

METHOD OF PREPARING GLUCOSYLCERAMIDE

SYNTHASE INHIBITORS

Background of the Invention

5 The invention relates to a method of preparing inhibitors of glucosylceramide synthase (GCS) useful for the treatment metabolic diseases, such as lysosomal storage diseases, either alone or in combination with enzyme replacement therapy, and for the treatment of cancer.

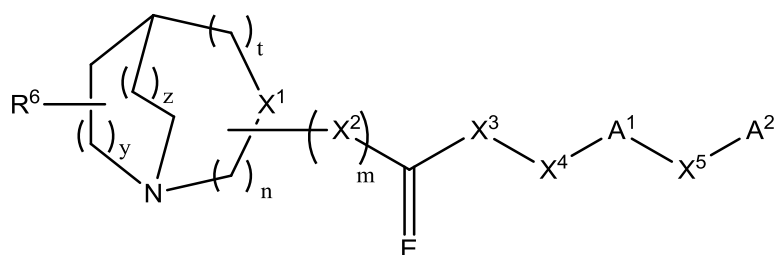
 Glucosylceramide synthase (GCS) is a pivotal enzyme which catalyzes the initial
10 glycosylation step in the biosynthesis of glucosylceramide-base glycosphingolipids (GSLs) namely via the pivotal transfer of glucose from UDP-glucose (UDP-Glc) to ceramide to form glucosylceramide. GCS is a transmembrane, type III integral protein localized in the cis/medial Golgi. Glycosphingolipids (GSLs) are believed to be integral for the dynamics of many cell membrane events, including cellular interactions, signaling
15 and trafficking. Synthesis of GSL structures has been shown (see, Yamashita et al., Proc. Natl. Acad. Sci. USA 1999, 96(16), 9142-9147) to be essential for embryonic development and for the differentiation of some tissues. Ceramide plays a central role in sphingolipid metabolism and downregulation of GCS activity has been shown to have marked effects on the sphingolipid pattern with diminished expression of
20 glycosphingolipids. Sphingolipids (SLs) have a biomodulatory role in physiological as well as pathological cardiovascular conditions. In particular, sphingolipids and their regulating enzymes appear to play a role in adaptive responses to chronic hypoxia in the neonatal rat heart (see, El Alwanit et al., Prostaglandins & Other Lipid Mediators 2005, 78(1-4), 249-263).

GCS inhibitors have been proposed for the treatment of a variety of diseases (see for example, WO2005068426). Such treatments include treatment of glycolipid storage diseases (e.g., Tay Sachs, Sandhoffs, GM2 Activator deficiency, GM1 gangliosidosis and Fabry diseases), diseases associated with glycolipid accumulation (e.g., Gaucher disease; Miglustat (Zavesca), a GCS inhibitor, has been approved for therapy in type 1 Gaucher disease patients, see, Treiber et al., *Xenobiotica* 2007, 37(3), 298-314), diseases that cause renal hypertrophy or hyperplasia such as diabetic nephropathy; diseases that cause hyperglycemia or hyperinsulemia; cancers in which glycolipid synthesis is abnormal, infectious diseases caused by organisms which use cell surface glycolipids as receptors, infectious diseases in which synthesis of glucosylceramide is essential or important, diseases in which synthesis of glucosylceramide is essential or important, diseases in which excessive glycolipid synthesis occurs (e.g., atherosclerosis, polycystic kidney disease, and renal hypertrophy), neuronal disorders, neuronal injury, inflammatory diseases or disorders associated with macrophage recruitment and activation (e.g., rheumatoid arthritis, Crohn's disease, asthma and sepsis) and diabetes mellitus and obesity (see, WO 2006053043).

In particular, it has been shown that overexpression of GCS is implicated in multi-drug resistance and disrupts ceramide-induced apoptosis. For example, Turzanski et al., (*Experimental Hematology* 2005, 33 (1), 62-72 have shown that ceramide induces apoptosis in acute myeloid leukemia (AML) cells and that P-glycoprotein (p-gp) confers resistance to ceramide-induced apoptosis, with modulation of the ceramide-glucosylceramide pathway making a marked contribution to this resistance in TF-1 cells. Thus, GCS inhibitors can be useful for treatment of proliferative disorders by inducing apoptosis in diseased cells.

Summary of the Invention

The present invention relates to a method of preparing a compound of the formula,



I

wherein:

n is 1, 2 or 3;

m is 1;

t is 0, 1 or 2;

y is 1 or 2;

z is 0, 1 or 2;

E is O;

X¹ is CR¹;

X² is O ;

X³ is -NH;

X⁴ is CR⁴R⁵, CH₂ CR⁴R⁵ or CH₂ -(C₁-C₆) alkyl-CR⁴R⁵;

X⁵ is a direct bond, O, S, SO₂, CR⁴R⁵; (C₁-C₆)alkyl, (C₁-C₆)alkyloxy, (C₁-

C₆)alkenyl, (C₁-C₆)alkenyloxy;

R is (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₁-C₆)alkyl, (C₂-C₉)heteroaryl(C₁-C₆)alkyl;

R¹ is H, CN, (C₁-C₆)alkylcarbonyl, or (C₁-C₆)alkyl;

R² and R³ are each independently -H, (C₁-C₆)alkyl optionally substituted by one or

more substituents selected from the group consisting of halogen, (C₁-C₆)alkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₁-C₆)alkyl(C₆-C₁₂)aryl, halo(C₆-C₁₂)aryl, and halo(C₂-C₉)heteroaryl,

or optionally when X² is -NR² and X³ is -NR³, R² and R³ may be taken together with the nitrogen atoms to which they are attached form a non-aromatic heterocyclic ring optionally substituted by with one or more substituents selected from halogen, (C₁-C₆)alkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₁-C₆)alkyl(C₆-C₁₂)aryl, halo(C₆-C₁₂)aryl, and halo(C₂-C₉)heteroaryl;

R^4 and R^5 are independently selected from H, (C₁-C₆)alkyl, or taken together with the carbon to which they are attached to form a spiro (C₃-C₁₀)cycloalkyl ring or spiro (C₃-C₁₀)cycloalkoxy ring;

R^6 is -H, halogen, -CN, (C₆-C₁₂)aryl, (C₆-C₁₂)aryloxy, (C₁-C₆)alkyloxy;

5 (C₁-C₆)alkyl optionally substituted by one to four halo or (C₁-C₆)alkyl;

A^1 is (C₂-C₆)alkynyl; (C₃-C₁₀)cycloalkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₂-C₉)heterocycloalkyl or benzo(C₂-C₉)heterocycloalkyl optionally substituted with one or more substituents selected from the group consisting of halo, (C₁-C₆)alkyl optionally substituted by one to three halo; (C₁-C₆)alkenyl, amino, (C₁-C₆)alkylamino, (C₁-

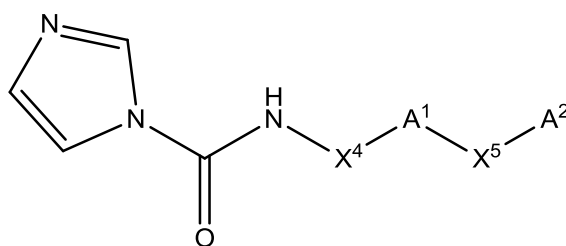
10 C₆)dialkylamino, (C₁-C₆)alkoxy, nitro, CN, -OH, (C₁-C₆)alkyloxy optionally substituted by one to three halo; (C₁-C₆)alkoxycarbonyl, and (C₁-C₆) alkylcarbonyl;

A^2 is H, (C₃-C₁₀)cycloalkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₂-C₉)heterocycloalkyl or benzo(C₂-C₉)heterocycloalkyl optionally substituted with one or more substituents selected from the group consisting of halo, (C₁-C₆)alkyl optionally

15 substituted by one to three halo; (C₁-C₆)alkylenyl, amino, (C₁-C₆) alkylamino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, O(C₃-C₆ cycloalkyl), (C₃-C₆) cycloalkoxy, nitro, CN, OH, (C₁-C₆)alkyloxy optionally substituted by one to three halo; (C₃-C₆) cycloalkyl, (C₁-C₆) alkoxycarbonyl, (C₁-C₆) alkylcarbonyl, (C₁-C₆) haloalkyl;

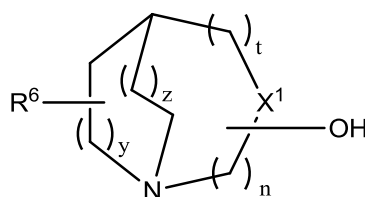
with the proviso that the sum of $n + t + y + z$ is not greater than 6;

20 comprising reacting the compound of Formula II



II

with the compound of Formula III



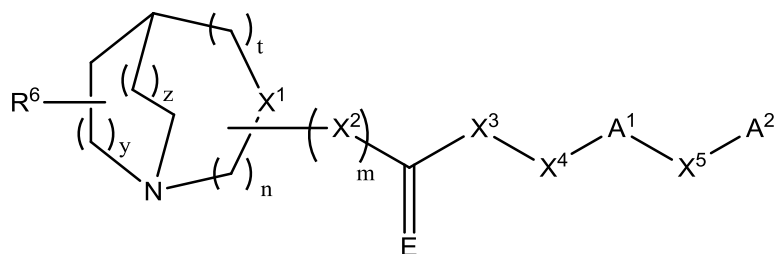
III

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wherein n, t, y, z, X⁴, A¹, X⁵ and A² are as defined above.

