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(12) **United States Patent**
Nakajima et al.(10) **Patent No.:** **US 9,585,889 B2**(45) **Date of Patent:** **Mar. 7, 2017**

- (54) **NITROGEN-CONTAINING BICYCLIC AROMATIC HETEROCYCLIC COMPOUND**
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(51) **Int. Cl.**

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CPC **A61K 31/519** (2013.01); **A61K 31/52** (2013.01); **A61K 31/5377** (2013.01); **C07D 473/00** (2013.01); **C07D 487/04** (2013.01)

(58) **Field of Classification Search**

None
See application file for complete search history.

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Primary Examiner — Golam M M Shameem*Assistant Examiner* — Laura Daniel(74) *Attorney, Agent, or Firm* — Oblon, McClelland, Maier & Neustadt, L.L.P.(57) **ABSTRACT**

On the basis of a cathepsin S inhibitory effect, an excellent agent for treating or preventing autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, systemic lupus erythematosus, or the like is provided. It was found that a nitrogen-containing bicyclic heterocyclic compound has an excellent cathepsin S inhibitory effect, thereby completing the invention. The compound of the present invention has an cathepsin S inhibitory effect, and can be used as an agent for preventing and/or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, systemic lupus erythematosus, or the like.

3 Claims, No Drawings

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**NITROGEN-CONTAINING BICYCLIC
AROMATIC HETEROCYCLIC COMPOUND**

CROSS REFERENCES TO RELATED
APPLICATIONS

This application is a Continuation Application of U.S. patent application Ser. No. 14/394,373, filed on Oct. 14, 2014, which was a 371 of International Patent Application No. PCT/JP2013/061348, filed on Apr. 17, 2013, and claims priority to Japanese Patent Application No. 2012-094278, filed on Apr. 17, 2012, and Japanese Patent Application No. 2013-051487, filed on Mar. 14, 2013.

TECHNICAL FIELD

The present invention relates to a pharmaceutical composition, for example, a nitrogen-containing bicyclic aromatic heterocyclic compound useful as an active ingredient of a pharmaceutical composition for preventing and/or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, or systemic lupus erythematosus (SLE).

BACKGROUND ART

Cathepsin S is mainly a lysosomal cysteine protease that is expressed on antigen-presenting cells such as dendritic cells, macrophages, and B-cells, and plays a role in the degradation of an invariant chain bonded when major histocompatibility gene complex class II (MHC class II) molecules are generated. MHC class II molecules are bonded to self-peptide or non-self peptide taken from the outside of a cell, and by presenting the non-self peptide or pathogenic self-peptide to CD4 positive T cells, secretion of various cytokines is induced. It is confirmed that by inhibiting cathepsin S or making cathepsin S deficient, loading of antigenic peptide to MHC class II molecules are inhibited, and antigen presentation to CD4 positive T cells is suppressed, and by these, an immune response with respect to a foreign antigen is decreased (Nakagawa et al, Immunity, 1999, Vol. 10, No. 2, pp. 207-217).

In this respect, an inhibitor of cathepsin S can be useful as an agent for preventing and/or treating inflammation and immune disease such as rheumatoid arthritis, multiple sclerosis, SLE, psoriasis and Crohn's disease, and as an agent for preventing and/or treating graft rejection.

It have been reported that the compounds represented by the following formulas (A) (Patent Documents 1 and 2), (B) (Patent Document 3), (C) (Patent Document 4), and (D) (Patent Document 5) exhibit cathepsin S inhibition, and are useful as an agent for treating inflammations such as rheumatism-immune disorders and the like. It have been reported that the compound represented by the following formula (E) exhibits cathepsin S inhibition, and is useful as an agent for treating inflammations such as rheumatism-immune disorders and the like (Patent Document 6).

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[Chem. 1]

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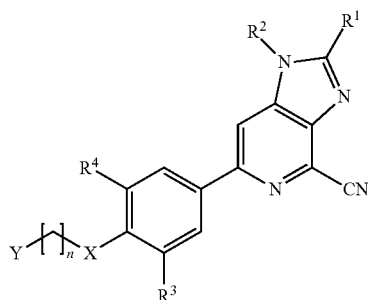
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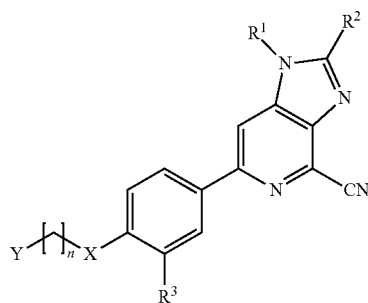
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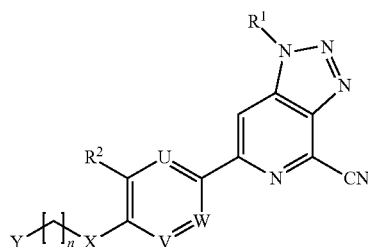
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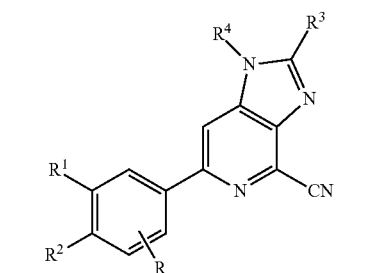
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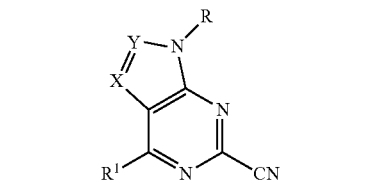
(B)



(C)



(D)



(E)

(Refer to this publication for the symbols in the formulas (A) to (D). In the formula (E), R is aryl or hetero aryl which may be substituted with a halogen or the like, H, C₁₋₆ alkyl, or C₃₋₆ cycloalkyl, Y is N, CH, or the like, R¹ is Y(CH₂)_pR⁷, Y in Y(CH₂)_pR⁷ is O or NR⁸, and R⁷ is a 5 to 6-membered saturated heterocycle.)

[Patent Document 1] Pamphlet of International Publication No. 2009/010491

[Patent Document 2] Pamphlet of U.S. Patent Application Publication No. 2009/0099172

[Patent Document 3] Pamphlet of International Publication No. 2010/081859

[Patent Document 4] Pamphlet of International Publication No. 2011/086125

[Patent Document 5] Pamphlet of International Publication No. 2007/080191

[Patent Document 6] Pamphlet of International Publication No. 2004/000843

DISCLOSURE OF INVENTION

Problems to Be Solved by the Invention

A pharmaceutical composition, for example, a nitrogen-containing bicyclic aromatic heterocyclic compound useful as an active ingredient of a pharmaceutical composition for preventing and/or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, or SLE is provided.

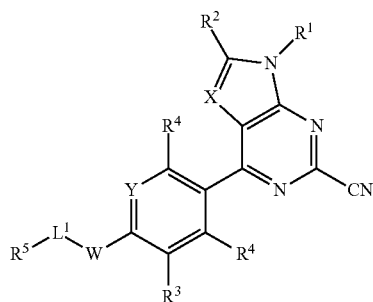
Means for Solving the Problems

As a result of intensive studies on a compound having the cathepsin S inhibitory effect, the present inventors found that a nitrogen-containing bicyclic aromatic heterocyclic compound has the cathepsin S inhibitory effect, thereby completing the invention.

That is, the present invention relates to the compound of the formula (I) or a salt thereof, and a pharmaceutical composition containing the compound of the formula (I) or a salt thereof, and a pharmaceutically acceptable excipient.

[1] A compound of the formula (I) or a salt thereof:

[Chem. 2]



(In the formula,

X is CH or N;

Y is CH, C-halogen, or N;

W is —O— or —S(O)_n—;

L¹ is a bond, -lower alkylene-, —O-lower alkylene-, —NH-lower alkylene-, or —C(O)-lower alkylene-;

R¹ is i) lower alkyl which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —O-lower alkyl, —NH₂, —NH-(lower alkyl), —N(lower alkyl)₂, —C(O)—NH₂, —C(O)—NH-

lower alkyl, —C(O)—N(lower alkyl)₂, —C(O)—NH—nonaromatic heterocycle which may be substituted, —C(O)—N(lower alkyl)-nonaromatic heterocycle which may be substituted, a nonaromatic heterocycle which may be substituted, and —C(O)-nonaromatic heterocycle which may be substituted;

ii) nonaromatic heterocycle which may be substituted; or iii) H;

R² is lower alkyl which may be substituted with a halogen, a halogen, or H;

R³ is lower alkyl which may be substituted with a halogen, a halogen, or —CN;

R⁴'s are the same as or different from each other and R⁴ is lower alkyl which may be substituted with a halogen, a halogen, —OH, —CN, or H.

R⁵ is lower alkyl which may be substituted with a substituent selected from the group consisting of —OH, a halogen, —NH₂, —NH-(lower alkyl), and —N(lower alkyl)₂, —O-(lower alkyl which may be substituted with a halogen), C₃₋₈ cycloalkyl which may be substituted, or a nonaromatic heterocycle which may be substituted; and n(s) is(are) the same as or different from each other, and n is an integer of 0 to 2.)

[2] The compound or a salt thereof according to [1], wherein X is N;

R¹ is lower alkyl which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —O-lower alkyl, and a nonaromatic heterocycle, in which the nonaromatic heterocycle may be substituted with lower alkyl;

R⁵ is a nonaromatic heterocycle which may be substituted with a substituent selected from a D2 group; in which the D2 group is

(1) a halogen;

(2) cycloalkyl which may be substituted with lower alkyl;

(3) an aromatic heterocycle which may be substituted with lower alkyl;

(4) a nonaromatic heterocycle which may be substituted with lower alkyl;

(5) —OH, —CN, or —NO₂;

(6) —C(O)—N(R⁰)₂ or —C(O)—N(lower alkyl)-nonaromatic heterocycle; and

(7) —C(O)-lower alkylene-OH;

—C(O)-(lower alkyl which may be substituted with one or more substituents selected from Z² or nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —CN, —O-lower alkyl, cycloalkyl, and a nonaromatic heterocycle); or

—C(O)-lower alkylene-(lower alkyl which may be substituted with one or more substituents selected from Z² or nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —CN, —O-lower alkyl, cycloalkyl, and a nonaromatic heterocycle); and

(8) lower alkyl or —O-lower alkyl which may be respectively substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (7);

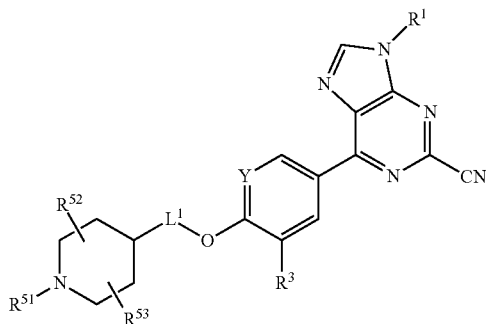
R⁰'s are the same as or different from each other, and R⁰ is H or lower alkyl; and

Z² is the group consisting of —OH—, —O-lower alkyl, and a halogen.

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- [3] The compound or a salt thereof according to [2], wherein W is —O— or —S—;
 L¹ is -lower alkylene-, —O-lower alkylene-, —NH-lower alkylene-, or —C(O)-lower alkylene-;
 R¹ is lower alkyl, and the lower alkyl may be substituted with tetrahydrofuranyl;
 R² is H;
 R³ is a halogeno-lower alkyl or —CN;
 R⁴ is H; and
 R⁵ is a nitrogen-containing heterocycloalkyl which may be substituted with a substituent selected from the D2 group.
- [4] The compound or a salt thereof according to [3], wherein W is —O—.
- [5] The compound or a salt thereof according to [4], wherein R⁵ is a group selected from the group consisting of azetidiny, pyrrolidinyl, piperidinyl, and octahydroindoliziny; in which each of azetidiny, pyrrolidinyl, piperidinyl, and octahydroindoliziny may be substituted with a substituent selected from the D2 group.
- [6] The compound or a salt thereof according to [5], which is represented by the formula (II):

[Chem. 3]



- (In the formula,
 R⁵¹ is H, lower alkyl, halogeno-lower alkyl, cycloalkyl, -lower alkylene-OH, -lower alkylene-CN, -lower alkylene-C(O)—N(R⁰)₂, -lower alkylene-C(O)-cyclic amino, —C(O)— heterocycloalkyl, —C(O)—N(R⁰)₂, —C(O)-lower alkylene-OH, or -lower alkylene-(aromatic heterocycle which may be substituted with lower alkyl); in which the heterocycloalkyl may be substituted with lower alkyl, —OH, or heterocycloalkyl;
 the cyclic amino may be substituted with a halogen or —CN; and
 R⁵² and R⁵³ are the same as or different from each other, and each of R⁵² and R⁵³ is H or lower alkyl.)
- [7] The compound or a salt thereof according to [6], wherein R¹ is a methyl.
- [8] The compound or a salt thereof according to [7], wherein L¹ is -lower alkylene-.
- [9] The compound or a salt thereof according to [8], wherein Y is N.
- [10] The compound or a salt thereof according to [9], wherein R³ is a trifluoromethyl.
- [11] The compound or a salt thereof according to [10],

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- wherein R⁵¹ is H, lower alkyl, -lower alkylene-C(O)—N(R⁰)₂, or —C(O)-(heterocycloalkyl which may be substituted with lower alkyl); and each of R⁵² and R⁵³ is H.
- [12] The compound or a salt thereof according to [11], wherein R⁵¹ is lower alkyl.
- [13] The compound or a salt thereof according to [6], which is selected from the following group consisting of:
 9-methyl-6-{6-[2-(piperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-9H-purine-2-carbonitrile;
 6-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-9-methyl-9H-purine-2-carbonitrile;
 9-methyl-6-[6-[2-[1-(1-methyl-L-prolyl)piperidin-4-yl]ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-9H-purine-2-carbonitrile;
 6-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl}-9-(tetrahydrofuran-3-ylmethyl)-9H-purine-2-carbonitrile; and
 6-{6-[2-(1-cyclobutylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-9-methyl-9H-purine-2-carbonitrile.
- [14] A pharmaceutical composition, containing:
 the compound or a salt thereof according to [1]; and
 a pharmaceutically acceptable excipient.
- [15] A pharmaceutical composition for preventing or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, SLE, or lupus nephritis, comprising:
 the compound or a salt thereof according to [1].
- [16] Use of the compound or a salt thereof according to [1] for the manufacture of a pharmaceutical composition for preventing or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, SLE, or lupus nephritis.
- [17] Use of the compound or a salt thereof according to [1] for preventing or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, SLE, or lupus nephritis.
- [18] The compound or a salt thereof according to [1] for preventing or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, SLE, or lupus nephritis.
- [19] A method for preventing or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, SLE, or lupus nephritis, comprising:
 administering to a subject an effective amount of the compound or a salt thereof according to [1].
- In addition, the present invention relates to a pharmaceutical composition for preventing and/or treating autoimmune disease as an aspect, allergic disease as another aspect, graft rejection of an organ, bone marrow or tissue as still another aspect, SLE as still another aspect, or lupus nephritis as further still another aspect, containing a compound of the formula (I) or a salt thereof. Moreover, the pharmaceutical composition contains an agent for preventing and/or treating autoimmune disease as an aspect, allergic disease as another aspect, graft rejection of an organ, bone marrow or tissue as still another aspect, SLE as still another aspect, or lupus nephritis as further still another aspect, containing a compound of the formula (I) or a salt thereof.
- In addition, the present invention relates to use of the compound of the formula (I) or a salt thereof for the manufacture of a pharmaceutical composition for preventing and/or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, SLE, or lupus nephritis, the compound of the formula (I) or a salt thereof for use in preventing and/or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or

tissue, SLE, or lupus nephritis, and a method for preventing and/or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, SLE, or lupus nephritis, comprising administering to a subject an effective amount of the compound of the formula (I) or a salt thereof.

Moreover, the "subject" is a human or animals in need of such prevention or treatment, and in an embodiment, the subject is a human in need of such prevention or treatment.

Moreover, unless otherwise described, in a case where the symbols in a chemical formula in the specification is also used in other chemical formulas, the same symbol indicates the same meaning.

Effects of the Invention

Moreover, the compound of the formula (I) or a salt thereof has a cathepsin S inhibitory effect, and can be used as an agent for preventing and/or treating autoimmune disease as an aspect, allergic disease as another aspect, graft rejection of an organ, bone marrow or tissue as still another aspect, SLE as still another aspect, or lupus nephritis as further still another aspect.

EMBODIMENTS FOR CARRYING OUT THE INVENTION

Hereinafter, the present invention will be described in detail.

In the present specification, "lower alkyl" is a linear or branched alkyl having 1 to 6 carbon atoms (hereinafter, referred to as C₁₋₆), and examples thereof include methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, n-pentyl and n-hexyl. The lower alkyl is C₁₋₄ alkyl as another aspect, is methyl, ethyl, isopropyl, isobutyl or tert-butyl as still another aspect, is methyl or ethyl as still another aspect, is methyl as still another aspect, and is ethyl as further still another aspect.

"Lower alkylene" is a linear or branched C₁₋₆ alkylene, and examples thereof include methylene, ethylene, trimethylene, tetramethylene, pentamethylene, hexamethylene, propylene, methylmethylene, ethylethylene, 1,2-dimethylethylene, 2,2-dimethylethylene and 1,1,2,2-tetramethylethylene. The lower alkylene is C₁₋₄alkylene as another aspect, is methylene, ethylene or trimethylene as still another aspect, is methylene or ethylene as still another aspect, is methylene as still another aspect, and is ethylene as further still another aspect.

"Halogen" is F, Cl, Br or I. The halogen is F as another aspect, and is Cl as still another aspect.

"Halogeno-lower alkyl" is lower alkyl substituted with one or more halogens. As another aspect, the halogeno-lower alkyl is lower alkyl substituted with 1 to 5 halogens, and as still another aspect, the halogeno-lower alkyl is trifluoromethyl.

"Cycloalkyl" is a C₃₋₁₀ saturated hydrocarbon cyclic group, which may have a bridge. Examples thereof include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl and adamantyl. The cycloalkyl is C₃₋₈ cycloalkyl as another aspect, is C₃₋₆ cycloalkyl as still another aspect, is cyclopropyl as still another aspect, is cyclobutyl as still another aspect, and is adamantyl as further still another aspect.

An "aromatic heterocyclic" group means a cyclic group selected from (i) 5 to 6-membered monocyclic aromatic heterocycle containing 1 to 4 hetero atoms selected from oxygen, sulfur and nitrogen, and (ii) a bicyclic or tricyclic heterocycle containing 1 to 5 hetero atoms selected from

oxygen, sulfur and nitrogen, which is formed by the monocyclic aromatic heterocycle being fused with one or two rings selected from the group consisting of a monocyclic aromatic heterocycle and a benzene ring. Oxide or dioxide may be formed by the oxidation of sulfur or nitrogen which is a ring atom. Specific examples thereof include pyridyl, pyrrolyl, pyrazinyl, pyrimidinyl, pyridazinyl, imidazolyl, triazolyl, triazinyl, tetrazolyl, thiazolyl, pyrazolyl, isothiazolyl, oxazolyl, isoxazolyl, thiadiazolyl, oxadiazolyl, thienyl, furyl, indolyl, isoindolyl, benzimidazolyl, indazolyl, quinolyl, isoquinolyl, quinazolyl, quinoxalyl, phthalazinyl, benzothiazolyl, benzisothiazolyl, benzothiadiazolyl, benzoxazolyl, benzisoxazolyl, benzofuranyl, benzothienyl, benzotriazolyl, carbazolyl, imidazopyridyl, imidazopyrimidinyl, imidazopyridazinyl, pyrazolopyrimidinyl and 1,8-naphthyridinyl. The aromatic heterocycle group is a 5 to 6-membered monocyclic aromatic heterocyclic group as another aspect, is pyridyl as still another aspect, and is imidazolyl as further still another aspect.

