tablet comprises 150 mg ± 2% of the compound of Formula II.

11. The use of any one of claims 6 to 10, wherein the tablet comprises 200 mg ± 5 % of the compound of Formula III.

12. The use of claim 11, wherein the tablet comprises 200 mg ± 2% of the compound of Formula III.

13. The use of any one of claims 6 to 12, wherein the tablet comprises 300 mg ± 5 % of the salt of Formula IV.

14. The use of claim 13, wherein the tablet comprises 300 mg ± 2% of the salt of Formula IV.

15. The use of any one of claims 1 to 14, wherein the tablet further comprises a pharmaceutically acceptable carrier.

16. The use of any one of claims 1 to 15, wherein the tablet is formulated in unit dose form.

17. The use of any one of claims 1 to 16, wherein the tablet is formulated for once a day dosing.

18. The use of claim 1, wherein the tablet core weight is 1350 mg and consists of the following components: 150.0 mg of the compound of Formula I; 150.0 mg of the compound of Formula II; 200.0 mg of the compound of Formula III; 300.0 mg of the salt of formula IV; 10.9 mg of lactose monohydrate; 267.1 mg of microcrystalline cellulose; 150.0 mg of colloidal silicon dioxide; 81.75 mg of croscarmellose sodium; 7.5 mg of hydroxypropyl cellulose; 11.25 mg of sodium lauryl sulfate; and 21.5 mg of magnesium stearate.