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The present invention relates to a method of preparing a compound of the formula,



I

wherein:

10

n is 1, 2 or 3;

m is 1;

t is 0, 1 or 2;

y is 1 or 2;

z is 0, 1 or 2;

15

E is O;

X¹ is CR¹;

X² is O ;

X³ is -NH;

X⁴ is CR⁴R⁵, CH₂ CR⁴R⁵ or CH₂ -(C₁-C₆) alkyl-CR⁴R⁵;

20

X⁵ is a direct bond, O, S, SO₂, CR⁴R⁵; (C₁-C₆)alkyl, (C₁-C₆)alkyloxy, (C₁-C₆)alkenyl, (C₁-C₆)alkenyloxy;

R is (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₁-C₆)alkyl, (C₂-C₉)heteroaryl(C₁-C₆)alkyl;

R¹ is H, CN, (C₁-C₆)alkylcarbonyl, or (C₁-C₆)alkyl;

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R² and R³ are each independently -H, (C₁-C₆)alkyl optionally substituted by one or more substituents selected from the group consisting of halogen, (C₁-C₆)alkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₁-C₆)alkyl(C₆-C₁₂)aryl, halo(C₆-C₁₂)aryl, and halo(C₂-C₉)heteroaryl, or optionally when X² is -NR² and X³ is -NR³, R² and R³ may be taken together with the nitrogen atoms to which they are attached form a non-aromatic heterocyclic ring optionally substituted by with one or more substituents selected from halogen, (C₁-C₆)alkyl, (C₆-

C₁₂)aryl, (C₂-C₉)heteroaryl, (C₁-C₆)alkyl(C₆-C₁₂)aryl, halo(C₆-C₁₂)aryl, and halo(C₂-C₉)heteroaryl;

R⁴ and R⁵ are independently selected from H, (C₁-C₆)alkyl, or taken together with the carbon to which they are attached to form a spiro (C₃-C₁₀)cycloalkyl ring or spiro (C₃-

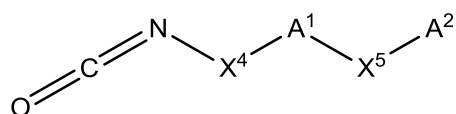
5 C₁₀)cycloalkoxy ring;

R⁶ is -H, halogen, -CN, (C₆-C₁₂)aryl, (C₆-C₁₂)aryloxy, (C₁-C₆)alkyloxy; (C₁-C₆)alkyl optionally substituted by one to four halo or (C₁-C₆)alkyl;

A¹ is (C₂-C₆)alkynyl; (C₃-C₁₀)cycloalkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₂-C₉)heterocycloalkyl or benzo(C₂-C₉)heterocycloalkyl optionally substituted with one or
10 more substituents selected from the group consisting of halo, (C₁-C₆)alkyl optionally substituted by one to three halo; (C₁-C₆)alkenyl, amino, (C₁-C₆)alkylamino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, nitro, CN, -OH, (C₁-C₆)alkyloxy optionally substituted by one to three halo; (C₁-C₆)alkoxycarbonyl, and (C₁-C₆) alkylcarbonyl;

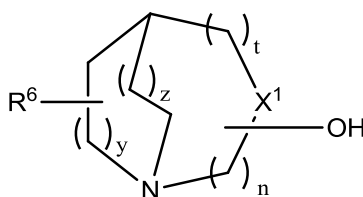
A² is H, (C₃-C₁₀)cycloalkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₂-C₉)heterocycloalkyl or benzo(C₂-C₉)heterocycloalkyl optionally substituted with one or
15 more substituents selected from the group consisting of halo, (C₁-C₆)alkyl optionally substituted by one to three halo; (C₁-C₆)alkylenyl, amino, (C₁-C₆) alkylamino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, O(C₃-C₆ cycloalkyl), (C₃-C₆) cycloalkoxy, nitro, CN, OH, (C₁-C₆)alkyloxy optionally substituted by one to three halo; (C₃-C₆) cycloalkyl, (C₁-
20 C₆) alkoxycarbonyl, (C₁-C₆) alkylcarbonyl, (C₁-C₆) haloalkyl;

with the proviso that the sum of n + t + y + z is not greater than 6;
comprising reacting the compound of Formula IV



IV

25 with a compound of Formula III

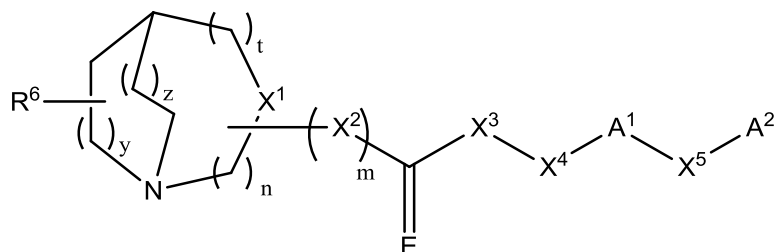


III

wherein n, t, y, z, X⁴, A¹, X⁵ and A² are as defined above.

The present invention relates to a method of preparing a compound of the formula,

5



I

wherein:

n is 1, 2 or 3;

m is 1;

10

t is 0, 1 or 2;

y is 1 or 2;

z is 0, 1 or 2;

E is O;

15

X¹ is CR¹;

X² is O ;

X³ is -NH;

X⁴ is CR⁴R⁵, CH₂ CR⁴R⁵ or CH₂ -(C₁-C₆) alkyl-CR⁴R⁵;

X⁵ is a direct bond, O, S, SO₂, CR⁴R⁵; (C₁-C₆)alkyl, (C₁-C₆)alkyloxy, (C₁-

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C₆)alkenyl, (C₁-C₆)alkenyloxy;

R is (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₁-C₆)alkyl, (C₂-C₉)heteroaryl(C₁-C₆)alkyl;

R¹ is H, CN, (C₁-C₆)alkylcarbonyl, or (C₁-C₆)alkyl;

R² and R³ are each independently -H, (C₁-C₆)alkyl optionally substituted by one or

more substituents selected from the group consisting of halogen, (C₁-C₆)alkyl, (C₆-C₁₂)aryl,

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(C₂-C₉)heteroaryl, (C₁-C₆)alkyl(C₆-C₁₂)aryl, halo(C₆-C₁₂)aryl, and halo(C₂-C₉)heteroaryl,

or optionally when X² is -NR² and X³ is -NR³, R² and R³ may be taken together with the nitrogen atoms to which they are attached form a non-aromatic heterocyclic ring optionally

substituted by with one or more substituents selected from halogen, (C₁-C₆)alkyl, (C₆-

C₁₂)aryl, (C₂-C₉)heteroaryl, (C₁-C₆)alkyl(C₆-C₁₂)aryl, halo(C₆-C₁₂)aryl, and halo(C₂-C₉)heteroaryl;

R⁴ and R⁵ are independently selected from H, (C₁-C₆)alkyl, or taken together with the carbon to which they are attached to form a spiro (C₃-C₁₀)cycloalkyl ring or spiro (C₃-