A "nonaromatic heterocyclic" group means a 3 to 15-membered monocyclic to tricyclic saturated heterocyclic group containing 1 to 4 hetero atoms selected from oxygen, sulfur and nitrogen, and as another aspect, a 4 to 10-membered monocyclic to tricyclic saturated heterocyclic group. Oxide or dioxide may be formed by the oxidation of sulfur or nitrogen which is a ring atom. A part of bond of a ring may be bridged. Specific examples thereof include azetidyl, pyrrolidinyl, piperidinyl, piperazinyl, azepanyl, morpholinyl, thiomorpholinyl, tetrahydropyridinyl, tetrahydropyrimidinyl, oxetanyl, tetrahydrofuranyl, tetrahydropyranyl, dioxolanyl, dioxanyl, tetrahydrothiopyranyl, azabicyclo[3.2.1]octyl and octahydroindolizyl. The nonaromatic heterocyclic group is a 4 to 10-membered monocyclic to tricyclic saturated heterocyclic group as another aspect, is a 4 to 8-membered monocyclic saturated heterocyclic group as still another aspect, is pyrrolidinyl, piperidinyl or piperazinyl as still another aspect, and is piperidinyl as further still another aspect.

A "heterocycloalkyl" group means a heterocyclic group in which bonds between ring atoms in the above-described "nonaromatic heterocyclic" group are only single bonds, and carbon atoms constituting the ring have bonding sites. Specific examples thereof include azetidin-3-yl, pyrrolidin-2-yl, pyrrolidin-3-yl, piperidin-3-yl, piperidin-4-yl, 3-azabicyclo[3.2.1]oct-8-yl, 8-azabicyclo[3.2.1]oct-3-yl, octahydroindolizin-7-yl, oxetane-3-yl, tetrahydrofuran-2-yl, tetrahydrofuran-3-yl, tetrahydropyran-3-yl, and tetrahydropyran-4-yl. As another aspect, the heterocycloalkyl group is oxetan-3-yl, tetrahydrofuran-2-yl, tetrahydrofuran-3-yl, tetrahydropyran-3-yl, or tetrahydropyran-4-yl.

A "nitrogen-containing heterocycloalkyl" group means a group having a nitrogen atom in the above-described "heterocycloalkyl", in which carbon atoms constituting the ring have bonding sites. Specific examples thereof include azetid-3-yl, pyrrolidin-2-yl, pyrrolidin-3-yl, piperidin-3-yl, piperidin-4-yl, 3-azabicyclo[3.2.1]oct-8-yl, 8-azabicyclo[3.2.1]oct-3-yl, and octahydroindolizin-7-yl. The nitrogen-containing heterocycloalkyl group is pyrrolidin-3-yl, piperidin-3-yl, piperidin-4-yl or octahydroindolizin-7-yl as another aspect, is pyrrolidin-2-yl or pyrrolidin-3-yl as still another aspect, is piperidin-4-yl as still another aspect, and is octahydroindolizin-7-yl as further still another aspect.

A "cyclic amino" group means a group having a nitrogen atom in the above-described "nonaromatic heterocycle", in which nitrogen atoms constituting the ring have bonding sites. Specific examples thereof include azetid-1-yl, pyrrolidin-1-yl, piperidin-1-yl, azepan-1-yl, oxazolidin-3-yl,

thiazolidin-3-yl, isoxazolidin-2-yl, isothiazolidin-2-yl, piperazin-1-yl, morpholin-4-yl, thiomorpholin-4-yl, 1,1-dioxidethiomorpholin-4-yl, indolin-1-yl, isoindolin-2-yl, and 1,2,3,4-tetrahydroquinolin-1-yl. As another aspect, the cyclic amino group is azetidin-1-yl, pyrrolidin-1-yl, piperidin-1-yl, piperazin-1-yl or morpholin-4-yl, and as still another aspect, is piperidin-1-yl.

In the present specification, for example, in a case where L^1 is —O-lower alkylene-, O is bonded to R^5 , and lower alkylene is bonded to W.

In the present specification, “may be substituted” means that it does not have any substituent, or has 1 to 5 substituents. Moreover, in a case of having a plurality of substituents, these substituents may be the same as or different from each other.

As acceptable substituents of “—C(O)—NH-nonaromatic heterocycle which may be substituted, —C(O)—N(lower alkyl)-nonaromatic heterocycle which may be substituted, a nonaromatic heterocycle which may be substituted, and —C(O)-nonaromatic heterocycle which may be substituted” in R^1 in the formula (I), or “a nonaromatic heterocycle” in “a nonaromatic heterocycle which may be substituted”, groups shown as the following D1 group can be exemplified.

The D1 group is

- (1) a halogen,
- (2) —OH, —CN, or —NO₂,
- (3) nonaromatic heterocycle, and
- (4) lower alkyl or —O-lower alkyl which may be respectively substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (3).

Another aspect of the D1 group is

- (1) a halogen, and
- (2) lower alkyl.

As acceptable substituents of “nonaromatic heterocycle which may be substituted” described in R^5 in the formula (I), groups shown as the following D2 group can be exemplified.

The D2 group is

- (1) a halogen,
- (2) cycloalkyl which may be substituted with lower alkyl,
- (3) an aromatic heterocycle which may be substituted with lower alkyl,
- (4) a nonaromatic heterocycle which may be substituted with lower alkyl,
- (5) —OH, —CN, or —NO₂,
- (6) —C(O)—N(R^0)₂ or —C(O)—N(lower alkyl)-nonaromatic heterocycle, and
- (7) —C(O)-lower alkylene-OH, —C(O)-(lower alkyl which may be substituted with one or more substituents selected from Z^2 or nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —CN, —O-lower alkyl, cycloalkyl and a nonaromatic heterocycle), or

—C(O)-lower alkylene-(lower alkyl which may be substituted with one or more substituents selected from Z^2 or nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —CN, —O-lower alkyl, cycloalkyl and a nonaromatic heterocycle), and

- (8) lower alkyl or —O-lower alkyl which may be respectively substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (7).

Here, R^0 's are the same as or different from each other, and R^0 is H or lower alkyl, and Z^2 is a group consisting of —OH—, —O-lower alkyl, and a halogen.

Another aspect of the D2 group is

- (1) a halogen,
- (2) cycloalkyl which may be substituted with lower alkyl,
- (3) an aromatic heterocycle which may be substituted with lower alkyl,
- (4) a nonaromatic heterocycle which may be substituted with lower alkyl,
- (5) —OH, —CN, or —NO₂,
- (6) —C(O)—NH₂, —C(O)—N(lower alkyl)₂, or —C(O)—N(lower alkyl)-nonaromatic heterocycle, and
- (7) —C(O)-lower alkylene-OH, —C(O)-(lower alkyl which may be substituted with one or more substituents selected from Z^2 or nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —CN, —O-lower alkyl, and cycloalkyl), or —C(O)-lower alkylene-(lower alkyl which may be substituted with one or more substituents selected from Z^2 or nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —CN, —O-lower alkyl, and cycloalkyl), and
- (8) lower alkyl or —O-lower alkyl which may be respectively substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (7).

Another aspect of the D2 group is (1) a halogen,

- (2) cycloalkyl,
- (3) an aromatic heterocycle which may be substituted with lower alkyl,
- (4) nonaromatic heterocycle,
- (5) oxo,
- (6) —OH,
- (7) —CN,
- (8) —C(O)—N(lower alkyl)₂,
- (9) —C(O)-lower alkylene-OH, and
- (10) —C(O)-(nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of lower alkyl, a halogen, —OH and —CN), and
- (11) lower alkyl which may be substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (10).

Another aspect of the D2 group is

- (1) a halogen,
- (2) cycloalkyl,
- (3) an aromatic heterocycle which may be substituted with lower alkyl,
- (4) nonaromatic heterocycle,
- (5) —OH,
- (6) —CN,
- (7) —C(O)—N(lower alkyl)₂,
- (8) —C(O)-lower alkylene-OH, and
- (9) —C(O)-(nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of lower alkyl, a halogen, —OH and —CN), and
- (10) lower alkyl which may be substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (9).

Another aspect of the D2 group is

- (1) cycloalkyl,
- (2) —OH,
- (3) —C(O)—N(lower alkyl)₂, and
- (4) —C(O)-(nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of lower alkyl, a halogen, —OH and —CN), and

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(5) lower alkyl which may be substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (4).

Another aspect of the D2 group is

- (1) —OH,
- (2) —C(O)—N(lower alkyl)₂, and
- (3) —C(O)-(nonaromatic heterocycle which may be substituted with one or more substituents selected from the group consisting of lower alkyl and —OH), and
- (4) lower alkyl which may be substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (3).

Another aspect of the D2 group is

- (1) lower alkyl,
- (2) -lower alkylene-OH,
- (3) -lower alkylene-C(O)—N(lower alkyl)₂, and
- (4) —C(O)-(nonaromatic heterocycle which may be substituted with lower alkyl).

As acceptable substituents of “C₃₋₈ cycloalkyl which may be substituted” or “aromatic heterocycle which may be substituted” in R⁵ in the formula (I), groups shown as the following D3 group can be exemplified.

The D3 group is

- (1) a halogen,
- (2) —OH, —CN, or —NO₂, and
- (3) lower alkyl or —O-lower alkyl which may be respectively substituted with one or more substituents selected from the group consisting of substituents according to the above-described (1) to (2).

Another aspect of the D3 group is

- (1) lower alkyl, and
- (2) a halogen.

Aspect of the present invention is shown below. “A compound” in this section means “a compound or a salt thereof”.

(1) A compound of the formula (I), in which X is CH. As another aspect, a compound in which X is N.

(2) A compound of the formula (I), in which Y is CH. As another aspect, a compound in which Y is N.

(3) A compound of the formula (I), in which W is —O— or —S—. As another aspect, a compound in which W is —O—.

(4) A compound of the formula (I), in which L¹ is -lower alkylene-, —O-lower alkylene-, or —C(O)-lower alkylene.

As another aspect, a compound in which L¹ is -lower alkylene- or —O-lower alkylene-.

As still another aspect, a compound in which L¹ is -lower alkylene-.

(5) A compound of the formula (I), in which R¹ is lower alkyl which may be substituted with a substituent selected from the group consisting of —OH, —O-lower alkyl, and a nonaromatic heterocycle, in which the nonaromatic heterocycle may be substituted with lower alkyl.

As another aspect, a compound in which R¹ is lower alkyl which may be substituted with —OH or —O-lower alkyl.

As still another aspect, a compound in which R¹ is lower alkyl which may be substituted with tetrahydrofuran.

As still another aspect, a compound in which R¹ is lower alkyl.

As further still another aspect, a compound in which R¹ is methyl.

(6) A compound of the formula (I), in which R² is lower alkylene which may be substituted with a halogen or H.

As another aspect, a compound in which R² is lower alkyl which may be substituted with a halogen.

As still another aspect, a compound in which R² is H.

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(7) A compound of the formula (I), in which R³ is halogeno-lower alkyl or —CN.

As another aspect, a compound in which R³ is halogeno-lower alkyl.

5 As still another aspect, a compound in which R³ is trifluoromethyl.

(8) A compound of the formula (I), in which R⁴⁺ are the same as or different from each other, and R⁴ is H or a halogen. As another aspect, a compound in which all R⁴ are H's.

10 (9) A compound of the formula (I), in which R⁵ is lower alkyl which may be substituted with —N(lower alkyl)₂, an aromatic heterocycle which may be substituted, or a non-aromatic heterocycle which may be substituted.

15 As another aspect, a compound in which R⁵ is lower alkyl which may be substituted with —N(lower alkyl)₂, or a nonaromatic heterocycle which may be substituted.

As still another aspect, a compound in which R⁵ is a nonaromatic heterocycle which may be substituted.

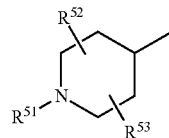
As still another aspect, a compound in which R⁵ is nitrogen-containing heterocycloalkyl which may be substituted.

25 As still another aspect, a compound in which R⁵ is azetidiny which may be substituted, pyrrolidiny which may be substituted, piperidiny which may be substituted, or octahydroindoliziny which may be substituted.

As still another aspect, a compound in which R⁵ is piperidiny which may be substituted.

30 (10) A compound of the formula (I), in which R⁵ is

[Chem. 4]



(10-1) A compound according to (10), in which R⁵¹ is H, lower alkyl, halogeno-lower alkyl, cycloalkyl, -lower alkylene-OH, or -lower alkylene-CN.

As another aspect, a compound according to (10), in which R⁵¹ is H, -lower alkylene-C(O)—N(R⁰)₂, or —C(O)-(heterocycloalkyl which may be substituted with lower alkyl, —OH or heterocycloalkyl).

As still another aspect, a compound according to (10), in which R⁵¹ is H, lower alkyl, -lower alkylene-C(O)—N(R⁰)₂, or —C(O)-(heterocycloalkyl which may be substituted with lower alkyl).

As still another aspect, a compound according to (10), in which R⁵¹ is lower alkyl.

As further still another aspect, a compound according to (10), in which R⁵¹ is H.

(10-2) a compound according to (10), in which each of R⁵² and R⁵³ is H.

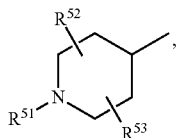
60 (11) A compound of the formula (I), in which arbitrary two or more among aspects described in the above-described (1) to (10-2) are combined.

Still another aspect of the present invention is shown below.

A compound of the formula (I) or a salt thereof in which X is N, Y is N, W is —O—, L¹ is -lower alkylene-, R¹ is methyl, R² is H, R³ is trifluoromethyl, all R⁴ are H's, R⁵ is

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[Chem. 5]



R⁵¹ is a lower alkyl, and each of R⁵² and R⁵³ is H.

Still another aspect of the present invention is shown below.

A compound of the formula (I) or a salt thereof in which X is CH;

R¹ is lower alkyl which may be substituted with one or more substituents selected from the group consisting of a halogen, —OH, —O-lower alkyl, and a nonaromatic heterocycle; in which the nonaromatic heterocycle may be substituted with lower alkyl; and

R⁵ is a nonaromatic heterocycle which may be substituted with a substituent selected from the D2 group.

Examples of a specific compound contained in the compound of the formula (I) or a salt thereof include the following compounds.

9-methyl-6-{6-[2-(piperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-9H-purine-2-carbonitrile,

6-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-9-methyl-9H-purine-2-carbonitrile,

9-methyl-6-[6-[2-[1-(1-methyl-L-prolyl)piperidin-4-yl]ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-9H-purine-2-carbonitrile,

6-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl}-9-(tetrahydrofuran-3-ylmethyl)-9H-purine-2-carbonitrile, and

6-{6-[2-(1-cyclobutylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-9-methyl-9H-purine-2-carbonitrile, and salts thereof.

Examples of another specific compound contained in the compound of the formula (I) or a salt thereof include the following compounds.

2-[4-(2-{4-[2-cyano-7-(2-hydroxyethyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidin-1-yl]-N,N-dimethylacetamide,

4-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile,

4-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl}-7-(2-hydroxyethyl)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile,

7-methyl-4-[6-(2-[[3R]-1-(1-methyl-D-prolyl)pyrrolidin-3-yl]oxy)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile, and

4-{6-[(1-[[1-(2-hydroxy-2-methylpropyl)piperidin-4-yl]carbonyl]piperidin-4-yl)methoxy]-5-(trifluoromethyl)pyridin-3-yl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile,

and salts thereof.

In the compound of the formula (I), tautomers or geometric isomers may be present depending on the types of substituents. In the present specification, the compound of the formula (I) is described as only one form of isomers, however the present invention also includes other isomers, and also includes a separated isomer or a mixture thereof.

In addition, the compound of the formula (I) may have an asymmetric carbon atom or axial asymmetry, and an optical isomer based on this may be present. The present invention

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also includes a separated optical isomer of the compound of the formula (I) or a mixture thereof.

Furthermore, the present invention also includes a pharmaceutically acceptable prodrug of the compound of the formula (I). The pharmaceutically acceptable prodrug is a compound having a group which can be converted into an amino group, a hydroxyl group, or a carboxyl group by solvolysis or under physiological conditions. As the group forming the prodrug, groups described in Prog. Med., 1985, Vol. 5, pp. 2157-2161, and "Pharmaceutical Research and Development, Drug Design" (Hirokawa Publishing Company), 1990, Vol. 7, pp. 163-198 are exemplified.

In addition, a salt of the compound of the formula (I) is a pharmaceutically acceptable salt of the compound of the formula (I), and an acid addition salt or a salt with a base may be formed depending on the types of substituents. Specifically, acid addition salts of inorganic acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, nitric acid, and phosphoric acid, and organic acids such as formic acid, acetic acid, propionic acid, oxalic acid, malonic acid, succinic acid, fumaric acid, maleic acid, lactic acid, malic acid, mandelic acid, tartaric acid, dibenzoyl tartaric acid, ditoluoyl tartaric acid, citric acid, methane sulfonic acid, ethane sulfonic acid, benzene sulfonic acid, p-toluene sulfonic acid, aspartic acid, and glutamic acid, salts of inorganic bases such as sodium, potassium, magnesium, calcium, and aluminum, and organic bases such as methylamine, ethylamine, ethanolamine, lysine, and ornithine, salts of various amino acids such as acetyl-leucine and the like and amino acid derivatives, and ammonium salt are exemplified.

Furthermore, the present invention also includes various hydrates or solvates of the compound of the formula (I) or a salt thereof, and crystal polymorphism substances. In addition, the present invention also includes a compound labeled with various radioactive or non-radioactive isotopes.

(Preparation Method)

The compound of the formula (I) and a salt thereof can be prepared by applying various known synthetic methods using the characteristics based on the basic structure thereof or the types of substituents. At that time, it may be effective in a preparing technology that the functional group is substituted with a suitable protective group (group which can be easily converted into the functional group) at the stage from a starting material to an intermediate depending on the types of functional groups. As such a protective group, the protective groups described in "Greene's Protective Groups in Organic Synthesis (4th edition, 2006)" written by P. G. M. Wuts and T. W. Greene can be exemplified, and these may be suitably selected and used depending on the reaction conditions. In such a method, the protective group is introduced, a reaction is performed, and the protective group is removed, if necessary. By doing this, it is possible to obtain a desired compound.