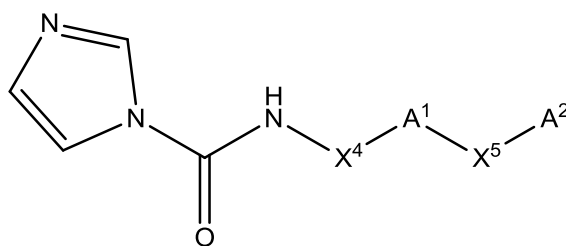
5 C₁₀)cycloalkoxy ring;

R⁶ is -H, halogen, -CN, (C₆-C₁₂)aryl, (C₆-C₁₂)aryloxy, (C₁-C₆)alkyloxy; (C₁-C₆)alkyl optionally substituted by one to four halo or (C₁-C₆)alkyl;

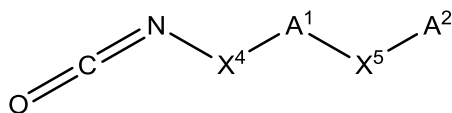
A¹ is (C₂-C₆)alkynyl; (C₃-C₁₀)cycloalkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₂-C₉)heterocycloalkyl or benzo(C₂-C₉)heterocycloalkyl optionally substituted with one or
10 more substituents selected from the group consisting of halo, (C₁-C₆)alkyl optionally substituted by one to three halo; (C₁-C₆)alkenyl, amino, (C₁-C₆)alkylamino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, nitro, CN, -OH, (C₁-C₆)alkyloxy optionally substituted by one to three halo; (C₁-C₆)alkoxycarbonyl, and (C₁-C₆) alkylcarbonyl;

A² is H, (C₃-C₁₀)cycloalkyl, (C₆-C₁₂)aryl, (C₂-C₉)heteroaryl, (C₂-
15 C₉)heterocycloalkyl or benzo(C₂-C₉)heterocycloalkyl optionally substituted with one or more substituents selected from the group consisting of halo, (C₁-C₆)alkyl optionally substituted by one to three halo; (C₁-C₆)alkylenyl, amino, (C₁-C₆) alkylamino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, O(C₃-C₆ cycloalkyl), (C₃-C₆) cycloalkoxy, nitro, CN, OH, (C₁-C₆)alkyloxy optionally substituted by one to three halo; (C₃-C₆) cycloalkyl, (C₁-
20 C₆) alkoxycarbonyl, (C₁-C₆) alkylcarbonyl, (C₁-C₆) haloalkyl;

with the proviso that the sum of n + t + y + z is not greater than 6;
comprising reacting the compounds of Formula II and Formula IV

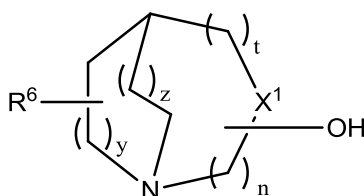


II



IV

with a compound of Formula III



III

5

wherein n, t, y, z, X⁴, A¹, X⁵ and A² are as defined above.

The present invention further relates to a method wherein n is 1; t is 0; y is 1 and z is 1.

10

The present invention further relates to a method wherein X⁴ is CR⁴R⁵.

The present invention further relates to a method wherein R⁴ and R⁵ are each methyl.

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The present invention further relates to a method wherein A¹ is (C₂-C₉)heteroaryl.

The present invention further relates to a method wherein A¹ is thiophene, thiazole, isothiazole, furane, oxazole, isoxazole, pyrrole, imidazole, pyrazole, triazole, pyridine, pymiridine, pyridazine, indole, benzotiazole, benzoisoxazole, benzopyrazole, benzoimidazole, benzofuran, benzooxazole or benzoisoxazole.

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The present invention further relates to a method wherein A¹ is thiazole.

The present invention further relates to a method wherein R⁶ is H.

The present invention further relates to a method wherein X⁵ is a direct bond.

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The present invention further relates to a method wherein A² is (C₆-C₁₂)aryl.

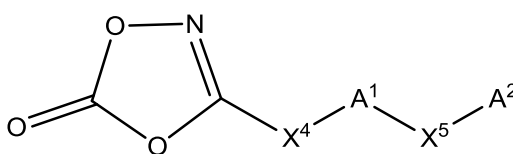
The present invention further relates to a method wherein A² is phenyl.

The present invention further relates to a method wherein the phenyl group is substituted by halo.

The present invention further relates to a method wherein the halo group is fluoro.

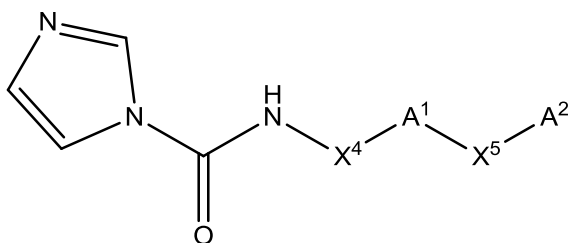
5 The present invention further relates to a method wherein R¹ is hydrogen.

The present invention further relates to a method including reacting the compound of Formula V



V

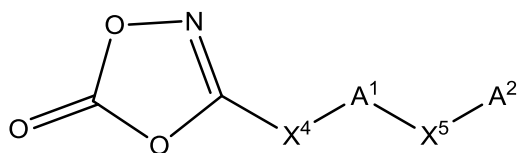
10 with imidazole to form the compound of Formula II



II

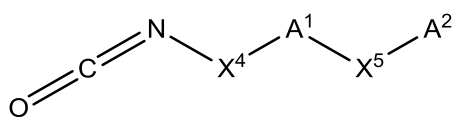
wherein X⁴, A¹, X⁵ and A² are as defined above.

15 The present invention further relates to a method including heating to reflux the compound of Formula V



V

to form the compound of Formula IV

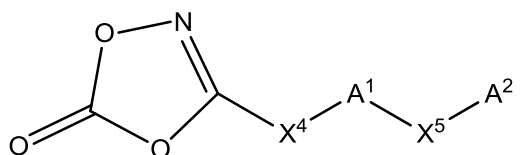


IV

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wherein X^4 , A^1 , X^5 and A^2 are as defined above.

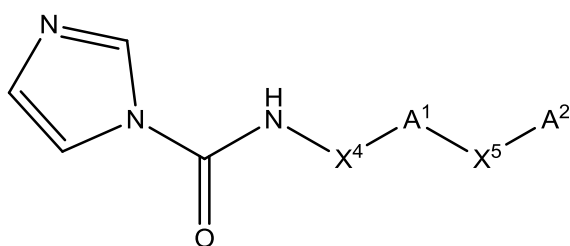
The present invention further relates to a method including reacting, while heating to reflux, the compound of Formula V



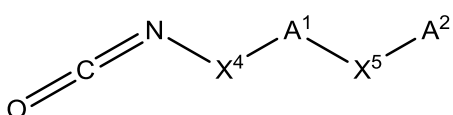
V

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with imidazole to form the compounds of Formula II and Formula IV



II ;

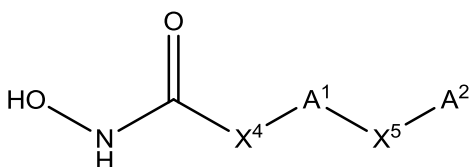


IV

wherein X^4 , A^1 , X^5 and A^2 are as defined above.

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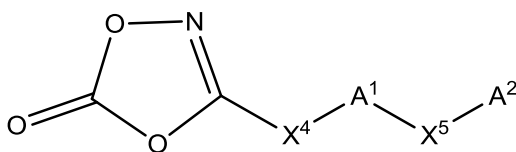
The present invention further relates to a method including reacting the compound of Formula VI



VI

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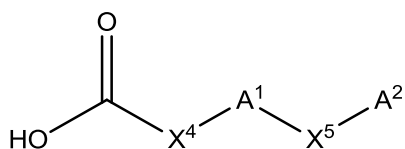
with N, N'- carbonyldiimidazole to form the compound of Formula V



V

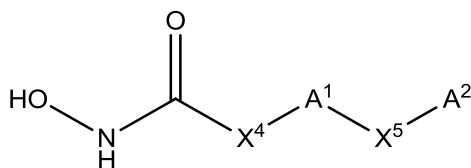
wherein X^4 , A^1 , X^5 and A^2 are as defined above.