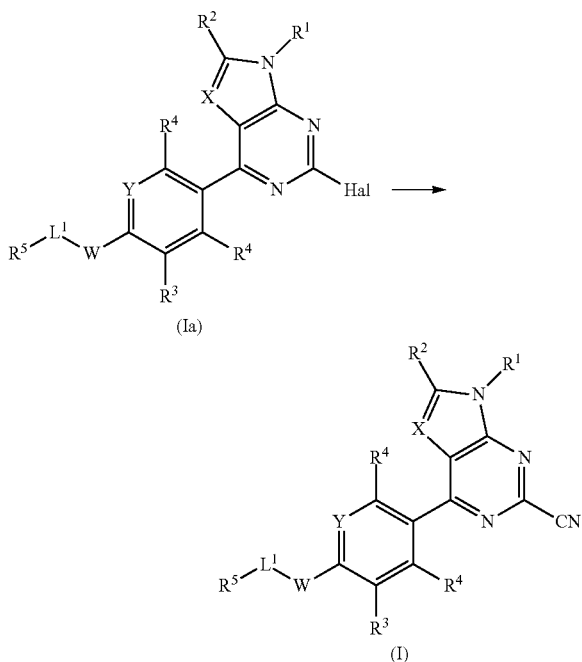
In addition, the prodrug of the compound of the formula (I) can be prepared by further performing the reaction by introducing a specific group at the stage from a starting material to an intermediate in the same manner as that of the above-described protective group, or using the obtained compound of the formula (I). The reaction can be performed by applying methods known to those skilled in the art such as general esterification, amidation, or dehydration.

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Hereinafter, representative preparation methods of the compound of the formula (I) will be described. Each preparation method can also be performed with reference to references described in the description. Moreover, the preparation method of the present invention is not limited to examples described below.

(First Preparation Method)

[Chem. 6]



(In the formula, Hal represents a halogen. The same shall apply hereinafter.)

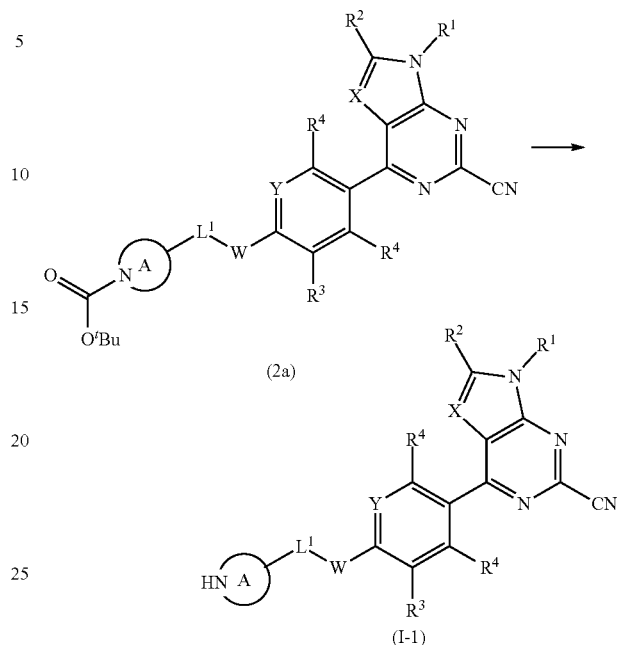
A compound (I) of the present invention can be obtained by subjecting a compound (1a) to a cyanation reaction.

In this reaction, the compound (1a) and a predetermined cyanation reagent are used in equivalent amounts, or either thereof in an excess amount, and the mixture thereof is stirred usually for 0.1 hours to 5 days under heating to reflux from room temperature, in a solvent which is inert to the reaction or in the absence of a solvent, in the presence of a predetermined catalyst. Examples of the solvent used in this reaction, which are not particularly limited, include aromatic hydrocarbons such as benzene, toluene and xylene, ethers such as diethylether, tetrahydrofuran, dioxane and dimethoxyethane, halogenated hydrocarbons such as dichloromethane, 1,2-dichloroethane and chloroform, N-methylpyrrolidone, N,N-dimethylformamide, N,N-dimethylacetamide, dimethyl sulfoxide, ethyl acetate, acetonitrile and a mixture thereof. Examples of the cyanation reagent include zinc cyanide. Examples of the predetermined catalyst include palladium (II) trifluoroacetate and tetrakis(triphenylphosphine)palladium (0). It is in some cases advantageous for smooth progress of the reaction to perform the reaction in the presence of phosphines such as triphenylphosphine or 2-(di-tert-butylphosphino)biphenyl.

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(Second Preparation Method)

[Chem. 7]



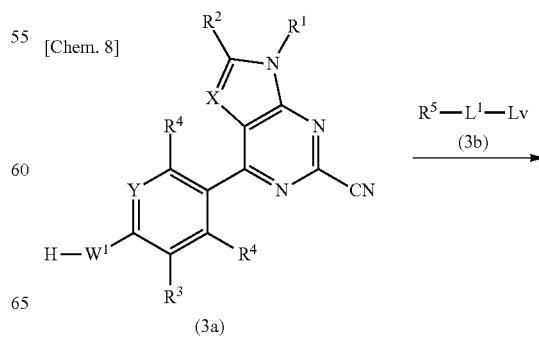
(In the formula, A represents nitrogen-containing heterocycloalkyl which may be substituted. The same shall apply hereinafter.)

A compound (I-1) of the present invention can be obtained by subjecting a compound (2a) to a Boc-elimination reaction.

In this reaction, the compound (2a) and an ethyl acetate solution of hydrogen chloride are used in equivalent amounts, or either thereof in an excess amount, and the mixture thereof is stirred usually for 0.1 hours to 5 days under heating to reflux from room temperature, in a solvent which is inert to the reaction or in the absence of a solvent. Examples of the solvent used in this reaction, which are not particularly limited, include aromatic hydrocarbons such as benzene, toluene and xylene, ethers such as diethylether, tetrahydrofuran, dioxane and dimethoxyethane, halogenated hydrocarbons such as dichloromethane, 1,2-dichloroethane and chloroform, N-methylpyrrolidone, N,N-dimethylformamide, N,N-dimethylacetamide, dimethyl sulfoxide, ethyl acetate, acetonitrile and a mixture thereof. In this reaction, a dioxane solution of hydrogen chloride, trifluoroacetic acid, or the like other than the ethyl acetate solution of hydrogen chloride can be used.

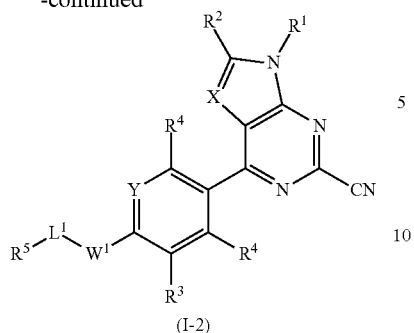
(Third Preparation Method)

[Chem. 8]



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-continued



(In the formula, Lv represents a leaving group, and W¹ represents —O— or —S—. The same shall apply hereinafter.)

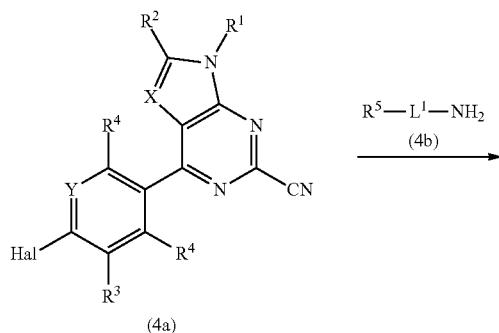
A compound (I-2) of the present invention can be obtained by subjecting a compound (3a) and a compound (3b) to an alkylation reaction. Here, examples of the leaving group include a halogen, a methanesulfonyloxy group and a p-toluenesulfonyloxy group.

In this reaction, the compound (3a) and the compound (3b) are used in equivalent amounts, or either thereof in an excess amount, and the mixture thereof is stirred usually for 0.1 hours to 5 days under heating to reflux from room temperature, in a solvent which is inert to the reaction or in the absence of a solvent, in the presence of a predetermined basic reagent. Examples of the solvent used in this reaction, which are not particularly limited, include aromatic hydrocarbons such as benzene, toluene and xylene, ethers such as diethylether, tetrahydrofuran, dioxane and dimethoxyethane, halogenated hydrocarbons such as dichloromethane, 1,2-dichloroethane and chloroform, N-methylpyrrolidone, N,N-dimethylformamide, N,N-dimethylacetamide, dimethyl sulfoxide, ethyl acetate, acetonitrile, and a mixture thereof. Examples of the basic reagent include organic bases such as triethylamine, N,N-diisopropyl ethylamine and N-methylmorpholine, or inorganic bases such as potassium carbonate, sodium carbonate, potassium hydroxide and sodium hydride. In addition, it is in some cases advantageous for smooth progress of the reaction to perform the reaction in the presence of potassium iodide or the like.

REFERENCES

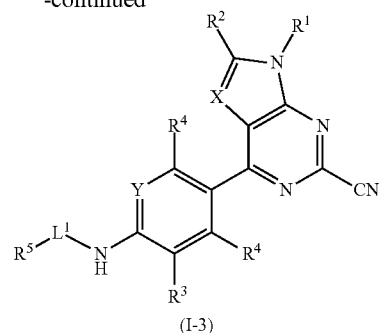
- “Organic Functional Group Preparations” written by S. R. Sandler and W. Karo, 2nd edition, Vol. 1, Academic Press Inc., 1991°
 “Courses in Experimental Chemistry” (5th edition) edited by The Chemical Society of Japan, Vol. 14 (2005) (Maruzen Co., Ltd.)
 (Fourth Preparation Method)

[Chem. 9]



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-continued



A compound (I-3) of the present invention can be obtained by subjecting a compound (4a) and a compound (4b) to an ipso substitution reaction.

In this reaction, the compound (4a) and the compound (4b) are used in equivalent amounts, or either thereof in an excess amount, and the mixture thereof is stirred usually for 0.1 hours to 5 days under heating to reflux from room temperature, in a solvent which is inert to the reaction or in the absence of a solvent, in the presence of a predetermined basic reagent. Examples of the solvent used in this reaction, which are not particularly limited, include aromatic hydrocarbons such as benzene, toluene and xylene, ethers such as diethylether, tetrahydrofuran, dioxane and dimethoxyethane, halogenated hydrocarbons such as dichloromethane, 1,2-dichloroethane and chloroform, N-methylpyrrolidone, N,N-dimethylformamide, N,N-dimethylacetamide, dimethyl sulfoxide, ethyl acetate, acetonitrile, and a mixture thereof. Examples of the basic reagent include organic bases such as triethylamine, N,N-diisopropyl ethylamine and N-methylmorpholine, or inorganic bases such as potassium carbonate, sodium carbonate, potassium hydroxide and sodium hydride. In addition, it is in some cases advantageous for smooth progress of the reaction to heat the reaction mixture by microwave irradiation.

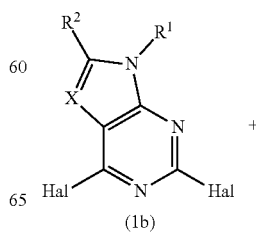
Various substituents on the groups R¹ and R⁵ in the compound of the formula (I) can be easily converted to other functional groups using the compound of the formula (I) as a starting material by using a reaction described in Examples described below, a reaction apparent to those skilled in the art, or modified methods thereof. For example, processes that those skilled in the art can usually employed such as reduction, halogenation, deprotection, hydrolysis, amidation, amination, oxidation, ureation, reductive amination, acylation, O-alkylation, N-alkylation, reductive alkylation, and epoxidation can be performed in arbitrary combination thereof.

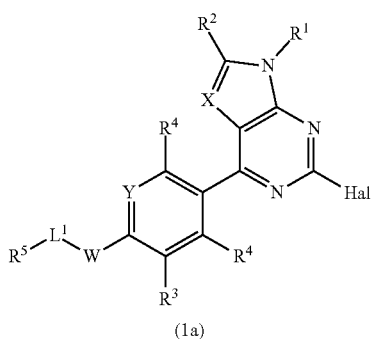
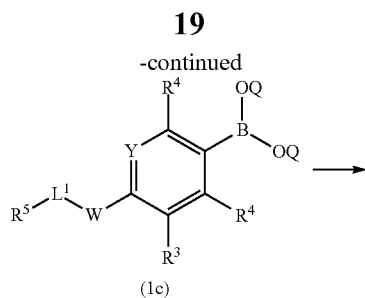
(Preparation of Starting Compound)

In the preparation methods above, a starting compound can be prepared by using, for example, the methods below, the methods described in Preparation Examples as described later, known methods, or modified methods thereof.

(Starting Material Synthesis 1)

[Chem. 10]





(In the formula, Hal's are the same as or different from each other, and Hal represents a halogen. Q's are the same as or different from each other, and Q represents H, lower alkyl, or lower alkylene in which two Q are combined to be one.)

The preparation method is a method of preparing a starting compound (1a) in the first preparation method.

A starting compound (1a) can be obtained by subjecting a compound (1b) and a compound (1c) to a coupling reaction.

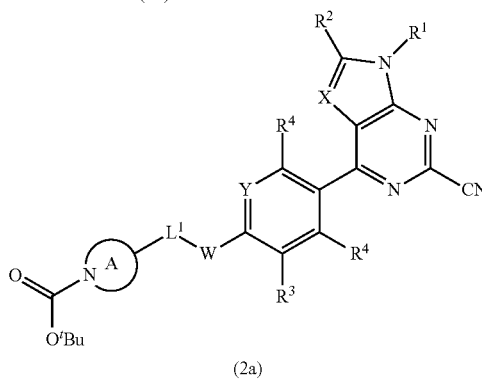
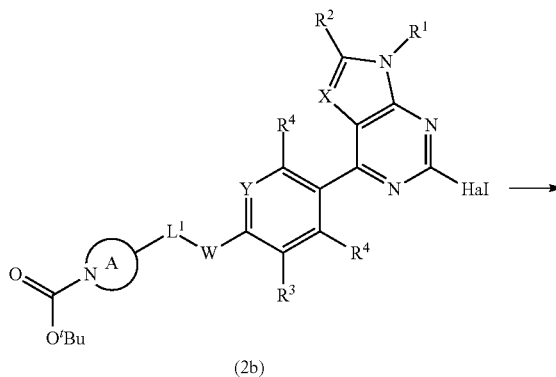
In this reaction, the compound (1b) and the compound (1c) are used in equivalent amounts, or either thereof in an excess amount, and the mixture thereof is stirred usually for 0.1 hours to 5 days under heating to reflux from room temperature, in a solvent which is inert to the reaction, in the presence of a basic reagent and a palladium catalyst. This reaction is preferably performed in an inert gas atmosphere. Examples of the solvent used in this reaction, which are not particularly limited, include aromatic hydrocarbons such as benzene, toluene and xylene, ethers such as diethylether, tetrahydrofuran, dioxane and dimethoxyethane, halogenated hydrocarbons such as dichloromethane, 1,2-dichloroethane and chloroform, alcohols such as methanol, ethanol, 2-propanol and butanol, N,N-dimethylformamide, dimethyl sulfoxide, water, and a mixed solvent thereof. Preferred examples of the base include inorganic bases such as potassium carbonate, sodium carbonate, cesium carbonate, potassium acetate and potassium hydroxide. Preferred examples of the palladium catalyst include tetrakis(triphenylphosphine)palladium, dichlorobis(triphenylphosphine)palladium, and [1,1'-bis(diphenylphosphino)ferrocene]palladium chloride. In addition, it is in some cases advantageous for smooth progress of the reaction to heat the reaction mixture by microwave irradiation.

20

REFERENCES

- “Metal-Catalyzed Cross-Coupling Reactions” edited by A. d. Meijere and F. Diederich, 1st edition, VCH Publishers Inc., 1997°
- “Courses in Experimental Chemistry” (5th edition) edited by The Chemical Society of Japan, Vol. 13 (2005) (Maruzen Co., Ltd.)
- (Starting Material Synthesis 2)

[Chem. 11]



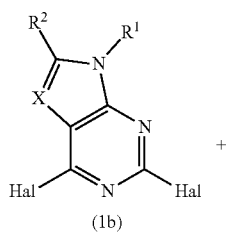
The preparation method is a method of preparing a starting compound (2a) in the second preparation method.

The starting compound (2a) can be obtained by subjecting a compound (2b) to a cyanation reaction.

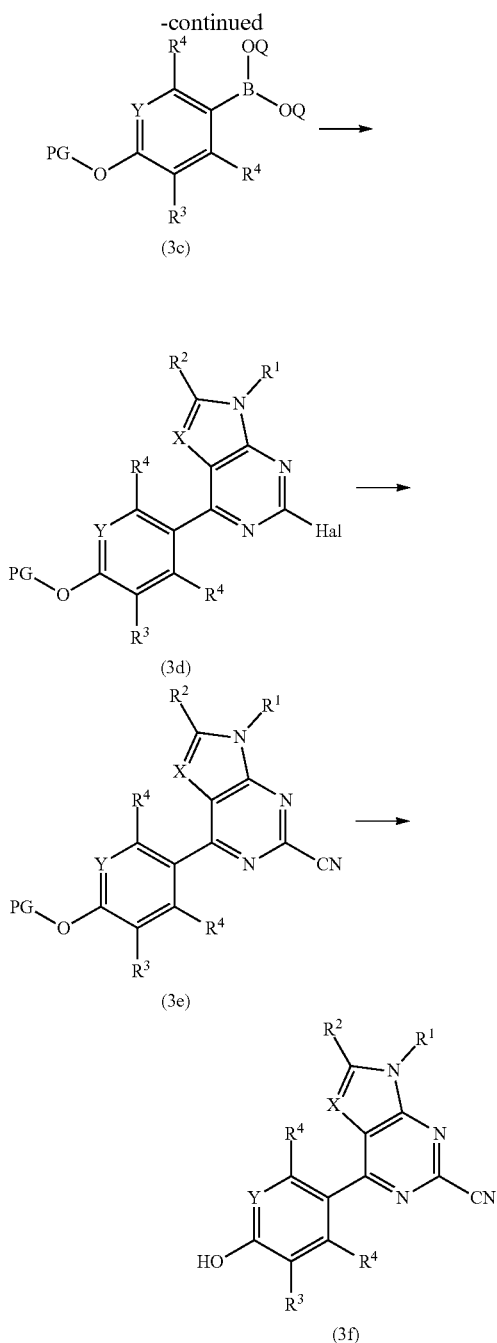
This reaction can be performed in the same manner as in the first preparation method. The compound (2b) is the compound (1a) which is nitrogen-containing heterocycloalkyl in which R⁵ is protected by Boc.

(Starting Material Synthesis 3)

[Chem. 12]



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(In the formula, PG represents a protective group.)

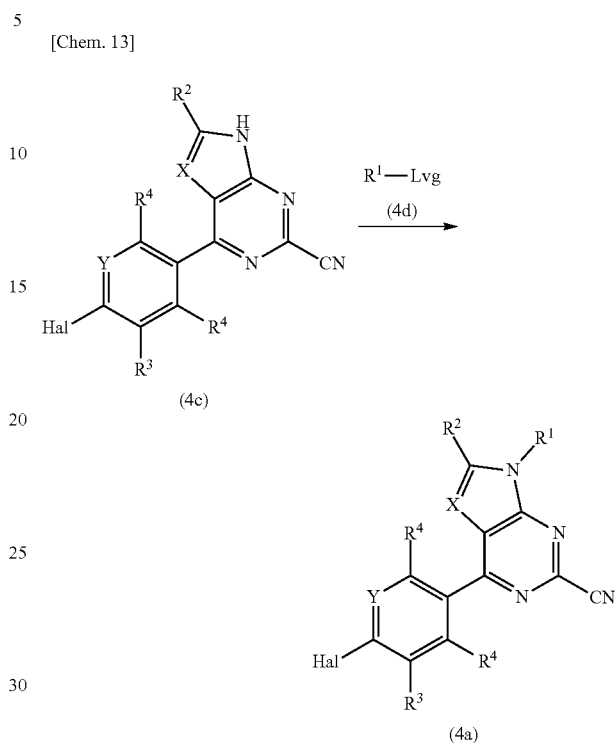
The preparation method is a method of preparing a compound (3f) in which W¹ in the starting compound (3a) in the third preparation method is —O—.

A compound (3d) can be obtained by subjecting the compound (1b) and a compound (3c) to a coupling reaction. Examples of the protective group include a p-methoxybenzyl group and the like. This reaction can be performed in the same manner as in the starting material synthesis 1.

A compound (3e) can be obtained by subjecting the compound (3d) to a cyanation reaction. This reaction can be performed in the same manner as in the first preparation method.

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The compound (3f) can be obtained by deprotecting the compound (3e).
(Starting Material Synthesis 4)



(In the formula, Lvg represents Lv or —OH.)

35 The preparation method is a method of preparing a starting compound (4a) in the fourth preparation method.

In a case where Lvg is Lv, the starting compound (4a) can be obtained by subjecting a compound (4c) and a compound (4d) to alkylation. This reaction can be performed in the same manner as in the third preparation method.

40 In a case where Lvg is —OH, the starting compound (4a) can be obtained by subjecting the compound (4c) and the compound (4d) to a Mitsunobu reaction. In this reaction, the compound (4c) and the compound (4d) are used in equivalent amounts, or either thereof in an excess amount, and the mixture thereof is stirred usually for 0.1 hours to 5 days under heating to reflux from room temperature, in a solvent which is inert to the reaction or in the absence of a solvent, in the presence of a predetermined organic phosphine, and ester or amide of azodicarboxylic acid. Examples of the solvent used in this reaction, which are not particularly limited, include aromatic hydrocarbons such as benzene, toluene and xylene, ethers such as diethylether, tetrahydrofuran, dioxane and dimethoxyethane, halogenated hydrocarbons such as dichloromethane, 1,2-dichloroethane, and chloroform, N-methylpyrrolidone, N,N-dimethylformamide, N,N-dimethylacetamide, dimethyl sulfoxide, ethyl acetate, acetonitrile, and a mixture thereof. Examples of the organic phosphine include triphenylphosphine and tributylphosphine. Examples of the ester or amide of azodicarboxylic acid include diethyl azodicarboxylate, bis(2-methoxyethyl) azodicarboxylate and 1,1'-(azodicarbonyl)dipiperidine.

50 The compound of the formula (I) is isolated and purified as a free compound, a salt thereof, a hydrate, a solvate, or a crystal polymorphism substance. The salt of the compound of the formula (I) can be prepared by a common salt formation reaction.

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Isolation and purification are carried out by applying usual chemical operations such as extraction, fractional crystallization, and various types of fractional chromatography.

Various isomers can be prepared by selecting a suitable starting compound, or can be separated using a difference in physicochemical properties among the isomers. For example, optical isomers are obtained by general optical resolution methods (for example, fractional crystallization leading to a diastereomeric salt with an optically active base or an acid, or chromatography using a chiral column) of a racemic mixture, and can also be prepared from a suitable optically active starting compound.

The pharmacological activity of the compound of the formula (I) was confirmed by the tests shown below.

Test Example 1

Measurement of a In Vitro Human Cathepsin S Inhibitory Activity

5 μ L of a human cathepsin S enzyme (R&D1183-CY-010) was added to a 96-well plate so as to be 20 ng/well. Next, 10-fold dilution of the test compound (a 10 mM DMSO solution) was serially performed with a buffer solution for assay (a solution of 50 mM sodium methoxide, 250 mM sodium chloride and 5 mM dithiothreitol (DTT), adjusted to pH=4.5 with 1 N hydrochloric acid) six times such that the final concentration becomes 0.1 nM to 10 μ M, then, 20 μ L of each of the solutions was added to the wells (the final DMSO concentration was 0.1%), and 25 μ L of synthetic substrate VVR-AMC (3211-V manufactured by PEPTIDE INSTITUTE, INC.) was added thereto such that the final concentration becomes 40 μ M, thereby starting an enzyme reaction. A fluorescence intensity (an excitation wavelength: 380 nm and a fluorescent wavelength: 460 nm) was measured at 37° C. for 10 minutes immediately after the start of the reaction using a fluorescence spectrophotometer (SPECTRA MAX GEMINI, manufactured by Molecular Devices Corporation), and the reaction rate at the time (5 minutes or 10 minutes) when linearity was observed was determined at each concentration of the test compound. A reaction rate at the time of not adding an enzyme in which the test compound was not added and a reaction rate at the time of adding an enzyme in which the test compound was not added were set to 100% inhibition and 0% inhibition, respectively, then, an inhibition rate at each concentration was determined, and IC₅₀ values were calculated by a linear regression method. The results are shown in Table 1. In the Table, Ex represents Example compound No., and Data 1 represents an IC₅₀ value (nM) of the human cathepsin S inhibitory activity.

TABLE 1

Ex	Data1
2	2.3
3	2.6
5	4.2
6	3.4
7	6.2
8	1.6
10	2.7
11	1.9
12	2.7
13	2.7
14	2.3
15	3.4
16	4.7

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TABLE 1-continued

Ex	Data1
19	1.7
20	3.8
21	1.9
23	1.9
24	0.76
25	1.6
26	1.2
27	1.9
28	1.4
29	3.5
30	4.7
31	2.7
32	2.3
33	6.1
34	1.7
35	2.1
36	8.9
37	3.6
38	4.5
39	1.0
40	3.1
41	4.8
42	2.9
44	3.6
45	2.6
47	2.1
50	2.7
51	3.4
52	2.6
54	3.9
56	3.3
58	2.6
65	3.1
66	2.9
68	2.2
70	2.2
71	2.0
72	1.6
73	2.0
74	2.2
75	2.5
76	3.0
77	2.7
78	2.6
79	1.4
80	3.5
81	3.2
82	4.5
83	3.1
84	1.3
85	3.3
86	1.9
87	2.7
88	2.4
89	3.1
90	3.0
91	2.1
92	2.0
93	2.1
94	3.6
95	2.3
96	4.0
97	3.8
101	2.8
102	2.0
103	3.7
104	7.8
105	2.2
106	2.8
107	3.1
108	3.2
109	2.4
110	2.3
111	4.3
112	8.9
113	1.9
114	2.9
115	3.3

25

TABLE 1-continued

Ex	Data1
116	2.5
117	2.3
118	2.3
119	2.7
120	2.1
121	2.4
122	1.7
135	2.1
136	1.5
137	1.0
138	2.7
139	2.1
140	2.1
141	2.0
142	1.5
143	1.5
144	2.8
145	1.0
146	3.2
147	2.4
148	2.8
149	2.7
150	1.7
152	2.2
153	1.6
154	2.4
155	1.5
156	2.4
157	2.1
158	2.4
159	2.7
160	2.4

Test Example 2

Evaluation of In Vitro MHC Class II Expression Inhibitory Effect Using Mouse Spleen Cells (Cell Evaluation System)

In antigen-presenting cells, the inhibition of cathepsin S inhibits the expression of MHC class II molecules. As a result, antibody production is inhibited, and decrease in an immune response occurs. For an increased expression of MHC class II in B cells, the inhibitory effect of the compound represented by formula (I) was examined.

First, spleen cells collected from C57BL/6 mice were seeded in a 96-well plate so as to be a 1×10^5 cells/well, then, 5-fold dilution of 10 mM DMSO solution of the test compound was serially performed with RPMI 1640 medium (including 10% fetal bovine serum (FCS), 5×10^{-5} M 2-mercaptoethanol, 50 IU/mL penicillin and 50 $\mu\text{g}/\text{mL}$ streptomycin) nine times such that the final concentration becomes 0.026 nM to 10 μM , and the resultant products were added thereto (the final DMSO concentration was 0.1%). At the same time, LPS (Sigma L4005) was added to the wells such that the final concentration becomes 2 $\mu\text{g}/\text{mL}$, and culture was performed at 37° C. for 48 hours in 5% CO₂. After culturing, the cells were stained at 4° C. for 20 minutes with PE-labeled streptavidin (BD BIOSCIENCE 554061) and biotin-labeled YAc antibody (EBIOSCIENCE 13-5741-85) together with FITC-labeled anti-mouse B220 antibody (BD BIOSCIENCE 553088), and an expression level (fluorescence intensity of YAc-biotin/streptavidin-PE) of MHC class II to which Ea peptide in B220 positive B cells was bonded was measured using a flow cytometry system (Guava EasyCyte Plus System, manufactured by Millipore Corporation). A value at the time of unstimulation of LPS in which the test compound was not added and a value at the

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time of stimulation of LPS in which the test compound was not added were set to 100% inhibition and 0% inhibition, respectively, then, an inhibition rate at each concentration was determined, and IC₅₀ values were calculated by a linear regression method. The results are shown in Table 2. In the Table, Ex represents Example compound No., and Data 2 represents an IC₅₀ value (nM).

TABLE 2

Ex	Data2
2	1.2
3	1.0
6	0.69
15	2.1
19	0.7
21	1.0
23	3.0
24	0.80
26	0.61
27	1.3
28	0.81
29	0.66
30	0.97
39	0.61
40	0.85
41	1.1
47	2.0
52	1.8
54	0.94
58	2.0
66	2.1
70	0.76
71	1.3
72	0.73
77	1.5
79	2.5
88	2.3
105	0.91
106	3.2
110	0.91
116	1.8
120	1.0
122	0.87
135	0.89
141	0.65
158	0.62

From the results of the above Test 2, it was confirmed that some compounds of the formula (I) inhibit the MHC class II increased expression in B cells.

Test Example 3

Evaluation of Ex Vivo MHC Class II Expression Inhibitory Effect Using Mouse Peripheral Blood

The expression inhibitory effect of MHC class II molecules was evaluated in an ex vivo system.

The test compound was orally administered into C57BL/6mice, and the inhibitory effect with respect to the increased expression of MHC class II in B cells in peripheral blood after oral administration was examined. That is, 10 mL/kg of the test compound dissolved in a vehicle (30% propylene glycol solvent (propylene glycol:cured castor oil (HCO40): Tween 80=4:2:1)/HCl (2 equivalents with respect to the test compound)/water) was orally administered (dose: 0.1 mg/kg, 0.3 mg/kg, 1 mg/kg, or 3 mg/kg) to a C57BL/6mice, and the peripheral blood was collected after 3 hours. 10 μL of RPMI 1640 medium (including 5×10^{-5} M of 2-mercaptoethanol, 50 IU/mL of penicillin, and 50 $\mu\text{g}/\text{mL}$ of streptomycin) or 10 μL of LPS adjusted with the medium such that the final concentration becomes 100 $\mu\text{g}/\text{mL}$ was added

to 90 μ L of the peripheral blood, and culture was performed at 37° C. for 15 hours in 5% CO₂. After culturing, the cells were stained at 4° C. for 30 minutes with FITC-labeled anti-mouse I-A/I-E antibody (BD BIOSCIENCE 553623) and PE-labeled B220 antibody (BD BIOSCIENCE 553090), and hemolysis and fixation were performed at 37° C. for 10 minutes using a buffer (BD BIOSCIENCE Phosflow Lyse/Fix Buffer 558049). After washing, using a flow cytometry system (FACSCanto II, manufactured by BD BIOSCIENCE), an expression level of MHC class II on a B220 positive B cell surface was measured by using a fluorescence intensity of FITC (hereinafter, referred to as MFI) as an indicator. The difference between a MFI at the time of stimulation of LPS and a MFI at the time of unstimulation of LPS was set to a Δ MFI, a Δ MFI of a mouse in which 10 mL/kg of only vehicle was administered was set to 1, and the inhibition rate of Δ MFI by the test compound administration was calculated.

From the results of the above Test 3, the compound of Example 28 exhibited 40 \pm 1.4% inhibition (0.3 mg/kg, p<0.001), the compound of Example 39 exhibited 61 \pm 1.2% inhibition (1 mg/kg, p<0.001), and the compound of Example 158 exhibited 59 \pm 1.5% inhibition (0.3 mg/kg, p<0.001), and thus, the increased expression of MHC class II was significantly inhibited. That is, it was confirmed that some compounds of the formula (I) inhibit the increased expression of MHC class II in peripheral blood B cells.

Test Example 4

Evaluation of Antibody Production Inhibitory Effect in Mouse OVA-Induced Antibody Production Model

Evaluation of an antibody production inhibitory effect in vivo was performed.

The BALB/c male mice were sensitized by intraperitoneal administration of 200 μ L of an emulsion consisting of 100 μ g of ovalbumin (OVA) (SIGMA A5503) and 100 μ L of a complete Freund's adjuvant (CFA) (DIFCO 231131) (Day 0), and a week after that, the mice were sensitized by intraperitoneal administration of 200 μ L of an emulsion consisting of 50 μ g of OVA and 100 μ L of an incomplete Freund's adjuvant (IFA) (DIFCO 263910) (Day 7). The test compound dissolved in a vehicle (30% propylene glycol solvent (propylene glycol:HCO40:Tween 80=4:2:1)/HCl (2 equivalents with respect to the test compound)/water) was orally administered (dose: 0.3 mg/kg, 1 mg/kg, or 3 mg/kg) at 10 mL/kg/day from Day 0 to Day 14, and an OVA specific IgG antibody titer in blood plasma at Day 14 was evaluated. Measurement of antibody titer was carried out using a blood plasma sample which was diluted 40,000 times by an ELISA method, and the antibody titer was represented as a ratio with respect to control blood plasma. An antibody titer of a mouse to which the vehicle was orally administered at 10 mL/kg/day from Day 0 to Day 14 was set to 0% inhibition, an antibody titer of a normal mouse to which the vehicle was orally administered at 10 mL/kg/day from Day 0 to Day 14 and which was not sensitized with OVA was set to 100% inhibition, and the inhibition rate at the time of the test compound administration was calculated.

From the results of the above Test 4, the compound of Example 2 exhibited 61 \pm 10% inhibition (1 mg/kg, p=0.02), the compound of Example 28 exhibited 83 \pm 2% inhibition (1 mg/kg, p<0.001), and the compound of Example 39 exhibited 74 \pm 8% inhibition (0.3 mg/kg, p<0.001), and thus, the antibody production was significantly inhibited. That is, it

was confirmed that some compounds of the formula (I) inhibit the antibody production.

Test Example 5

Evaluation of SLE Onset Inhibitory Effect in Spontaneous Model Using NZB/W F1 Mouse

NZB/W F1 mice are used as a model of SLE that spontaneously develops a condition close to humans (Helyer et al, Proceedings of the university otago medical school (New Zealand), 1961, Vol. 39, p. 3; Advances in Immunology, 1985, Vol. 37, p. 269-390; Perry et al, Journal of Biomedicine and Biotechnology, 2011, Article ID 271694, Review Article).

NZB/W F1 female mice were grouped on the basis of a urine protein level at the time of 20 weeks old, an anti-dsDNA antibody titer (IgG, IgM), an expression level of MHC class II in peripheral blood B cells, and a body weight. At this time, an individual having greater than 100 mg/dL of urine protein was excluded. After 20 weeks old, the test compound was orally administered once a day, urine collection and blood collection were performed every 4 weeks thereafter, and time-dependent transitions of urine protein, an anti-dsDNA antibody titer, and an expression level of MHC class II in peripheral blood B cells were evaluated. The evaluation ended at the time when urine protein-increased individual in the vehicle administration group reached the majority (at the time of 41 weeks old).

From the results of the above Test 5, at the time of 41 weeks old, urine protein of individuals of 6 cases among 8 cases in the vehicle administration group was 1,000 mg/dL or more. In the compound administration group (dose: 0.3 mg/kg) of Example 39, urine protein of all individuals of 10 cases was 100 mg/dL or less. Therefore, it was confirmed that the compound of Example 39 included in the formula (I) inhibits an increase of urine protein, and maintains kidney functions.

In addition, in the above Test 5, an anti-dsDNA IgG antibody titer at the time of 41 weeks old was measured by the ELISA method. The average of an anti-dsDNA IgG antibody titer of the vehicle administration group 6 cases was 69,375 \pm 20,607 (mU/mL). In the compound administration group (dose: 0.3 mg/kg) of Example 39, the average of an anti-dsDNA IgG antibody titer of 10 cases was 9,263 \pm 1,700 (mU/mL). Therefore, it was confirmed that the compound of Example 39 included in the formula (I) inhibits an increase of an anti-dsDNA antibody titer.

Furthermore, a pathology evaluation of kidney tissue in the above Test 5 was performed. A formalin-fixed paraffin-embedded section of kidney tissue was prepared, then, a specimen was prepared by periodic acid-methenamine-silver (PAM) staining, and the specimen was observed with a microscope. In accordance with histological classification of lupus nephritis by WHO, a histopathological image was classified into Class I (normal glomerulus), Class II (mesangium lesion), Class III (focal segmental glomerulonephritis), Class IV (diffuse glomerulonephritis), and Class VI (progressive sclerosing lupus nephritis).

From the results of the above Test 5, it was confirmed that the compound of Example 39 included in the formula (I) can be used for preventing and/or treating SLE.

From the above results, the compound of the formula (I) or salts thereof can be used for preventing and/or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, SLE, or lupus nephritis.

As an aspect of the graft rejection of an organ, bone marrow or tissue, antibody-related type rejection in which B cells are involved and T cell-related type rejection in which T cells are involved can be exemplified. As another aspect, antibody-related type rejection in which B cells are involved can be exemplified. As an aspect of the autoimmune disease, rheumatoid arthritis, psoriasis, ulcerative colitis, Crohn's disease, systemic lupus erythematosus, Sjogren's syndrome, multiple sclerosis, and the like can be exemplified. As an aspect of the allergic disease, asthma, atopic dermatitis, and the like can be exemplified. As an aspect of SLE, lupus nephritis, central nervous system lupus, and the like can be exemplified. As an aspect of the lupus nephritis, nephrotic syndrome, renal failure, and the like can be exemplified.

A pharmaceutical composition which contains one or two or more kinds of the compound of the formula (I) or a salt thereof as an active ingredient can be prepared by methods which are commonly used using excipients commonly used in the art, that is, pharmaceutical excipients or pharmaceutical carriers.

Administration may be any form of an oral administration by a tablet, a pill, a capsule, a granule, powder or a liquid medicine, or a parenteral administration by intraarticular, intravenous or intramuscular injections, a suppository, an eye drop, an eye ointment, a transdermal solution, an ointment, a transdermal patch, a transmucosal solution, a transmucosal patch or an inhalant.