5 The present invention further relates to a method including reacting the compound of Formula VII



VII

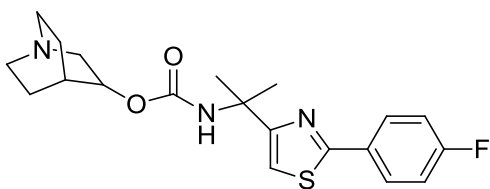
10 with N, N'- carbonyldiimidazole and hydroxylamine to form the compound of Formula VI



VI

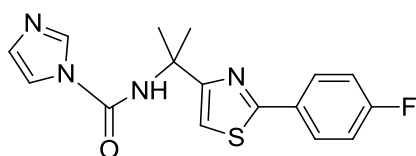
15 wherein X^4 , A^1 , X^5 and A^2 are as defined above.

The present invention relates to a method of preparing the compound of Formula VIII



VIII

20 comprising reacting a compound of Formula IX

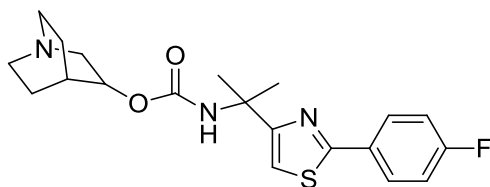


IX

with quinuclidinol.

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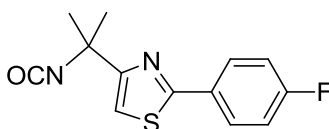
The present invention relates to a method of preparing the compound of Formula VIII



VIII

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comprising reacting a compound of Formula X



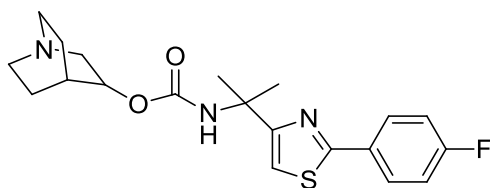
X

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with quinuclidinol.

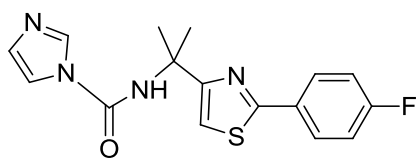
The present invention relates to a method of preparing the compound of Formula VIII

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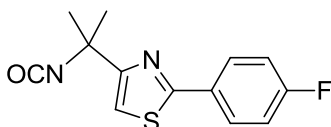
VIII

comprising reacting a compounds of Formula IX and Formula X



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IX ;

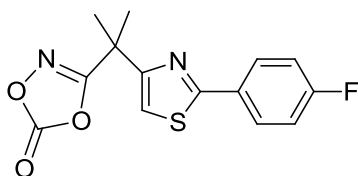


X

10

with quinuclidinol.

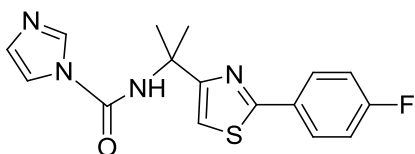
The present invention further relates to a method including reacting the compound of Formula XI



XI

15

with imidazole to form the compound of Formula IX

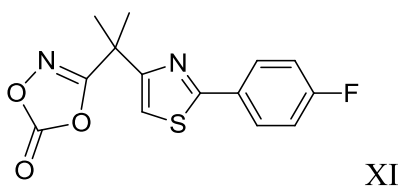


IX

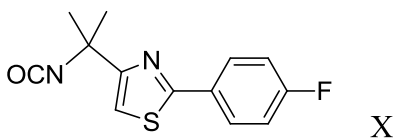
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The present invention further relates to a method including heating to reflux the compound of Formula XI

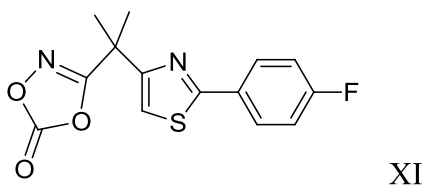


to form the compound of Formula X

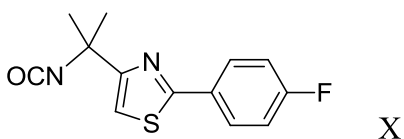
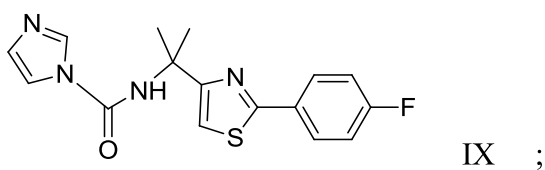


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The present invention further relates to a method including reacting, while heating to reflux, the compound of Formula XI

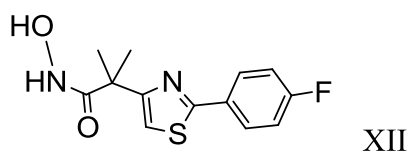


10 with imidazole to form the compounds of Formula IX and Formula X



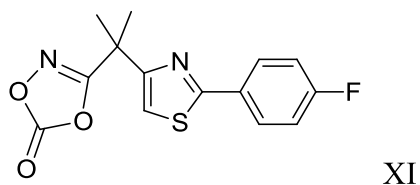
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The present invention further relates to a method including reacting the compound of Formula XII



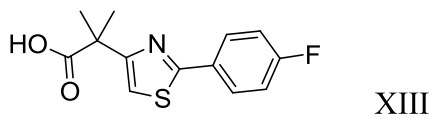
with N, N'- carbonyldiimidazole to form the compound of Formula XI

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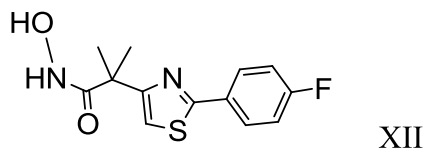
The present invention further relates to a method including reacting the compound of Formula XIII

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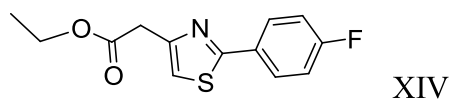
with N, N'- carbonyldiimidazole and hydroxylamine to form the compound of Formula XII

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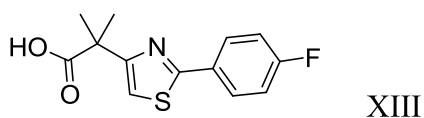


The present invention further relates to a method including reacting the compound of Formula XIV

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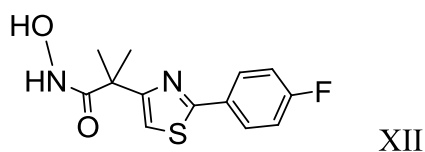


with potassium tert-butoxide and methyl iodine followed by reacting the ethyl ester so formed with lithium hydroxide to form the compound of Formula XIII



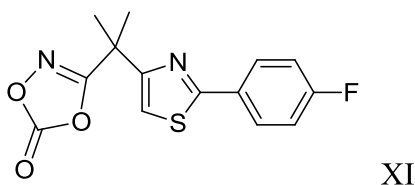
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The present invention relates to a compound of Formula XII



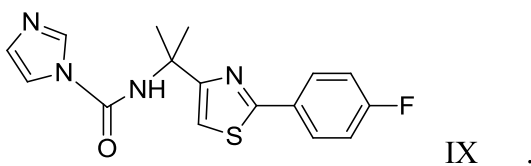
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The present invention relates to a compound of Formula XI