As a solid composition for the oral administration, a tablet, powder, a granule or the like can be used. In such a solid composition, one or two or more kinds of active ingredients are mixed with at least one of inert excipients. According to commonly used methods in the related art, the composition may contain an inert additive, for example, a lubricant, a disintegrant, a stabilizer or a solubilizer. The tablet or pill, if necessary, may be coated with a film of sugar, or a stomach-soluble or enteric-soluble substance.

A liquid composition for oral administration contains an emulsion, a solution preparation, a suspension, a syrup or an elixir which is a pharmaceutically acceptable, and contains a generally used inert diluent, for example, purified water or ethanol. In addition to the inert diluent, the liquid composition may contain adjuvants such as a solubilizing agent, a wetting agent and a suspension, a sweetener, a flavor, a fragrance or a preservative.

The injection for parenteral administration contains a sterile aqueous or non-aqueous solution preparation, a suspension or an emulsion. As the aqueous solvent, for example, distilled water for injection or physiological saline are included. As the non-aqueous solvent, for example, alcohols such as ethanol are included. Such a composition may further include an isotonic agent, a preservative, a wetting agent, an emulsifier, a dispersant, a stabilizer or a solubilizer. For example, these are sterilized by a filtration through a bacteria-catching filter, mixing of a germicide, or irradiation. In addition, these also can be used by preparing a sterile solid composition, and before using, dissolving or suspending the composition in sterile water or a sterile solvent for injection.

As an external preparation, an ointment, a plaster, a cream, a jelly, a poultice, a spray, a lotion, an eye drop and an eye ointment are contained. A generally used ointment base, lotion base, aqueous or non-aqueous liquid medicine, suspension and emulsion are contained.

The transmucosal agent such as an inhalant and a pernasal agent can be used in a solid, liquid or semi-solid form, and can be prepared according to methods known in the related art. For example, a known excipient, a pH adjuster, a

preservative, a surfactant, a lubricant, a stabilizer and a thickener may be suitably added. In administration, it is possible to use a device for a suitable inhalation or insufflation. For example, using a known device such as a metered dose inhaler or a nebulizer, administration can be performed as a compound alone or a powder of a prescribed mixture, or as a solution or a suspension in combination with a carrier which is pharmaceutically acceptable. A dry powder inhaler may be an inhaler for single or multiple administrations, and it is possible to use dry powder or a powder-containing capsule. Alternatively, a suitable propellant, for example, a form of a pressurized aerosol spray using a suitable gas such as chlorofluoroalkane, hydrofluoroalkane or carbon dioxide may be used.

In a case of a normal oral administration, a suitable daily dose is about 0.001 mg/kg to 100 mg/kg of body weight, preferably 0.1 mg/kg to 30 mg/kg, and more preferably 0.1 mg/kg to 10 mg/kg, and this dose is administered once or 2 to 4 times. In a case of an intravenous administration, a daily dose is suitably about 0.0001 mg/kg to 10 mg/kg of body weight, and this dose is administered once or several times per day. In addition, as the transmucosal agent, about 0.001 mg/kg to 100 mg/kg of body weight is administered once or several times per day. The dose is suitably determined according to individual cases in consideration of symptoms, age and gender.

The dose differs depending on the types of an administration route, a dosage form, an administration site, an excipient and an additive, and the pharmaceutical composition of the present invention contains one or more kinds of the compound of the formula (I) or a salt thereof in which the active ingredient is 0.01% by weight to 100% by weight, and 0.01% by weight to 50% by weight as a certain aspect.

The compound of the formula (I) or a salt thereof can be used in combination with various therapeutic agents or preventive agents for which it is believed that the compound of the formula (I) or a salt thereof described above shows effectiveness. In the combined use, co-administration or separate administration in succession may be performed, or administration may be performed at a desired time interval. The co-administration formulation may be separately formulated, or may be a pharmaceutical composition including various therapeutic agents or preventive agents for which it is believed that the compound of the formula (I) or a salt thereof described above shows effectiveness and the compound of the formula (I) or a salt thereof.

EXAMPLES

Hereinafter, preparation methods for the compound of the formula (I) or salts thereof will be described in more detail with reference to Examples. Moreover, the present invention is not limited to compounds described in the following Examples. In addition, preparing methods for the starting compounds will be described in Preparation Examples. In addition, the preparation method for the compound of the formula (I) is not limited to only the preparation methods in specific Examples shown below, and the compound of the formula (I) can also be prepared by using a combination of the preparation methods or a method apparent to those skilled in the art.

The following abbreviations may be used in some cases in Examples, Preparation Examples, and Tables below.

PEX: Preparation Example No, Ex: Example No, PSyn: Preparation Example No prepared by the same method, Syn: Example No prepared in the same method, Str: chemical structural formula, DAT: physicochemical data.

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ESI+: m/z value in mass spectrometry (ionization method ESI, (M+H)⁺ unless otherwise specified), ESI-: m/z value (ionization method ESI, (M-H)⁻ unless otherwise specified), EI: m/z value in mass spectrometry (ionization method EI, (M)⁺ unless otherwise specified), CI: m/z value in mass spectrometry (ionization method CI, (M+H)⁺ unless otherwise specified), APCI+: m/z value in mass spectrometry (atmospheric pressure chemical ionization method APCI, (M+H)⁺ unless otherwise specified), APCI/ESI+: m/z value (simultaneous ionization of APCI and ESI, (M+H)⁺ unless otherwise specified), NMR1: δ (ppm) at ¹H NMR in dimethyl sulfoxide-d₆, NMR2: δ (ppm) at ¹H NMR in CDCl₃, s: singlet (spectrum), d: doublet (spectrum), t: triplet (spectrum), q: quartet (spectrum), m: multiplet (spectrum), br: broad line (spectrum) (example: br-s).

Me: methyl, Et: ethyl, iPr: isopropyl, tBu: tert-butyl, Boc: tert-butoxycarbonyl, Bn: benzyl.

Hex: n-hexane, EtOAc: ethyl acetate, DMF: N,N-dimethylformamide, DMAc: N,N-dimethylacetamide, MeOH: methanol, EtOH: ethanol, CH₂Cl₂: dichloromethane, CHCl₃: chloroform, THF: tetrahydrofuran.

Na₂CO₃: sodium carbonate, NaHCO₃: sodium bicarbonate, Na₂SO₄: anhydrous sodium sulfate, MgSO₄: anhydrous magnesium sulfate.

M: mol/L.

HCl in a structural formula represents hydrochloride and Fum represents fumarate. The number before HCl represents a molar ratio. For example, HCl means monohydrochloride and 2HCl means dihydrochloride.

A compound marked with "Chiral" in a structural formula represents a compound which is an optically active substance.

In the following Tables, there are cases where PSyn No. is described as Syn, and it shows that the Example compound is prepared by the same method as in the compound of PEx No. For example, in the following Tables, there is a description of PSyn3 in Syn of Ex. 123, and it shows that Example 123 is prepared by the same method as in Preparation Example 3.

A description of "3+4" in PSyn and Syn shows that first, preparation is performed by the same method as in Example 3, and subsequently, using the product as a starting material, preparation is performed by the same method as in Example 4. For example, in the following Tables, there is a description of PSyn24+11 in Syn of Ex. 102, and it shows that Example 102 is prepared by the same method as in Preparation Example 24, and subsequently, preparation is performed by the same method as in Example 11.

In the following Tables, a compound marked with "#" in Example No. represents a compound which is not included in definition of Claims.

Thermal analysis (TG/DTA) was measured under the following conditions. DSC Q 2000 manufactured by TA Instruments, room temperature to 300° C., a temperature increase rate of 10° C./minutes, N₂ (50 mL/minutes), a sample pan made of aluminum. For a description of a heat absorption peak temperature, "near" means 2° C., and as another aspect, means ±1° C.

Preparation Example 1

2-[4-ethoxy-3-(trifluoromethyl)phenyl]-4,4,5,5-tetraethyl-1,3,2-dioxaborolane (188 mg), Na₂CO₃ (217 mg), and tetrakis(triphenylphosphine)palladium (0) (29 mg) were added to a mixture of 2,4-dichloro-7-methyl-7H-pyrrolo[2,3-d]pyrimidine (100 mg), dioxane (2 mL), and water (0.3 mL) in an argon atmosphere, and the reaction mixture was

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stirred at 100° C. for 6 hours. After the reaction mixture was cooled to room temperature, extraction thereof was performed using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=9:1 to 7:3), whereby 2-chloro-4-[4-ethoxy-3-(trifluoromethyl)phenyl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine (130 mg) was obtained as a light yellow solid.

Preparation Example 2

After 60% sodium hydride (17 mg) was added to a solution of tert-butyl 4-(2-hydroxyethyl)piperidine-1-carboxylate (109 mg) in DMF (2 mL) at 0° C. in a nitrogen atmosphere, the mixture was stirred at room temperature for 30 minutes. After the reaction mixture was cooled to 0° C., 4-bromo-1-fluoro-2-(methylsulfonyl)benzene (100 mg) was added thereto. The reaction mixture was stirred at 0° C. for 30 minutes, and stirred at room temperature for 3 hours. After a saturated aqueous ammonium chloride solution was added to the reaction mixture, extraction thereof was performed using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=80:20 to 55:45), whereby tert-butyl 4-{2-[4-bromo-2-(methylsulfonyl)phenoxy]ethyl}piperidine-1-carboxylate (169 mg) was obtained as a colorless liquid.

Preparation Example 3

Zinc (29 mg), 2-(di-tert-butylphosphino)biphenyl (59 mg), zinc cyanide (63 mg), and palladium (II) trifluoroacetate (30 mg) were added to a solution of tert-butyl 4-{2-[4-(2-chloro-7-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-2-(trifluoromethyl)phenoxy]ethyl}piperidine-1-carboxylate (480 mg) in DMAc (10 mL) at room temperature in an argon atmosphere, and the mixture was stirred at 110° C. for 5 hours. After the reaction mixture was cooled to room temperature, EtOAc was added thereto, followed by stirring, and the insoluble material was separated by filtration using Celite. This filtrate was washed with a saturated aqueous NaHCO₃ solution, water, and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=9:1 to 1:1), whereby tert-butyl 4-{2-[4-(2-cyano-7-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-2-(trifluoromethyl)phenoxy]ethyl}piperidine-1-carboxylate (236 mg) was obtained as a light yellow solid.

Preparation Example 4

Potassium carbonate (25.8 g) and 1-(chloromethyl)-4-methoxybenzene (14.4 mL) were added to a mixture of 4-bromo-2-(trifluoromethyl)phenol (15 g) and N-methyl pyrrolidone (100 mL) at room temperature, and the mixture was stirred at 90° C. for 3 hours. After the reaction mixture was cooled to room temperature, water was added thereto, then extraction thereof was performed using EtOAc, and the extract was washed with water. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=9:1 to 6:1),

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whereby 4-bromo-1-[(4-methoxybenzyl)oxy]-2-(trifluoromethyl)benzene (22 g) was obtained as a colorless oil.

Preparation Example 5

Zinc (64 mg), 2-(di-tert-butylphosphino)biphenyl (163 mg), zinc cyanide (230 mg), and palladium (II) trifluoroacetate (81 mg) were added to a DMAc (16.5 mL) solution of 2-chloro-6-{4-[(4-methoxybenzyl)oxy]-3-(trifluoromethyl)phenyl}-9-methyl-9H-purine (1.1 g) at room temperature in an argon atmosphere, and the mixture was stirred at 100° C. for 3 hours. After the reaction mixture was cooled to room temperature, water was added thereto, then extraction thereof was performed using EtOAc, and the extract was washed with water. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=1:1 to 1:4), whereby a mixture of 6-{4-[(4-methoxybenzyl)oxy]-3-(trifluoromethyl)phenyl}-9-methyl-9H-purine-2-carbonitrile and 6-[4-hydroxy-3-(trifluoromethyl)phenyl]-9-methyl-9H-purine-2-carbonitrile (502 mg) was obtained.

A mixture of the above mixture (501 mg), trifluoroacetic acid (3.5 mL), and anisole (0.62 mL) was stirred at room temperature for 2 hours. After the reaction mixture was concentrated under reduced pressure, the residue was purified by silica gel column chromatography (Hex:EtOAc=1:2 to 1:6), whereby 6-[4-hydroxy-3-(trifluoromethyl)phenyl]-9-methyl-9H-purine-2-carbonitrile (243 mg) was obtained as a white solid.

Preparation Example 6

tert-Butyl 4-(2-[[4-(methylphenyl)sulfonyl]oxy]ethyl)piperidine-1-carboxylate (2.65 g) and potassium carbonate (2.0 g) were added to a mixture of 4-bromo-2-(trifluoromethyl)benzenethiol (1.8 g) and DMF (20 mL) at room temperature, and the mixture was stirred at the same temperature overnight. The reaction mixture was further stirred at 80° C. for 3 hours. After the reaction mixture was cooled to room temperature, water (50 mL) and EtOAc (30 mL) were added thereto, then extraction thereof was performed using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=100:0 to 80:20), whereby tert-butyl 4-(2-[[4-bromo-2-(trifluoromethyl)phenyl]sulfonyl]ethyl)piperidine-1-carboxylate (2.83 g) was obtained as a colorless oil.

Preparation Example 7

tert-Butyl 4-(2-hydroxyethyl)piperidine-1-carboxylate (4.8 g) and triphenylphosphine (5.7 g) were added to a solution of 4-bromo-2-(trifluoromethyl)phenol (5 g) in THF (50 mL) at room temperature. Under ice-cooling, bis(2-methoxyethyl) azodicarboxylate (5.1 g) was added to the mixture at an internal temperature of 10° C. or lower. The mixture was stirred at room temperature for 6 hours. After the reaction mixture was ice-cooled, water (50 mL) was added dropwise thereto, and the mixture was stirred for 1 hour. Extraction was performed on the mixture using EtOAc, and the extract was washed with water. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. After Hex was added to the residue, the mixture was stirred, and the insoluble

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material was separated by filtration. The filtrate was concentrated under reduced pressure, and the residue was purified by basic silica gel column chromatography (Hex:EtOAc=9:1 to 6:1), whereby tert-butyl 4-(2-[[4-bromo-2-(trifluoromethyl)phenoxy]ethyl]piperidine-1-carboxylate (7.46 g) was obtained as a colorless oil.

Preparation Example 8

Palladium (II) acetate (62 mg) was added to a mixture of 1-bromo-2-(methoxymethoxy)benzene (1.2 g), 2-dicyclohexylphosphino-2',6'-dimethoxybiphenyl (227 mg), and THF (24 mL) in a nitrogen atmosphere. After the mixture was ice-cooled, a 0.5 M solution of cyclobutyl zinc bromide in THF (22.1 mL) was added dropwise thereto, and the mixture was stirred at room temperature overnight. After the reaction mixture was concentrated under reduced pressure, EtOAc was added to the residue. A saturated aqueous NaHCO₃ solution was added to the mixture, followed by stirring, and the insoluble material was separated by filtration using Celite. Extraction was performed on the filtrate using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=10:0 to 9:1), whereby 1-cyclobutyl-2-(methoxymethoxy)benzene (713 mg) was obtained as a colorless oil.

Preparation Example 9

10% palladium on carbon (about 50% wet) (170 mg) was added to a mixture of tert-butyl 3-(2-ethoxy-2-oxoethylidene)-8-azabicyclo[3.2.1]octane-8-carboxylate (1.7 g) and EtOH (34 mL), and the mixture was stirred at room temperature for 4 hours in a hydrogen atmosphere at 3 atm. The reaction mixture was separated by filtration using Celite, and the filtrate was concentrated under reduced pressure, whereby tert-butyl 3-(2-ethoxy-2-oxoethyl)-8-azabicyclo[3.2.1]octane-8-carboxylate (1.7 g) was obtained as a colorless oil.

Preparation Example 10

60% sodium hydride (109 mg) was added to a mixture of triethyl phosphonoacetate (1.02 g) and THF (12 mL) at 0° C. in a nitrogen atmosphere, and the mixture was stirred at room temperature for 10 minutes. 3-Benzyl-3-azabicyclo[3.2.1]octan-8-one (978 mg) and DMF (5 mL) were added to the mixture, and the mixture was stirred at room temperature overnight. After water was added to the reaction mixture, extraction thereof was performed using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=97:3 to 80:20), whereby ethyl (3-benzyl-3-azabicyclo[3.2.1]oct-8-ylidene)acetate (362 mg) was obtained as a colorless liquid.

Preparation Example 11

After a solution of tert-butyl 3-(2-ethoxy-2-oxoethyl)-8-azabicyclo[3.2.1]octane-8-carboxylate (1.7 g) in THF (10 mL) was cooled to -78° C. in an argon atmosphere, a 1.02 M solution of diisobutylaluminum hydride in toluene (17.5 mL) was added dropwise thereto, then, the mixture was

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stirred at the same temperature for 1 hour, and further stirred at 0° C. for 30 minutes. An aqueous Rochelle salt solution and EtOAc were added the reaction mixture, followed by stirring for 1 hour. After the temperature of the reaction mixture was raised to room temperature, extraction thereof was performed using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=97:3 to 80:20), whereby tert-butyl 3-(2-hydroxyethyl)-8-azabicyclo[3.2.1]octane-8-carboxylate (0.83 g) was obtained as a colorless oil.

Preparation Example 12

After concentrated hydrochloric acid (3.1 mL) was added to a solution of 1-cyclobutyl-2-(methoxymethoxy)benzene (1.43 g) in MeOH (29 mL) at room temperature, the mixture was stirred at 60° C. for 2 hours. After the reaction mixture was cooled to room temperature, a standard buffer solution of pH 6.86 was added thereto, and EtOAc was added. Extraction was performed on the mixture using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure, whereby 2-cyclobutylphenol (1.1 g) was obtained as a colorless oil.

Preparation Example 13

Tetra-n-butylammonium tribromide (4.72 g) was added to a solution of 2-cyclobutylphenol (1.21 g) in CH₂Cl₂ (24.2 mL) at room temperature, and the mixture was stirred at the same temperature overnight. After the reaction mixture was poured into a saturated aqueous sodium sulfite solution, extraction was performed on the mixture using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=100:0 to 84:16), whereby 4-bromo-2-cyclobutylphenol (1.85 g) was obtained as a colorless oil.

Preparation Example 14

Di-tert-butyl dicarbonate (300 mg) and 20% palladium hydroxide on carbon (about 50% wet) were added to a mixture of ethyl (3-benzyl-3-azabicyclo[3.2.1]oct-8-ylidene)acetate (350 mg) and EtOH (10 mL) in a nitrogen atmosphere, and the mixture was stirred at room temperature for 16 hours in a hydrogen atmosphere at 4 atm. The reaction mixture was separated by filtration using Celite, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=97:3 to 85:15), whereby tert-butyl 8-(2-ethoxy-2-oxoethyl)-3-azabicyclo[3.2.1]octane-3-carboxylate (371 mg) was obtained as a colorless liquid.

Preparation Example 15

Triethylamine (0.27 mL) and p-toluenesulfonyl chloride (269 mg) were added to a mixture of tert-butyl 3-(2-hydroxyethyl)-8-azabicyclo[3.2.1]octane-8-carboxylate (300 mg) and CH₂Cl₂ (6 mL) at room temperature, and the mixture was stirred at the same temperature overnight. After CHCl₃ and water were added to the reaction mixture, extraction thereof was performed using CHCl₃, and the extract was

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washed with water. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=95:5 to 70:30), whereby tert-butyl 3-(2-[[[4-methylphenyl)sulfonyl]oxy]ethyl)-8-azabicyclo[3.2.1]octane-8-carboxylate (375 mg) was obtained as a light brown oil.

Preparation Example 16

tert-Butyl 3-(2-[[[4-methylphenyl)sulfonyl]oxy]ethyl)-8-azabicyclo[3.2.1]octane-8-carboxylate (150 mg) and potassium carbonate (100 mg) were added to a solution of 4-[4-hydroxy-3-(trifluoromethyl)phenyl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (100 mg) in DMAc (2 mL) at room temperature, and the mixture was stirred at 80° C. for 1 hour. After the reaction mixture was cooled to room temperature, extraction thereof was performed using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=80:20 to 50:50), whereby tert-butyl 3-{2-[4-(2-cyano-7-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-2-(trifluoromethyl)phenoxy]ethyl}-8-azabicyclo[3.2.1]octane-8-carboxylate (146 mg) was obtained as a white solid.

Preparation Example 17

A 2 M solution of methylamine and triethylamine (1.8 mL) in THF (4.6 mL) were added to a solution of 2,4,6-trichloro-5-nitropyrimidine (2 g) in THF (30 mL) under ice-cooling, and the mixture was stirred for 1 hour under ice-cooling. After water was added to the reaction mixture, extraction thereof was performed using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=99:1 to 90:10), whereby 2,6-dichloro-N-methyl-5-nitropyrimidine-4-amine (680 mg) was obtained as a pale yellow solid.

Preparation Example 18

Tin (II) chloride dihydrate (2.0 g) was added to a solution of 2,6-dichloro-N-methyl-5-nitropyrimidine-4-amine (680 mg) in EtOH (13 mL) at room temperature. The mixture was stirred at room temperature for 3 hours, and stirred at 90° C. for 2 hours. The reaction mixture was cooled to room temperature, and concentrated under reduced pressure. EtOAc and a saturated aqueous NaHCO₃ solution were added to the residue, and the insoluble material was separated by filtration using Celite. Extraction was performed on the filtrate using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=80:20 to 50:50), whereby 2,6-dichloro-N⁴-methylpyrimidine-4,5-diamine (482 mg) was obtained as a pale yellow solid.

Preparation Example 19

Potassium carbonate (442 mg) and methyl iodide (0.17 mL) were added to a solution of 2,6-dichloro-8-methyl-7H-purine (500 mg) in THF (5 mL) under ice-cooling, and the

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mixture was stirred at room temperature for 5 hours. After the reaction mixture was ice-cooled, water was added thereto, extraction thereof was performed using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=9:1 to 8:2), whereby 2,6-dichloro-8,9-dimethyl-9H-purine (310 mg) was obtained as a white solid.

Preparation Example 20

Tetrahydrofuran-3-ylmethanol (8 mL) and tributylphosphine (22 mL) were added to a suspension of 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (10 g) in CH₂C₂ (150 mL) at room temperature. 1,1'-(azodicarbonyl)dipiperidine (22 g) was added thereto under ice-cooling, and the mixture was stirred at room temperature overnight. After Hex (300 mL) was added to the reaction mixture, the mixture was stirred, and the insoluble material was separated by filtration. The filtrate was concentrated under reduced pressure, and the residue was purified by silica gel column chromatography (Hex:EtOAc=9:1 to 6:4), whereby 2,4-dichloro-7-(tetrahydrofuran-3-ylmethyl)-7H-pyrrolo[2,3-d]pyrimidine (5.9 g) was obtained as a dark brown oil.

Preparation Example 21

1-(tert-Butoxycarbonyl)-4-piperidinemethanol (400 mg) was added to a mixture of 60% sodium hydride (90 mg) and DMF (11 mL) under ice-cooling in a nitrogen atmosphere, and the mixture was stirred for 10 minutes. A solution of 4-bromo-1-(bromomethyl)-2-(trifluoromethyl)benzene (550 mg) in DMF (2 mL) was added to the mixture under ice-cooling. After the reaction mixture was stirred for 30 minutes under ice-cooling, the temperature of the reaction mixture was raised to room temperature, and stirring was performed for 30 minutes. After a saturated aqueous ammonium chloride solution was added to the reaction mixture, extraction thereof was performed using an EtOAc-Hex mixed solvent (1:1), and the extract was washed with water and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=99:1 to 90:10), whereby tert-butyl 4-({[4-bromo-2-(trifluoromethyl)benzyl]oxy}methyl)piperidine-1-carboxylate (598 mg) was obtained as a pale yellow oil.

Preparation Example 22

A mixture of 2,6-dichloro-N⁴-methylpyrimidine-4,5-diamine (400 mg) and trifluoroacetic acid (8 mL) was stirred for 8 hours while heating to reflux. Furthermore, the reaction mixture was stirred at 120° C. for 1.5 hours under microwave irradiation. After the reaction mixture was concentrated under reduced pressure, phenylphosphonic dichloride (1.8 mL) was added to the residue, and the mixture was stirred 180° C. for 2 hours. After the reaction mixture was cooled to room temperature, cold water (20 mL) was added thereto, then, the mixture was stirred, and the precipitate was collected by filtration. The obtained solid was purified by silica gel column chromatography (Hex:EtOAc=100:0 to 90:10), whereby 2,6-dichloro-9-methyl-8-(trifluoromethyl)-9H-purine (435 mg) was obtained as a white solid.

Preparation Example 23

After tert-butyl 4-{2-[4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-2-(trifluoromethyl)phenoxy]ethyl}piperidine-

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1-carboxylate (266 mg), Na₂CO₃ (169 mg), and tetrakis(triphenylphosphine)palladium (0) (31 mg) were added to a mixture of 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (100 mg), dioxane (1.6 mL), and water (0.6 mL), the mixture was stirred at 150° C. for 2 hours under microwave irradiation, and cooled to room temperature. After water and EtOAc were added to the reaction mixture, the mixture was stirred, filtered, and washed with EtOAc, whereby tert-butyl 4-{2-[4-(2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-2-(trifluoromethyl)phenoxy]ethyl}piperidine-1-carboxylate (121 mg) was obtained as a brown solid.

Preparation Example 24

A 4 M solution of hydrogen chloride in EtOAc (5 mL) was added to a suspension of tert-butyl 4-(2-{4-[2-chloro-7-(2-methoxyethyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidine-1-carboxylate (500 mg) in EtOAc (5 mL), and the mixture was stirred at the same temperature for 1 hour. The reaction mixture was concentrated under reduced pressure, whereby 2-chloro-7-(2-methoxyethyl)-4-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl}-7H-pyrrolo[2,3-d]pyrimidine monohydrochloride (445 mg) was obtained as a pale yellow solid.

Preparation Example 25

Tris(dimethylamino)sulfonium difluorotrimethylsilicate (239 mg) was added to a mixture of 4-[4-fluoro-3-(trifluoromethyl)phenyl]-7-[(4-methylphenyl)sulfonyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (200 mg) and DMAc (3 mL), and the mixture was stirred at 100° C. for 1 hour under microwave irradiation. After the reaction mixture was cooled to room temperature, the reaction mixture was purified by silica gel column chromatography (Hex:EtOAc=98:2 to 60:40), whereby 4-[4-fluoro-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (81 mg) was obtained as a pale yellow solid.

Preparation Example 26

Triethylamine (0.35 mL) was added to a suspension of 2-chloro-7-(2-methoxyethyl)-4-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl}-7H-pyrrolo[2,3-d]pyrimidine monohydrochloride (440 mg) in dichloroethane (8 mL) at room temperature, and the mixture was stirred at the same temperature for 15 minutes. After the reaction mixture was ice-cooled, acetic acid (0.24 mL), sodium triacetoxyborohydride (359 mg), and acetaldehyde (0.14 mL) were added thereto, and the mixture was stirred for 1 hour under ice-cooling. After a saturated aqueous NaHCO₃ solution was added to the reaction mixture, extraction thereof was performed using CHCl₃, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (CHCl₃), whereby 2-chloro-4-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7-(2-methoxyethyl)-7H-pyrrolo[2,3-d]pyrimidine (431 mg) was obtained as a colorless oil.

Preparation Example 27

Bis(pinacolato)diboron (17.2 g), potassium acetate (13.3 g), and dichlorobis(triphenylphosphine)palladium (II) (1.9 g) were added to a solution of tert-butyl 4-{2-[4-bromo-2-(trifluoromethyl)phenoxy]ethyl}piperidine-1-carboxylate

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(20.4 g) in dioxane (204 mL) at room temperature in an argon atmosphere, and the mixture was stirred at 100° C. for 4 hours. After the reaction mixture was cooled to room temperature, extraction thereof was performed using EtOAc, and the extract was washed with water. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=6:1 to 4:1), whereby tert-butyl 4-{2-[4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-2-(trifluoromethyl)phenoxy]ethyl}piperidine-1-carboxylate (22.0 g) was obtained as a colorless oil.

Preparation Example 28

tert-Butyl 4-(2-hydroxyethyl)piperidine-1-carboxylate (7.7 g) and 1,1'-(azodicarbonyl)dipiperidine (11.6 g) were added to a mixture of 5-bromo-3-(trifluoromethyl)pyridine-2-ol (7 g) and toluene (124 mL) at room temperature. Tributylphosphine (11.6 mL) was added thereto under ice-cooling, and the mixture was stirred at room temperature overnight. After diisopropylether was added to the reaction mixture, the mixture was stirred, and the insoluble material was separated by filtration. The filtrate was concentrated under reduced pressure, and the residue was purified by silica gel column chromatography (Hex:EtOAc=100:0 to 70:30), whereby tert-butyl 4-(2-{[5-bromo-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate (8.93 g) was obtained as a colorless oil.

Preparation Example 29

Sodium cyanoborohydride (2.2 g) was added to a mixture of ethyl 4-piperidine acetate (2.0 g), (1-ethoxycyclopropoxy)trimethylsilane (6.1 g), acetic acid (6.7 mL), a molecular sieve 3 A (2.0 g), MeOH (20 mL), and THF (20 mL) at room temperature, and the mixture was stirred at 65° C. for 6 hours. The reaction mixture was cooled to room temperature, and concentrated under reduced pressure. After EtOAc was added to the residue, the mixture was stirred, and the insoluble material was separated by filtration. After water was added to the filtrate, potassium carbonate was added to alkalinize the mixture. Extraction was performed on the mixture using EtOAc, and the extract was washed with a saturated aqueous NaHCO₃ solution and saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (CHCl₃:MeOH=100:0 to 95:5), whereby ethyl (1-cyclopropylpiperidin-4-yl)acetate (2.1 g) was obtained as a colorless oil.

Preparation Example 30

Lithium tetrahydroborate (730 mg) was added to a solution of ethyl (1-cyclopropylpiperidin-4-yl)acetate (2.1 g) in THF (53 mL) under ice-cooling. MeOH (2.4 mL) was added thereto, and the mixture was stirred at room temperature overnight. After water and EtOAc were added to the reaction mixture, the mixture was stirred for 30 minutes, and extraction was performed using EtOAc. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (Hex:EtOAc=80:20 to 60:40), whereby 2-(1-cyclopropylpiperidin-4-yl)ethanol (1.3 g) was obtained as a pale yellow oil.

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Preparation Example 31

Chloroacetyl chloride (0.43 mL) was added to a mixture of a solution of 4-cyanopiperidine (500 mg) in CH₂Cl₂ (5 mL) and an aqueous 2 M Na₂CO₃ solution (5.7 mL) under ice-cooling, and the mixture was stirred for 30 minutes under ice-cooling. After the reaction mixture was further stirred at room temperature for 4 hours, CH₂Cl₂ and water were added thereto, then, extraction was performed on the mixture using CH₂Cl₂, and the extract was washed with saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure, whereby 1-(chloroacetyl)piperidine-4-carbonitrile (845 mg) was obtained as a light yellow oil.

Preparation Example 32

Cyanomethylenetriethylphosphorane (1.1 mL) was added to a mixture of tert-butyl 4-{2-[4-(2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-2-(trifluoromethyl)phenoxy]ethyl}piperidine-1-carboxylate (1.0 g), tetrahydro-2H-pyran-4-ol (0.36 mL), and toluene (30 mL) at room temperature in a nitrogen atmosphere, and the reaction mixture was stirred overnight while heating to reflux. The reaction mixture was cooled to room temperature, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=67:33 to 33:67), whereby tert-butyl 4-(2-{4-[2-chloro-7-(tetrahydro-2H-pyran-4-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidine-1-carboxylate (1.04 g) was obtained as a yellow amorphous material.

Preparation Example 33

Bis(pinacolato)diboron (4.12 g), potassium acetate (4.35 g), and dichlorobis(triphenylphosphine)palladium (II) (1.04 g) were added to a solution of 4-bromo-1-[(4-methoxybenzyl)oxy]-2-(trifluoromethyl)benzene (5.87 g) in dioxane (90 mL) at room temperature in an argon atmosphere, and the mixture was stirred at 100° C. for 2 hours. After the reaction mixture was cooled to room temperature, 2,6-dichloro-9-methyl-9H-purine (3 g), Na₂CO₃ (4.7 g), tetrakis(triphenylphosphine)palladium (0) (854 mg), and water (15 mL) were added thereto. The reaction mixture was stirred at 100° C. for 4 hours. After the reaction mixture was cooled to room temperature, EtOAc was added thereto, followed by stirring, and the insoluble material was separated by filtration using Celite. Extraction was performed on the filtrate using EtOAc, and the extract was washed with water. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=4:1 to 1:1), whereby 2-chloro-6-{4-[(4-methoxybenzyl)oxy]-3-(trifluoromethyl)phenyl}-9-methyl-9H-purine (1.1 g) was obtained as a white solid.

Preparation Example 34

Acetic acid (0.1 mL) and a 1.0 M solution of tetrabutylammonium fluoride in THF (1 mL) were added to a solution of tert-butyl 4-(2-{4-[7-(2-{[tert-butyl(dimethyl)silyl]oxy}ethyl)-2-cyano-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidine-1-carboxylate (390 mg) in THF (5 mL) at room temperature, and the mixture was stirred at the same temperature for 7 hours. After a saturated aqueous NaHCO₃ solution was added to the reaction mixture, extraction thereof was performed using

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EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO_4 and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=70:30 to 40:60), whereby tert-butyl 4-(2-{4-[2-cyano-7-(2-hydroxyethyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidine-1-carboxylate (323 mg) was obtained as a colorless oil.

Preparation Example 48

2,6-Dichloro-9-methyl-9H-purine (5.6 g), Na_2CO_3 (11.7 g), water (16.8 mL), and tetrakis(triphenylphosphine)palladium (0) (1.6 g) were added to a solution of tert-butyl 4-(2-{[5-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate (13.8 g) in dioxane (112 mL) at room temperature in an argon atmosphere, and the mixture was stirred at an internal temperature of 85° C. for 5 hours. After the reaction mixture was cooled to room temperature, water (700 mL) was added thereto, and the mixture was stirred for 3 hours. The precipitate was collected by filtration, then, washed with water and diisopropyl ether, and dried. The obtained solid was purified by silica gel column chromatography (Hex:EtOAc:CHCl₃=1:1:1), whereby tert-butyl 4-(2-{[5-(2-chloro-9-methyl-9H-purin-6-yl)-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate (9.5 g) was obtained as a solid.

As a different method, the compound was obtained from a reaction of tert-butyl 4-(2-{[5-bromo-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate and 2,6-dichloro-9-methyl-9H-purine by the same method as in Preparation Example 33.

Preparation Example 110

Bis(pinacolato)diboron (11.5 g), potassium acetate (9.6 g), and dichlorobis(triphenylphosphine)palladium (II) (1.1 g) were added to a solution of tert-butyl 4-(2-{[5-bromo-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate (14.7 g) in dioxane (294 mL) at room temperature in an argon atmosphere, and the mixture was stirred at 110° C. for 2 hours. After the reaction mixture was cooled to room temperature, EtOAc (200 mL) was added thereto, followed by stirring. Then, the insoluble material was separated by filtration, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=98:2 to 90:10), whereby tert-butyl 4-(2-{[5-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate (14.0 g) was obtained as an oil.

Preparation Example 133

Zinc (498 mg), 2-di-tert-butylphosphino-2',4',6'-triisopropylbiphenyl (1.3 g), zinc cyanide (1.8 g), and palladium (II) trifluoroacetate (498 mg) were added to a solution of tert-butyl 4-(2-{[5-(2-chloro-9-methyl-9H-purin-6-yl)-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate (8.3 g) in DMAc (125 mL) at room temperature in an argon atmosphere, and the mixture was stirred at 50° C. for 3 hours. After the reaction mixture was cooled to room temperature, EtOAc (100 mL) and Celite were added thereto, and the mixture was stirred at room temperature. The insoluble material was separated by filtration, and a saturated aqueous NaHCO_3 solution, saturated brine, and water was added to the filtrate. Extraction was performed on

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the mixture using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO_4 and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=70:30 to 30:70), whereby tert-butyl 4-(2-{[5-(2-cyano-9-methyl-9H-purin-6-yl)-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate (7.6 g) was obtained as a solid.

Preparation Example 137

A solution of tert-butyl (3R)-3-hydroxypyrrolidine-1-carboxylate (7.0 g) in THF (30 mL) was added to a suspension of 60% sodium hydroxide (2.3 g) in THF (40 mL) under ice-cooling. At the same temperature, a THF (30 mL) solution of ethyl bromoacetate (13 mL) was added to the mixture, and the mixture was stirred overnight. After water and EtOAc were added to the reaction mixture, extraction thereof was performed using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over MgSO_4 and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=99:1 to 70:30), whereby tert-butyl (3R)-3-(2-ethoxy-2-oxoethoxy)pyrrolidine-1-carboxylate (6.6 g) was obtained as an oil.

Preparation Example 138

O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate (546 mg) was added to a solution of 1-(tert-butoxycarbonyl)piperidine-4-carboxylic acid (329 mg) in DMF (10 mL) at room temperature, and the mixture was stirred at the same temperature for 15 minutes. 7-Methyl-4-[6-(piperidin-4-yl methoxy)-5-(trifluoromethyl)pyridin-3-yl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (500 mg) and N,N-diisopropylethylamine (0.38 mL) were added to the reaction mixture at room temperature, and the mixture was stirred at the same temperature overnight. After EtOAc and a saturated aqueous NaHCO_3 solution were added to the reaction mixture, extraction thereof was performed using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO_4 and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (CHCl₃:MeOH=100:0 to 96:4), whereby tert-butyl 4-([4-{[5-(2-cyano-7-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-3-(trifluoromethyl)pyridin-2-yl]oxy}methyl)piperidin-1-yl]carbonyl)piperidine-1-carboxylate (488 mg) was obtained as a solid.