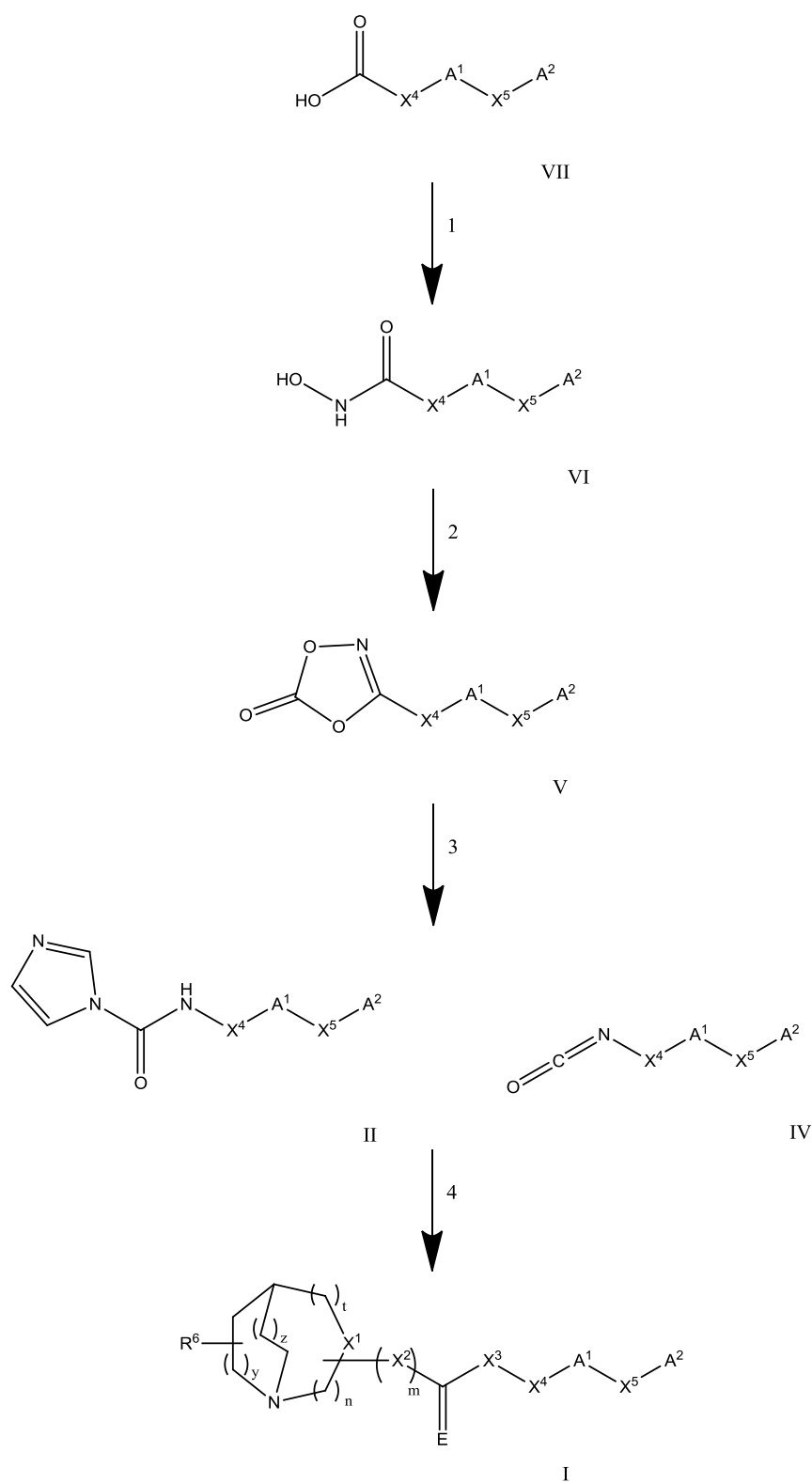
The present invention relates to a compound of Formula IX

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Detailed Description of the Invention

Scheme 1



In reaction 1 of Scheme 1, the carboxylic acid compound of Formula **VII** is converted to the corresponding hydroxamic acid compound of Formula **VI** by reacting **VII** with N, N'- carbonyldiimidazole (i.e. CDI) in a polar aprotic solvent, such as

tetrahydrofuran (THF). The solution is stirred at a temperature between about -5°C to about 25°C, preferable about 20°C, for a time period between about 5 minutes to about 30 minutes, preferably about 10 to 15 minutes. The resulting solution mixture is allowed to warm to room temperature and stirred for an additional time period between about 30 minutes to about 2 hours, preferably about 1 hour. Hydroxylamine is then added to the solution mixture at a temperature between about -5°C to about 10°C, preferable about 3°C. The resulting reaction mixture is stirred under inert atmosphere (i.e., nitrogen) for a time period between about 5 min to about 8 hours, preferably about 10 min.

In reaction 2 of Scheme 1, the hydroxamic acid compound of Formula **VI** is converted to the corresponding compound of Formula **V** by the addition of N, N'-carbonyldiimidazole to a solution of **VI** in toluene under inert atmosphere (i.e., nitrogen) and stirred for a time period between about 30 minutes to about 4 hours, preferably about 2.5 hours.

In reaction 3 of Scheme 1, the compound of Formula **V** is converted to the corresponding compounds of Formula **II** and Formula **IV** by reacting **V** with imidazole in the presence of a aprotic solvent, such as toluene. The reaction mixture is heated to reflux for a time period between about 4 hours to about 28 hours, preferable about 6 hours.

In reaction 4 of Scheme 1, a mixture of the compounds of Formula **II** and Formula **IV** (or each intermediate separately) is converted to the corresponding compound of Formula **I** by reacting **II** and **IV** with (S)-(+)-quinuclidinol in the presence of a aprotic solvent, such as toluene. The reaction mixture is heated to reflux for a time period between about 12 hours to about 24 hours, preferable about 18 hours.

Preparation A

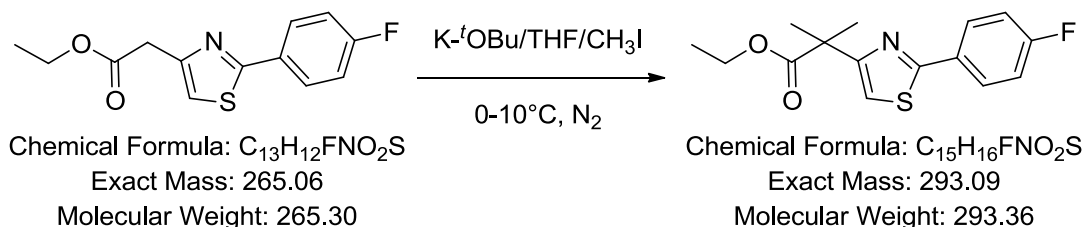
To 4-Fluorophenylthioamide (50.35g, 1 eq.) was added 8.6 weight volumes of 200 proof ethanol (based on thioamide) (430mL) and ethyl 4-chloroacetoacetate (68.2g, 1.1 eq.). The mixture was place under a nitrogen atmosphere. It was heated under reflux for 5h and allowed to cool to room temperature. The solution was concentrated to an oil and TBME (10 volumes, 500mL) and 6 volumes of saturated NaHCO₃ (300mL) added. The aqueous layer was back extracted with 5 volumes (250mL) of TBME. The combined organic layer was washed with water and then concentrated to an oil and then dried to a solid. The product was crystallized from 3 weight volumes of hot hexanes. Yield 89 % Product 98.7% pure by HPLC (area %).

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Example 1

(S)-Quinuclidin-3-yl (2-(2-(4-fluorophenyl)thiazol-4-yl)propan-2-yl)carbamate

Step 1: Dimethylation with methyl iodide



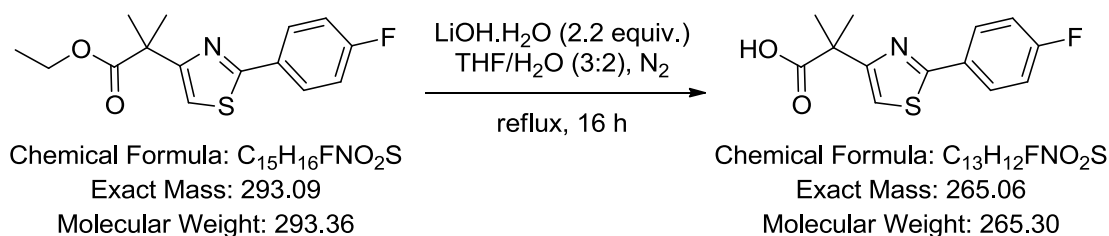
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Procedure: In a 100 L reactor was added tetrahydrofuran (THF, 28.4 Kg) and potassium tert-butoxide (MW 112.21, 2.28 Kg g, 4.0 equiv.). This mixture was cooled to 0-2°C (internal temperature). The starting ester (MW 265.3, 2.0 Kg, 1.0 equiv.) was dissolved in THF (4 L) and transferred to the reactor over a period of 10-60 min, keeping the internal temperature below 10°C during the addition. The reaction mixture was stirred at 3-9 °C for 15- 60 min. A solution of methyl iodide (MW 141.94, 1.88 L, 4.0 equiv.) in THF (4.8 L) was added to the reactor over 30-120 min keeping the internal temperature

20

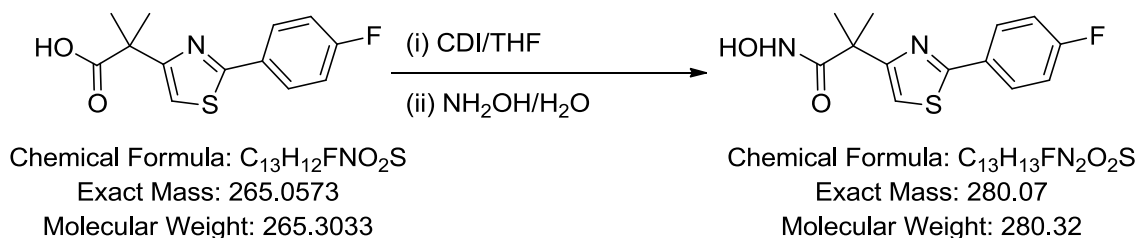
below 10°C. A solution of NaCl (2.0 Kg) in water (14L) was added over 10 min and the mixture was stirred for at least 10 min more. The reaction was made acidic by the addition of 1 M HCl (~ 1.44L). The layers were separated and the aqueous layer was back extracted with THF (6.2 kg). The combined organic layers were vacuum distilled to ~ 16 L. This THF solution of the Step1 product was used in the next reaction.