Preparation Example 139

Trifluoroacetic acid (2 mL) was added to a solution of tert-butyl (3R)-3-(2-{[5-(2-cyano-7-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-3-(trifluoromethyl)pyridin-2-yl]oxy}ethoxy)pyrrolidine-1-carboxylate (240 mg) in CH_2Cl_2 (4 mL)-acetonitrile (1 mL) at room temperature, and the mixture was stirred at the same temperature for 1 hour. After the reaction mixture was concentrated under reduced pressure, CH_2Cl_2 (4 mL) and a 4 M solution of hydrogen chloride in EtOAc (0.4 mL) were added to the residue at room temperature. The reaction mixture was concentrated under reduced pressure, and diisopropyl ether was added to the residue. The precipitate was collected by filtration, whereby 7-methyl-4-[6-{2-[(3R)-pyrrolidin-3-yloxy]

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ethoxy}-5-(trifluoromethyl)pyridin-3-yl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (208 mg) was obtained as a solid.

In the same manner as in Preparation Examples described above, compounds of Preparation Examples shown in the following tables were prepared. Structures, preparing methods, and physicochemical data of respective Preparation Example compounds are shown in Tables 3 to 24, respectively.

Example 1

Zinc (11 mg), 2-(di-tert-butylphosphino)biphenyl (23 mg), zinc cyanide (25 mg), and palladium (II) trifluoroacetate (12 mg) were added to a solution of 2-chloro-4-[4-ethoxy-3-(trifluoromethyl)phenyl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine (125 mg) in DMAc (2.5 mL) at room temperature in an argon atmosphere, and the mixture was stirred at 110° C. for 5 hours. After the reaction mixture was cooled to room temperature, EtOAc was added thereto, followed by stirring, and the insoluble material was separated by filtration using Celite. Extraction was performed on the filtrate using EtOAc, and the extract was washed with a saturated aqueous NaHCO₃ solution, water and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (Hex:EtOAc=9:1 to 1:1), whereby 4-[4-ethoxy-3-(trifluoromethyl)phenyl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (52.9 mg) was obtained as a pale yellow solid.

Example 2

A 4 M solution of hydrogen chloride in EtOAc (2 mL) was added to a solution of tert-butyl 4-{2-[4-(2-cyano-7-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-2-(trifluoroethyl)phenoxy]ethyl}piperidine-1-carboxylate (232 mg) in EtOAc (2 mL) at room temperature, and the mixture was stirred at the same temperature for 1 hour. The precipitate was collected by filtration, whereby 7-methyl-4-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (198 mg) was obtained as a white solid.

Example 3

Potassium carbonate (133 mg), potassium iodide (160 mg), and 2-chloro-N,N-dimethylacetamide (34 μL) were added to a suspension of 7-methyl-4-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (150 mg) in acetonitrile (6 mL) at room temperature, and the mixture was stirred at 60° C. for 2 hours. After the reaction mixture was cooled to room temperature, water was added thereto, extraction thereof was performed using EtOAc, and the extract was washed with a saturated aqueous ammonium chloride solution, water, and saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (Hex:EtOAc=1:3). The obtained product was washed with diisopropyl ether, and collected by filtration, whereby 2-(4-{2-[4-(2-cyano-7-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)-2-(trifluoromethyl)phenoxy]ethyl}piperidin-1-yl)-N,N-dimethylacetamide (118 mg) was obtained as a white solid.

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Example 4

Trifluoroacetic acid (0.41 mL) was added to a solution of tert-butyl {2-cyano-4-[4-ethoxy-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidin-7-yl}acetate (160 mg) in CH₂Cl₂ (1.6 mL) at room temperature, and the mixture was stirred at the same temperature for 5 hours. After the reaction mixture was concentrated under reduced pressure, the residue was purified by silica gel column chromatography (CHCl₃:MeOH=100:0 to 80:20), whereby {2-cyano-4-[4-ethoxy-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidin-7-yl}acetic acid (137 mg) was obtained as a white solid.

Example 5

1-Methylpiperazine (42 μL), N,N-diisopropylethylamine (66 μL), and (1-cyano-2-ethoxy-2-oxoethylidene aminoxy) dimethylamino-morpholino-carbenium hexafluorophosphate (187 mg) were added to a mixture of {2-cyano-4-[4-ethoxy-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidin-7-yl}acetic acid (100 mg) and DMF (4 mL) at room temperature, and the mixture was stirred at the same temperature for 2 hours. After water was added to the reaction mixture, the precipitate was collected by filtration, and washed with water. After the obtained solid was dissolved in CHCl₃, the solution was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (CHCl₃:MeOH=100:0 to 50:50), whereby 4-[4-ethoxy-3-(trifluoromethyl)phenyl]-7-[2-(4-methylpiperazin-1-yl)-2-oxoethyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (82 mg) was obtained as a white solid.

Example 6

A saturated aqueous NaHCO₃ and water were added to a suspension of 9-methyl-6-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-9H-purine-2-carbonitrile monohydrochloride (100 mg) in CHCl₃ (20 mL), and the mixture was stirred. Extraction was performed on the mixed solution using CHCl₃, and the extract was washed with water. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. After DMF (2 mL) was added to the residue, N-methyl-D-proline (41 mg) and O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate (122 mg) were added thereto, and the mixture was stirred at room temperature overnight. Extraction was performed on the reaction mixture using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (Hex:CHCl₃=50:50 to 0:100), whereby 9-methyl-6-[4-{2-[1-(1-methyl-D-prolyl)piperidin-4-yl]ethoxy]-3-(trifluoromethyl)phenyl]-9H-purine-2-carbonitrile (76 mg) was obtained as a white amorphous material.

Example 7

A saturated aqueous NaHCO₃ and water were added to a suspension of 9-methyl-6-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-9H-purine-2-carbonitrile monohydrochloride (100 mg) in CHCl₃ (20 mL), and the mixture was stirred. Extraction was performed on the mixed solution using CHCl₃, and the organic layer was washed with water.

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The organic layer was dried over MgSO_4 and filtered, and the filtrate was concentrated under reduced pressure. After CH_2Cl_2 (3 mL) was added to the residue to suspend the residue, dimethylcarbamoyl chloride (26 μL) and triethylamine (60 μL) were added thereto, and the mixture was stirred overnight at room temperature. Extraction was performed on the reaction mixture using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over Na_2SO_4 and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (CHCl_3 : MeOH=100:0 to 95:5), whereby 4-{2-[4-(2-cyano-9-methyl-9H-purin-6-yl)-2-(trifluoromethyl)phenoxy]ethyl}-N,N-dimethylpiperidine-1-carboxamide (83 mg) was obtained as a pale yellow solid.

Example 8

1-Methylpiperazine (28 μL), N,N-diisopropylethylamine (43 μL), and O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetraethyluronium hexafluorophosphate (109 mg) were added to a solution of (2-cyano-4-{4-[2-(tetrahydro-2H-pyran-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidin-7-yl}cetic acid (80 mg) in DMF (3.2 mL) at room temperature, and the mixture was stirred at the same temperature for 2 hours. After water was added to the reaction mixture, the precipitate was collected by filtration, and washed with water. After the obtained solid was dissolved in CHCl_3 , the solution was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (CHCl_3 :MeOH=100:0 to 50:50). After the obtained amorphous material was dissolved in CH_2Cl_2 (1 mL), a 4 M solution of hydrogen chloride in EtOAc (84 μL) was added thereto at room temperature. The mixture was concentrated under reduced pressure, and diisopropyl ether was added to the residue. The precipitate was washed with diisopropyl ether and collected by filtration, whereby 7-[2-(4-methylpiperazin-1-yl)-2-oxoethyl]-4-{4-[2-(tetrahydro-2H-pyran-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (89 mg) was obtained as a white solid.

Example 9

2-(1-Methylpiperidin-4-yl)ethanamine (93 mg) and N,N-diisopropylethylamine (0.22 mL) were added to a mixture of 4-[4-fluoro-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (100 mg) and N-methylpyrrolidone (2 mL), and the mixture was stirred at 140° C. for 1 hour under microwave irradiation. After the reaction mixture was cooled to room temperature, the reaction mixture was purified by silica gel column chromatography (CHCl_3 : MeOH=98:2 to 85:15), whereby 4-[4-{[2-(1-methylpiperidin-4-yl)ethyl]amino}-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (66 mg) was obtained as a white solid.

Example 10

Trifluoroacetic acid (1 mL) was added to a solution of tert-butyl 4-{2-[4-(2-cyano-8,9-dimethyl-9H-purin-6-yl)-2-(trifluoromethyl)phenoxy]ethyl}piperidine-1-carboxylate (100 mg) in CH_2Cl_2 (2 mL)-acetonitrile (0.5 mL) at room temperature, and the mixture was stirred at the same temperature for 1 hour. The reaction mixture was concentrated under reduced pressure, and CH_2Cl_2 (2 mL) was added to the residue. A 4 M solution of hydrogen chloride in EtOAc

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(0.2 mL) was added to the mixture at room temperature, and the precipitate was collected by filtration, whereby 8,9-dimethyl-6-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl}-9H-purine-2-carbonitrile monohydrochloride (59 mg) was obtained as a white solid.

Example 11

Triethylamine (114 μL) was added to a suspension of 8,9-dimethyl-6-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl}-9H-purine-2-carbonitrile monohydrochloride (128 mg) in dichloroethane (3 mL) at room temperature, and the mixture was stirred at the same temperature for 15 minutes. After the reaction mixture was ice-cooled, acetic acid (76 μL), sodium triacetoxyborohydride (113 mg), and acetaldehyde (45 μL) were added thereto, and the mixture was stirred for 1 hour under ice-cooling. After a saturated aqueous NaHCO_3 solution was added to the reaction mixture, extraction thereof was performed using CHCl_3 , and the extract was washed with water and saturated brine. The organic layer was dried over MgSO_4 and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (CHCl_3). The obtained product was washed with diisopropyl ether, and collected by filtration, whereby 6-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl}-8,9-dimethyl-9H-purine-2-carbonitrile (79 mg) was obtained as a pale yellow solid.

Example 12

Triethylamine (90 μL) was added to a solution of 7-methyl-4-{6-[2-(piperidin-3-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (100 mg) in dichloroethane (2 mL) at room temperature, and the mixture was stirred at the same temperature for 20 minutes. After mixture was ice-cooled, acetic acid (61 μL), sodium triacetoxyborohydride (91 mg), and acetaldehyde (36 μL) were added thereto, and the mixture was stirred for 1 hour under ice-cooling. After a saturated aqueous NaHCO_3 solution was added to the reaction mixture, extraction thereof was performed using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over MgSO_4 and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (Hex:EtOAc=80:20 to 40:60). The obtained product was dissolved in EtOAc (1 mL), then a 4 M solution of hydrogen chloride in EtOAc (0.3 mL) was added thereto at room temperature, and the mixture was stirred at the same temperature for 10 minutes. The precipitate was collected by filtration, whereby 4-{6-[2-(1-ethylpiperidin-3-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (92 mg) was obtained as a pale yellow solid.

Example 13

A 4 M solution of hydrogen chloride in EtOAc (0.12 mL) was added to a solution of 4-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7-(2-methoxyethyl)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (80 mg) in CH_2Cl_2 (1.5 mL) at room temperature, and the mixture was stirred at the same temperature for 0.5 hours. The reaction mixture was concentrated under reduced pressure, and diisopropyl ether was added to the residue. The precipitate was washed with diisopropyl ether, and collected by filtration,

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whereby 4-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7-(2-methoxyethyl)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (50 mg) was obtained as a pale yellow solid.

Example 14

4-[4-Hydroxy-3-(trifluoromethyl)phenyl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (100 mg) and potassium carbonate (87 mg) were added to a solution of 2-(1-cyclopropylpiperidin-4-yl)ethyl 4-methylbenzenesulfonate (152 mg) in DMAc (2 mL) at room temperature, and the mixture was stirred at 80° C. for 1 hour. After the reaction mixture was cooled to room temperature, EtOAc, water, and a saturated aqueous NaHCO₃ solution were added thereto, and the mixture was stirred. Extraction was performed on the mixture using EtOAc, and the extract was washed with water and saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (Hex:CHCl₃=50:50 to 0:100), whereby 4-{4-[2-(1-cyclopropylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (54 mg) was obtained as pale yellow powder.

Example 15

1-Methyl-L-proline (33 mg), O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate (85 mg), and N,N-diisopropylethylamine (64 μL) were added to a solution of 4-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7-(tetrahydrofuran-2-yl methyl)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (80 mg) in DMF (1.6 mL) at room temperature, and the mixture was stirred at the same temperature overnight. After the reaction mixture was diluted with EtOAc, a saturated aqueous NaHCO₃ solution and water were added thereto, then extraction thereof was performed using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (Hex:CHCl₃=50:50 to 0:100). The obtained product was dissolved in EtOAc (1 mL), then a 4 M solution of hydrogen chloride in EtOAc (31 μL) was added thereto at room temperature, and the mixture was stirred at the same temperature for 1 hour. The precipitate was collected by filtration, whereby 4-[4-{2-[1-(1-methyl-L-prolyl)piperidin-4-yl]ethoxy}-3-(trifluoromethyl)phenyl]-7-(tetrahydrofuran-2-ylmethyl)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (77 mg) was obtained as a pale yellow solid.

Example 16

Potassium carbonate (62 mg), potassium iodide (25 mg), and 2-chloro-N,N-dimethylacetamide (18 μL) were added to a mixture of 4-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7-(tetrahydrofuran-2-ylmethyl)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (80 mg) and acetonitrile (3.2 mL) at room temperature, and the mixture was stirred at 60° C. for 2 hours. After the reaction mixture was cooled to room temperature, the reaction mixture was diluted with EtOAc. Then, a saturated aqueous NaHCO₃ solution and water were added thereto, then extraction thereof was performed using EtOAc, and the extract was washed with saturated brine. The organic layer was

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dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (Hex:CHCl₃=50:50 to 0:100). The obtained product was dissolved in EtOAc (1 mL), then a 4 M solution of hydrogen chloride in EtOAc (37 μL) was added thereto at room temperature, and the mixture was stirred at the same temperature for 1 hour. The precipitate was collected by filtration, whereby 2-[4-(2-{4-[2-cyano-7-(tetrahydrofuran-2-yl methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidin-1-yl]-N,N-dimethylacetamide monohydrochloride (57 mg) was obtained as a pale yellow solid.

Example 17

2-(1-Methylpyrrolidin-3-yl)ethylamine (32 mg) and N,N-diisopropylethylamine (45 μL) were added to a solution of 4-[4-fluoro-3-(trifluoromethyl)phenyl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (75 mg) in DMAc (1.5 mL), then the mixture was stirred at 150° C. for 6 hours under microwave irradiation, and the mixture was further stirred at 160° C. for 3 hours. After the reaction mixture was cooled to room temperature, the reaction mixture was purified by basic silica gel column chromatography (Hex:EtOAc=50:50 to 30:70). The obtained product was dissolved in EtOH (1 mL), and fumaric acid (7 mg) was added thereto. The mixture was concentrated under reduced pressure, and diisopropyl ether was added to the residue. The precipitate was washed with diisopropyl ether, and collected by filtration, whereby 7-methyl-4-[4-{2-(1-methylpyrrolidin-3-yl)ethylamino}-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monofumarate (23 mg) was obtained as an amorphous material.

Example 18

A 4 M solution of hydrogen chloride in dioxane (1 mL) was added to a solution of tert-butyl 4-(2-{4-[7-(2-{[tert-butyl(dimethyl)silyl]oxy}ethyl)-2-cyano-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidine-1-carboxylate (150 mg) in dioxane (1 mL) at room temperature, and the mixture was stirred at the same temperature for 1 hour. After the reaction mixture was concentrated under reduced pressure, the precipitate was washed with diisopropyl ether, and collected by filtration, whereby 7-(2-hydroxyethyl)-4-{4-[2-(piperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (110 mg) was obtained as a pale yellow solid.

Example 28

Trifluoroacetic acid (40 mL) was added to a solution of tert-butyl 4-(2-{[5-(2-cyano-9-methyl-9H-purin-6-yl)-3-(trifluoromethyl)pyridin-2-yl]oxy}ethyl)piperidine-1-carboxylate (5.8 g) in CH₂Cl₂ (80 mL)-acetonitrile (20 mL) at room temperature, and the mixture was stirred at the same temperature for 1 hour. The reaction mixture was concentrated under reduced pressure, and CH₂Cl₂ (10 mL) was added to the residue. A 4 M solution of hydrogen chloride in EtOAc (10 mL) was added to the mixture at room temperature, and the resultant mixture was stirred at the same temperature for 10 minutes. EtOAc (5 mL) and diisopropyl ether (150 mL) were added to the reaction mixture at room temperature, and the resultant mixture was stirred for 30 minutes. The precipitate was collected by filtration, whereby 9-methyl-6-{6-[2-(piperidin-4-yl)ethoxy]-5-(trifluoromethyl)-

49

ethylpyridin-3-yl]-9H-purine-2-carbonitrile monohydrochloride (5.1 g) was obtained as a solid.

Example 39

Triethylamine (2.7 mL) was added to a suspension of 9-methyl-6-{6-[2-(piperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-9H-purine-2-carbonitrile monohydrochloride (3.0 g) in dichloroethane (60 mL) at room temperature, and the mixture was stirred at the same temperature for 20 minutes. After the reaction mixture was ice-cooled, acetic acid (1.8 mL), sodium triacetoxyborohydride (2.7 g) were added thereto, and a solution of acetaldehyde (1.1 mL) in dichloroethane (5 mL) was added thereto. The mixture was stirred for 1 hour under ice-cooling. After a saturated aqueous NaHCO₃ solution was added to the reaction mixture, extraction thereof was performed using CHCl₃, and the extract was washed with saturated brine. The organic layer was dried over Na₂SO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was suspended in EtOH (80 mL), and the suspension was dissolved by stirring for 30 minutes while heating to reflux. The mixture was cooled to room temperature, and the precipitate was collected by filtration, whereby 6-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-9-methyl-9H-purine-2-carbonitrile (1.8 g) was obtained as a crystal (heat absorption peak temperature obtained by a TG/DTA analysis: near 191° C.).

Example 103

By the same method as in Example 13 except that one equivalent fumaric acid was used instead of a solution of hydrogen chloride in EtOAc, 7-ethyl-4-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monofumarate was obtained.

Example 114

tert-Butyl 4-(2-{4-[2-cyano-7-(oxetan-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidine-1-carboxylate was reacted in the same method as in Preparation Example 24. Subsequently, by the same method as in Example 12 except that one equivalent fumaric acid was used instead of a solution of hydrogen chloride in EtOAc, 4-{4-[2-(1-ethylpiperidin-4-yl)ethoxy]-3-(trifluoromethyl)phenyl]-7-(oxetan-3-yl)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monofumarate was obtained.