Step 2: Hydrolysis of the ethyl ester with LiOH monohydrate



Procedure: To the ester in THF was added a solution of LiOH.H₂O (MW 41.96, 0.695 Kg, 2.2 equiv.) in water (9.3 L) was added. The mixture was heated at reflux for 8-16 hours. After the reactions was judge complete by HPLC, water (12 L) was added and the mixture was vacuum distilled to ~ 16L. TBME (5.9 kg) was added and after stirring the layers were separated. The aqueous layer containing the product was washed a second time with TBME (5.9 Kg). TBME was added to the aqueous layer and the mixture was made acidic (pH ≤ 3) by the addition of 5 M HCl (~ 3.67 Kg). The layers were separated and the aqueous layer was extracted a second time with TBME (4.5 Kg). Heptane (15 Kg) was added to the combined organic layers and the mixture was vacuum distilled to ~ 16L. After heating and cooling to 5-25 °C and stirring for at least 3 h, the product was filtered, washed with heptane, and vacuum dried. Yield 85.8 % (2.15 Kg) HPLC purity (area %) 99.72%

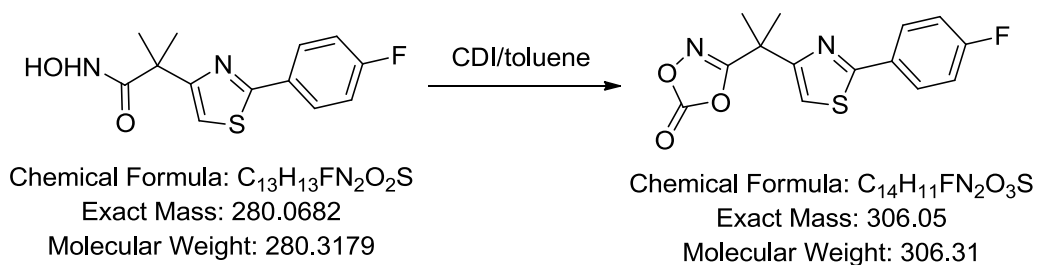
Reaction 1 : Formation of hydroxamic acid with NH₂OH



5

Procedure: To a 100 L reactor was added THF (14.2 Kg) and N, N'-carbonyldiimidazole (CDI; MW 162.15, 1.34 Kg, 1.1 equiv.). The acid from reaction 2 (2.0 Kg, 1.0 equiv) dissolved in THF (4L) was added over 15- 20 min. The mixture was stirred at room temperature for 2.5-3 h. The reaction was cooled to 0-3 °C. Aqueous hydroxylamine (50% aqueous; 1.7 L, 4.0 equiv.) was added over 5-15 min keeping the internal temperature less than 18 °C. After the addition was complete, the layers were separated and the organic layer was washed with water (12 Kg) and a solution of sodium chloride (2.0 Kg) in water (12L). The separated organic layer was vacuum distill to ~ 16 l. Toluene(13.8 Kg) was added and the mixture was again vacuum distilled to ~ 16L. Heptane (11 kg) was added and the mixture was stirred at room temperature for at least 16 h. The resulting solid was filtered , washed with heptane (11 Kg) and vacuum dried at room temperature. The yield was 1.58 Kg (74.8%).

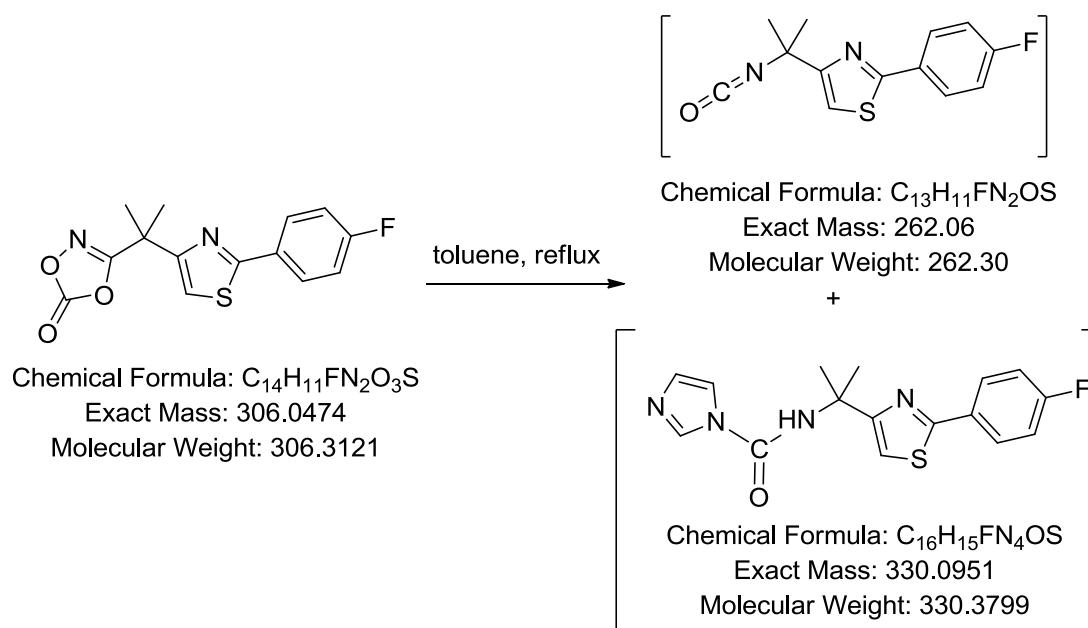
Reaction 2 : Conversion of hydroxamic acid to a dioxazolone



5 **Procedure:** Toluene (17.3 Kg) and the hydroxamic acid from, reaction 1 (MW 280.32, 2.0 Kg) was transferred to a 100 L reactor. After stirring at room temperature for at least 15 min carbonyl diimidazole CDI (MW 162.15, 1.27 Kg, 1.1 equiv.) was added. The mixture was stirred at room temperature for 1-4 h until the reaction was judge complete by HPLC.

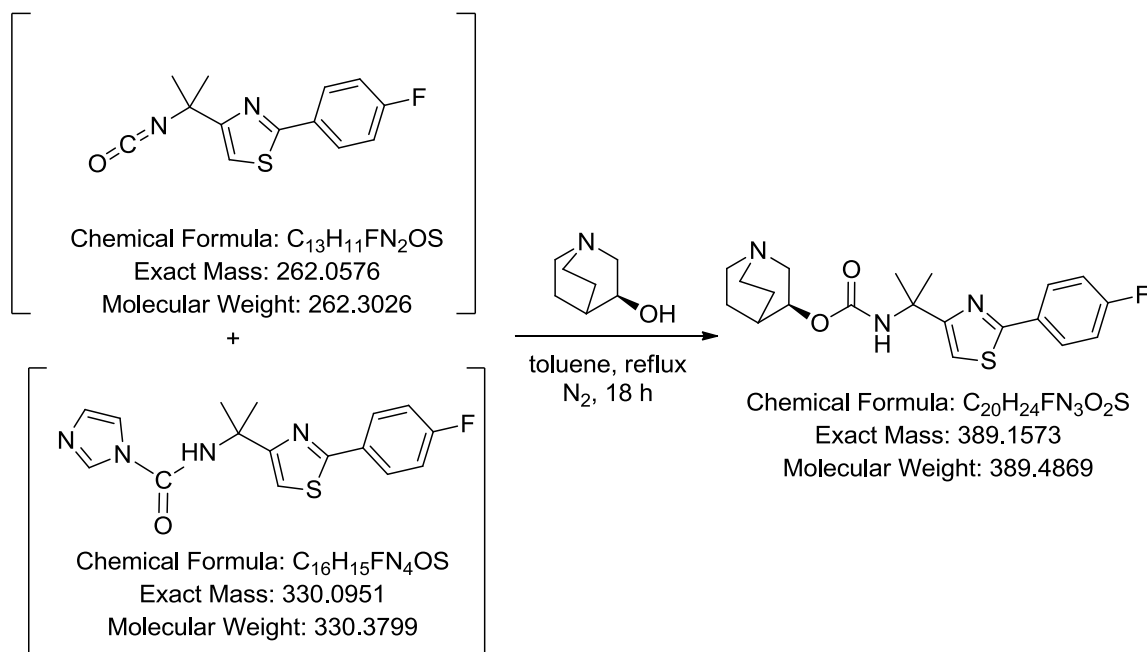
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Reaction 3 Conversion of the dioxazolone to a mixture of the imidazole urea and isocyanate



Procedure : The solution of the dioxazolone (reaction 2) was heated at 60 °C for 6-16 hours to complete the conversion to a mixture of the isocyanate and imidazole urea as judge by HPLC analysis.

Reaction 4 : Final conversion to the carbamate



5

Procedure: (S)-(+)-3- quinuclidinol (1.14 Kg , 1.18 equiv.) was added to the mixture of the isocyanate and imidazole urea toluene solution (reaction 3) and the solution was heated at 100-110 °C for 18-28 h. Toluene (8.6 Kg) was added to the reaction and the mixture was washed twice with water (20 Kg). The product was removed from the organic layer with two extractions of aqueous 1M HCl.(19.7 Kg). Isopropyl acetate (34.8 Kg) was added to the combined acidic aqueous layers. The mixture was cooled to 5-10 °C and 10M aqueous NaOH (5.3 Kg) was added. The layers were separated and the organic layer was vacuum distilled to ~ 16 L. Heptane (21.4 Kg) was added to the remaining isopropyl acetate solution and again the solution was distilled to ~ 16 L. the resulting suspension was stirred for at least 4 h. The product was filtered, washed with heptane (13.7 Kg) and vacuum dried at room temperature. The yield was 2.3 Kg (82.8 % yield). HPLC purity (Area %) 99.7 % .

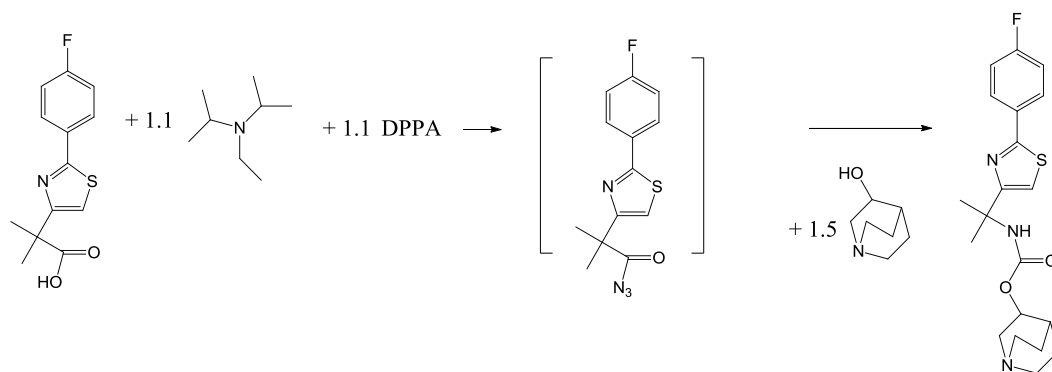
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¹H NMR (400 MHz, CDCl₃) δ 8.04 – 7.83 (m, 2H), 7.20 – 6.99 (m, 3H), 5.53 (s, 1H), 4.73 – 4.55 (m, 1H), 3.18 (dd, *J* = 14.5, 8.4 Hz, 1H), 3.05 – 2.19 (m, 5H), 2.0 – 1.76 (m, 11H). ¹³C NMR (100 MHz, CDCl₃) δ 166.38, 165.02, 162.54, 162.8-155.0 (d, C-F), 130.06, 128.43, 128.34, 116.01, 115.79, 112.46, 71.18, 55.70, 54.13, 47.42, 46.52, 27.94, 25.41, 24.67, 19.58.

Example 2

(S)-Quinuclidin-3-yl (2-(2-(4-fluorophenyl)thiazol-4-yl)propan-2-yl)carbamate

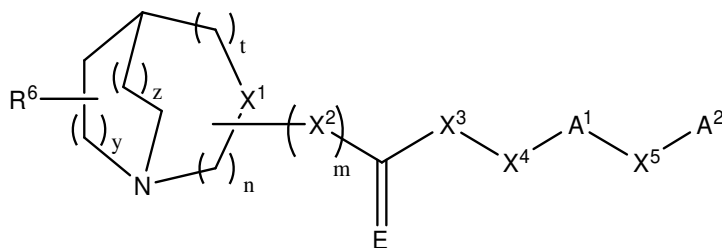


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15 2-(2-(4-fluorophenyl)thiazol-4-yl)-2-methylpropanoic acid (1g) and diisopropylethyl
amine (0.57 ml) were dissolved in toluene and stirred at 110 °C under N₂. DPPA (0.9 ml)
was added dropwise. The mixture was stirred for 3 hours at 110 °C to complete the
conversion of the acetyl azide and isocyanate. Quinuclidin-3-ol (0.72 g) was added and
20 stirred for 18 hours. The result mixture was diluted with toluene (50 ml) and washed with
saturated sodium bicarbonate solution. The organic layer was concentrated to oil. Product
of quinuclidin-3-yl (2-(2-(4-fluorophenyl)thiazol-4-yl)propan-2-yl)carbamate was
purified by crystallization from EtOAc (0.6 g). Claims

CLAIMS

1. A method of preparing a compound of the formula,



I

wherein:

n is 1, 2 or 3;

m is 1;

t is 0, 1 or 2;

y is 1 or 2;

z is 0, 1 or 2;

E is O;

X¹ is CR¹;

X² is O;

X³ is -NH;

X⁴ is CR⁴R⁵, CH₂ CR⁴R⁵ or CH₂ -(C₁-C₆) alkyl-CR⁴R⁵;

X⁵ is a direct bond;

R¹ is H;

R⁴ and R⁵ are independently selected from H or (C₁-C₆)alkyl; or

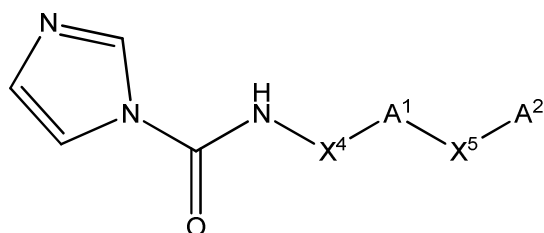
R⁴ and R⁵ may be taken together with the carbon to which they are attached to form a spiro (C₃-C₁₀)cycloalkyl ring or spiro (C₃-C₁₀)cycloalkoxy ring;

R⁶ is -H;

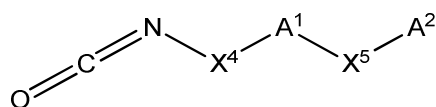
A¹ is (C₂-C₉)heteroaryl, optionally substituted with one or more substituents selected from the group consisting of halo, (C₁-C₆)alkyl optionally substituted by one to three halo; (C₁-C₆)alkenyl, amino, (C₁-C₆)alkylamino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, nitro, CN, -OH, (C₁-C₆)alkyloxy optionally substituted by one to three halo; (C₁-C₆)alkoxycarbonyl, and (C₁-C₆) alkylcarbonyl;

A² is (C₆-C₁₂)aryl, optionally substituted with one or more substituents selected from the group consisting of halo, (C₁-C₆)alkyl optionally substituted by one to three halo; (C₁-C₆)alkylenyl, amino, (C₁-C₆) alkylamino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, O(C₃-C₆ cycloalkyl), (C₃-C₆) cycloalkoxy, nitro, CN, OH, (C₁-C₆)alkyloxy optionally substituted by one to three halo; (C₃-C₆) cycloalkyl, (C₁-C₆) alkoxycarbonyl, (C₁-C₆) alkylcarbonyl, (C₁-C₆) haloalkyl;

with the proviso that the sum of n + t + y + z is not greater than 6;
comprising reacting the compounds of Formula II and Formula IV

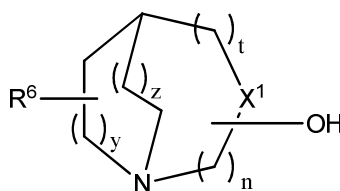


II



IV

with a compound of Formula III

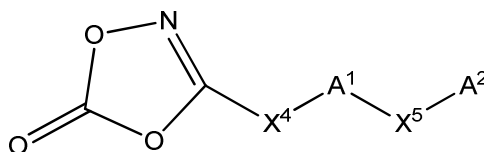


III

wherein n, t, y, z, X⁴, A¹, X⁵ and A² are as defined above.

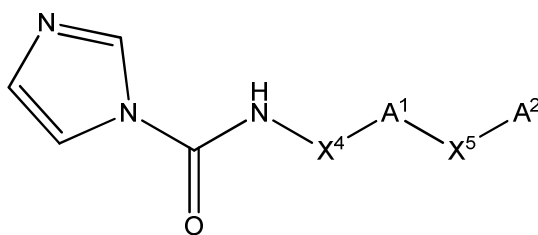
2. A method according to claim 1, wherein n is 1; t is 0; y is 1 and z is 1.
3. A method according to claim 2, wherein X⁴ is CR⁴R⁵.

4. A method according to claim 3, wherein R⁴ and R⁵ are each methyl.
5. A method according to claim 4, wherein A¹ is thiophene, thiazole, isothiazole, furane, oxazole, isoxazole, pyrrole, imidazole, pyrazole, triazole, pyridine, pymiridine, pyridazine, indole, benzothiazole, benzoisoxazole, benzopyrazole, benzoimidazole, benzofuran or benzooxazole.
6. A method according to claim 5, wherein A¹ is thiazole.
7. A method according to claim 6, wherein A² is phenyl.
8. A method according to claim 7, wherein the phenyl group is substituted by halo.
9. A method according to claim 8, wherein the halo group is fluoro.
10. A method according to claim 1, further including reacting, while heating to reflux, the compound of Formula V

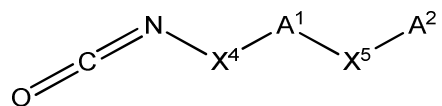


V

with imidazole to form the compounds of Formula II and Formula IV



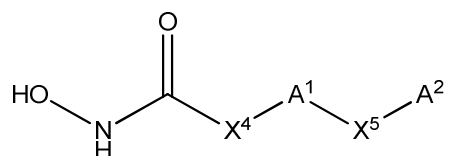
II



IV

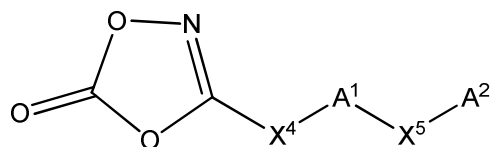
wherein X^4 , A^1 , X^5 and A^2 are as defined above.

11. A method according to claim 10, further including reacting the compound of Formula VI



VI

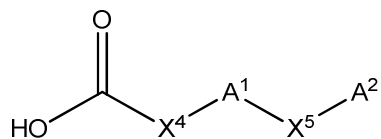
with N, N'- carbonyldiimidazole to form the compound of Formula V



V

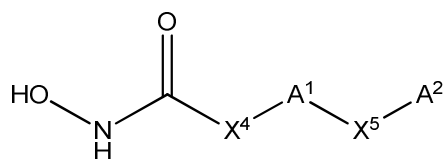
wherein X^4 , A^1 , X^5 and A^2 are as defined above.

12. A method according to claim 11, further including reacting the compound of Formula VII



VII

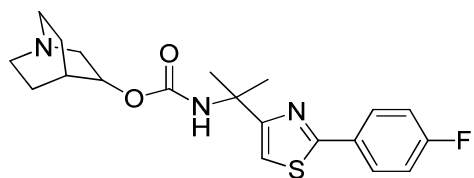
with N, N'- carbonyldiimidazole and hydroxylamine to form the compound of Formula VI



VI

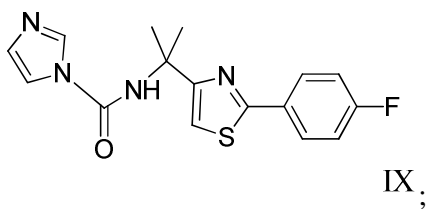
wherein X^4 , A^1 , X^5 and A^2 are as defined above.

13. A method of preparing the compound of Formula VIII

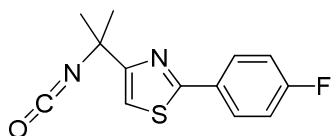


VIII

comprising reacting compounds of Formula IX and Formula X



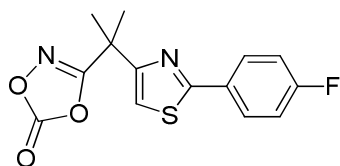
IX;



X

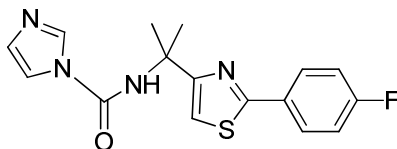
with quinuclidinol.

14. A method according to claim 13, further including reacting, while heating to reflux, the compound of Formula XI

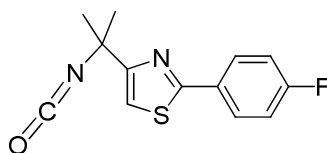


XI

with imidazole to form the compounds of Formula IX and Formula X

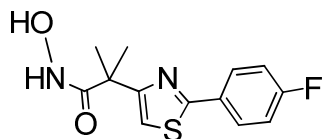


IX;



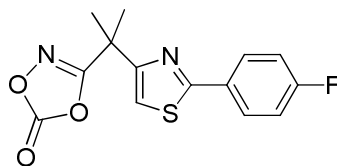
X.

15. A method according to claim 14, further including reacting the compound of Formula XII



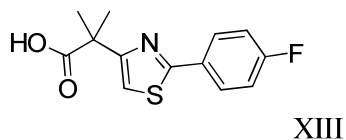
XII

with N, N'- carbonyldiimidazole to form the compound of Formula XI

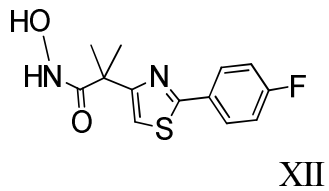


XI.

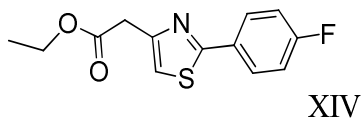
16. A method according to claim 15, further including reacting the compound of Formula XIII



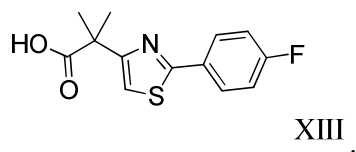
with N, N'- carbonyldiimidazole and hydroxylamine to form the compound of Formula XII



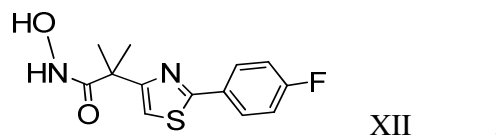
17. A method according to claim 16, further including reacting the compound of Formula XIV



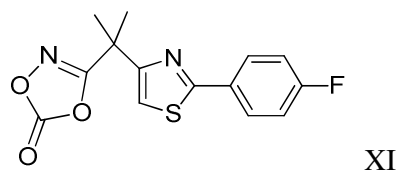
with potassium tert-butoxide and methyl iodine followed by reacting the ethyl ester so formed with lithium hydroxide to form the compound of Formula XIII



18. A compound of Formula XII



19. A compound of Formula XI



20. A compound of Formula IX