Example 115

tert-Butyl 4-(2-{4-[2-cyano-7-(oxetan-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl]-2-(trifluoromethyl)phenoxy}ethyl)piperidine-1-carboxylate was reacted in the same method as in Preparation Example 24. Subsequently, by the same method as in Example 15 except that one equivalent fumaric acid was used instead of a solution of hydrogen chloride in EtOAc, 4-[4-{2-[1-(1-methyl-L-prolyl)piperidin-4-yl]ethoxy}-3-(trifluoromethyl)phenyl]-7-(oxetan-3-yl)-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monofumarate was obtained.

Example 134

A 4 M solution of hydrogen chloride in EtOAc (4.2 mL) was added dropwise to a suspension of 6-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-9-

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methyl-9H-purine-2-carbonitrile (3.9 g) in EtOH (58.5 mL) at room temperature, whereby a solution was obtained. After the solution was concentrated under reduced pressure, the residue was suspended in EtOH (117 mL). The suspension was stirred at 80° C. to obtain a solution, and the solution was cooled to room temperature while stirring. The solution was stirred at room temperature overnight. The precipitated solid was filtered, and the solid was washed with EtOH, whereby 6-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-9-methyl-9H-purine-2-carbonitrile monohydrochloride (3.2 g) was obtained as a crystal (heat absorption peak temperature obtained by a TG/DTA analysis: near 220° C.).

Example 135

Isobutylene oxide (38 μL) and N,N-diisopropylethylamine (73 μL) were added to a solution of 7-methyl-4-[6-{[1-(piperidin-4-yl carbonyl)piperidin-4-yl]methoxy}-5-(trifluoromethyl)pyridin-3-yl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monohydrochloride (120 mg) in EtOH (2.4 mL)-DMF (0.5 mL) at room temperature, and the mixture was stirred at 60° C. overnight. The reaction mixture was cooled to room temperature, and EtOAc and water were added thereto. Extraction was performed on the mixture using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by basic silica gel column chromatography (Hex:CHCl₃=70:30 to 20:80), whereby 4-{6-[(1-{1-(2-hydroxy-2-methylpropyl)piperidin-4-yl]carbonyl}piperidin-4-yl)methoxy]-5-(trifluoromethyl)pyridin-3-yl]-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile (93 mg) was obtained as a solid.

Example 136

Triethylamine (30 μL), 1H-benzotriazole-1-methanol (48 mg), and sodium triacetoxyborohydride (91 mg) were added to a solution of 9-methyl-6-{6-[2-(piperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-9H-purine-2-carbonitrile monohydrochloride (100 mg) in THF (1.5 mL)-EtOH (1.5 mL), and the mixture was stirred at room temperature for 16 hours. After a saturated aqueous NaHCO₃ solution and EtOAc were added to the reaction mixture, extraction thereof was performed using EtOAc, and the extract was washed with saturated brine. The organic layer was dried over MgSO₄ and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (CHCl₃:MeOH=100:0 to 80:20), whereby 9-methyl-6-{6-[2-(1-methylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-9H-purine-2-carbonitrile (57 mg) was obtained as a solid.

Example 143

By the same method as in Example 15 except that one equivalent fumaric acid was used instead of a solution of hydrogen chloride in EtOAc, 7-methyl-4-[6-(2-[(3S)-1-(1-methyl-D-prolyl)pyrrolidin-3-yl]oxy)ethoxy]-5-(trifluoromethyl)pyridin-3-yl]-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile monofumarate was obtained.

By the same method as in Examples described above, compounds of Examples shown in the following tables were prepared. Structures of respective Example compounds are shown in Tables 25 to 51, and preparing methods, and physicochemical data are shown in Tables 52 to 64.

TABLE 3

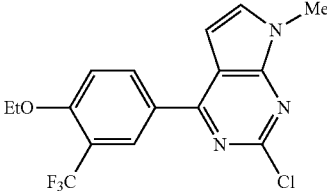
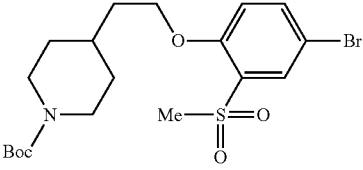
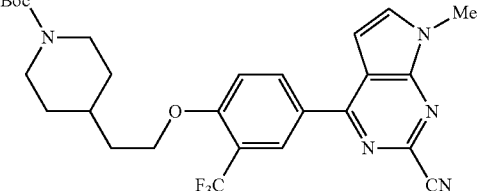
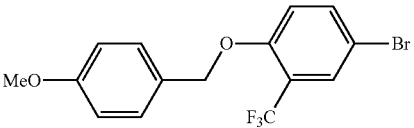
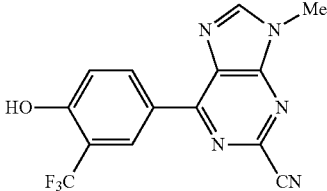
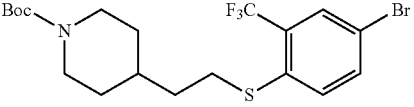
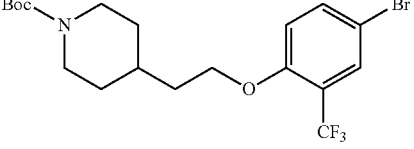
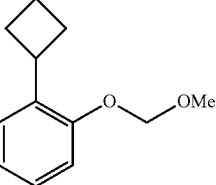
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4	4		Cl+: 361, 363
5	5		ESI+: 320
6	6		ESI+: 490, 492 [M + Na]
7	7		ESI+: 474, 476 [M + Na]
8	8		ESI+: 215 [M + Na]

TABLE 4

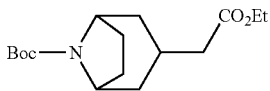
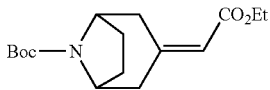
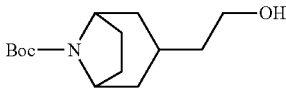
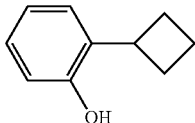
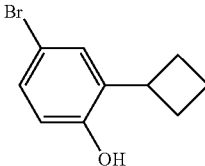
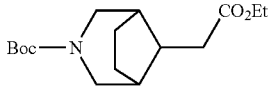
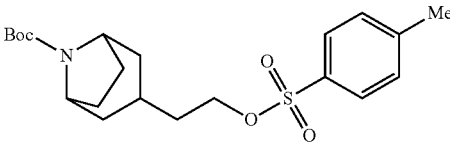
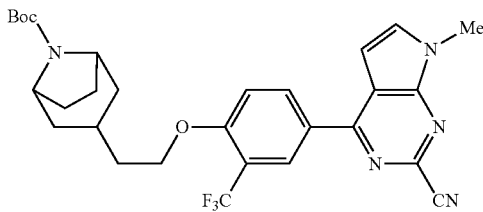
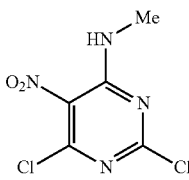
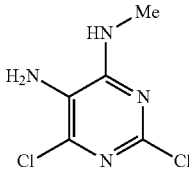
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11	11		APCI/ESI+: 256
12	12		ESI+: 149
13	13		EI: 226, 228
14	14		ESI+: 298
15	15		ESI+: 432 [M + Na]
16	16		ESI+: 556
17	17		ESI+: 223, 225, 227
18	18		ESI+: 193, 195, 197

TABLE 5

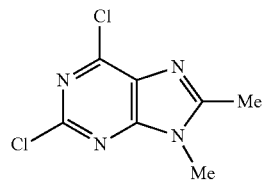
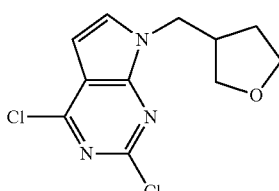
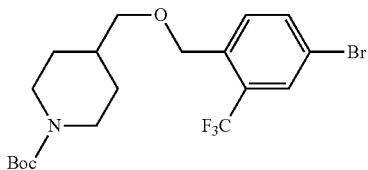
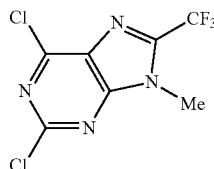
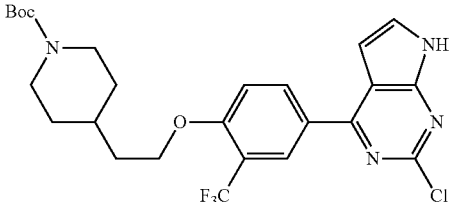
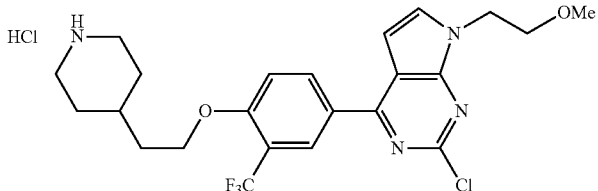
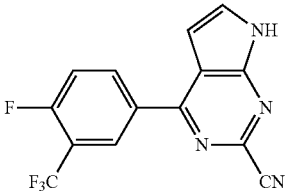
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21	21		ESI+: 452, 454
22	22		ESI+: 271, 273
23	23		ESI+: 525, 527
24	24		ESI+: 483, 485
25	25		ESI+: 307

TABLE 6

PEX	PSyn	Str	DAT
26	26		ESI+: 511, 513
27	27		ESI+: 522 [M + Na]
28	28		ESI+: 475, 477 [M + Na]
29	29		ESI+: 212
30	30		ESI+: 170
31	31		ESI+: 187, 189
32	32		ESI+: 609, 611
33	33		ESI+: 449, 451

TABLE 7

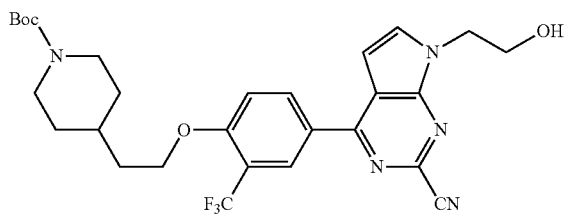
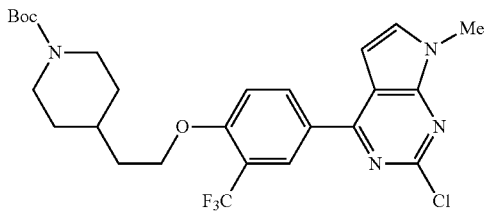
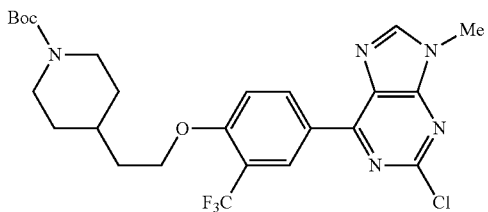
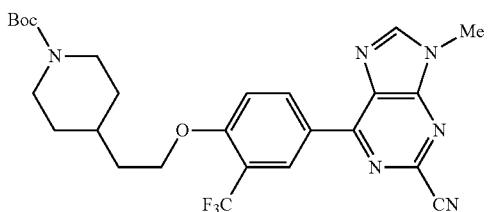
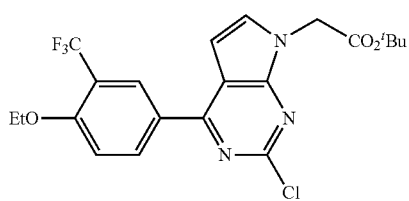
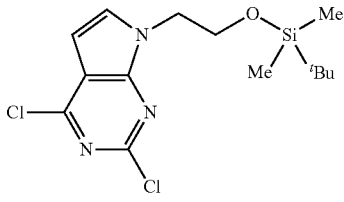
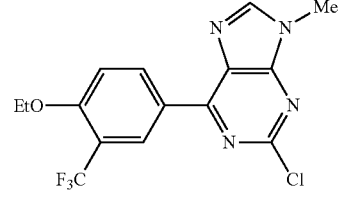
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35	1		ESI+: 539, 541
36	1		ESI+: 540, 542
37	3		ESI+: 531
38	1		ESI+: 456, 458
39	19		ESI+: 346, 348, 350
40	1		ESI+: 357, 359

TABLE 8

PEX	PSyn	Str	DAT
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42	3		ESI+: 674
43	33		ESI+: 540, 542
44	27		ESI+: 510
45	3		ESI+: 531
46	33		ESI+: 555, 557
47	4		EI: 216, 218

TABLE 9

PEx	PSyn	Str	DAT
48	48		ESI+: 541, 543
49	3		ESI+: 546
50	4		ESI+: 353, 355
51	19		ESI+: 302, 304, 306
52	15		APCI/ESI+: 344 [M - tBu + H]
53	16		ESI+: 546
54	27		ESI+: 423 [M + Na]
55	11		ESI+: 256

TABLE 10

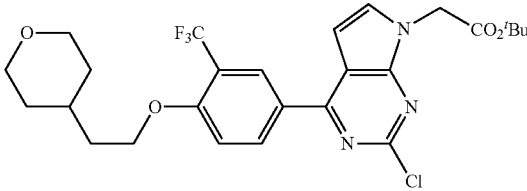
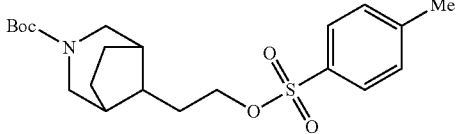
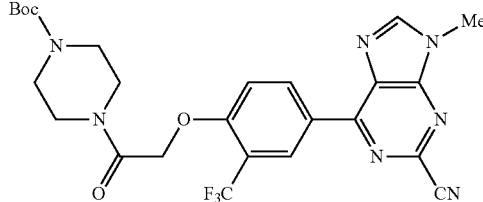
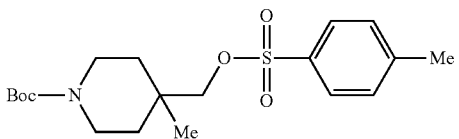
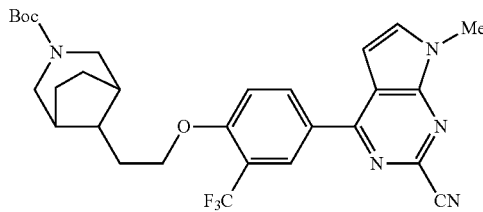
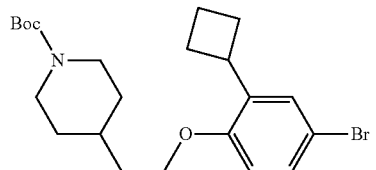
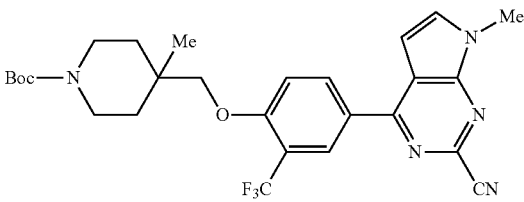
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58	16		ESI+: 546
59	15		ESI+: 406 [M + Na]
60	16		ESI+: 556
61	7		ESI+: 460, 462 [M + Na]
62	16		ESI+: 530

TABLE 11

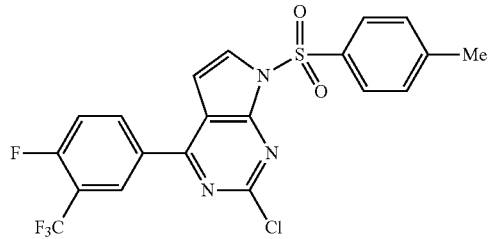
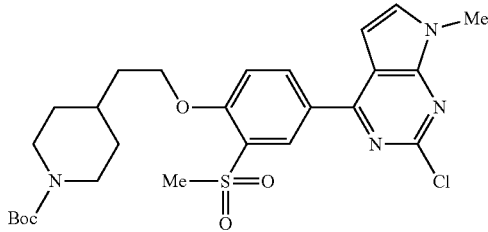
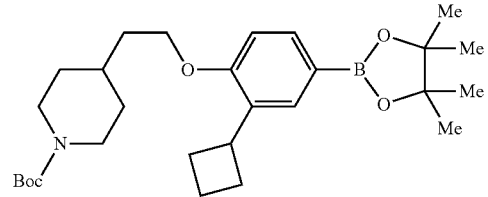
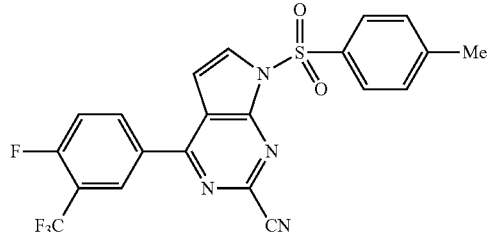
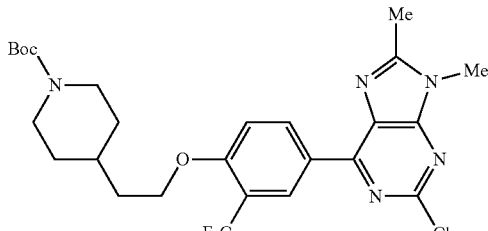
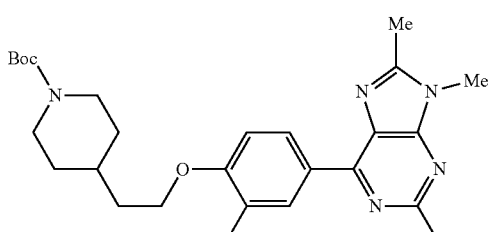
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65	27		ESI+: 508 [M + Na]
66	3		ESI+: 461
67	1		ESI+: 554, 556
68	3		ESI+: 545

TABLE 12

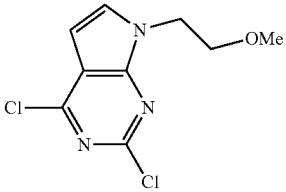
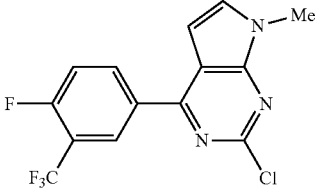
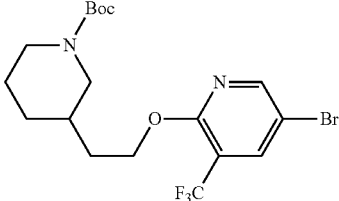
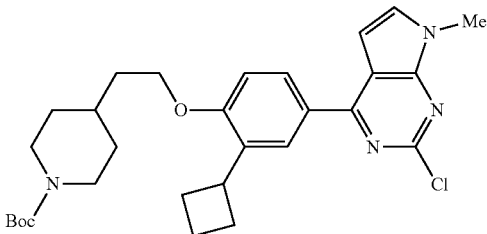
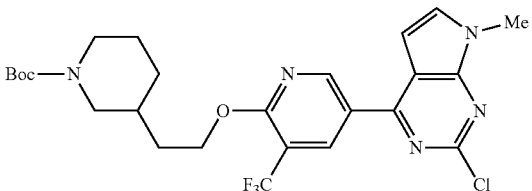
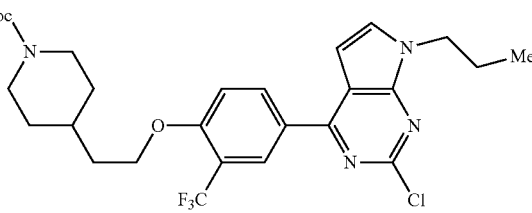
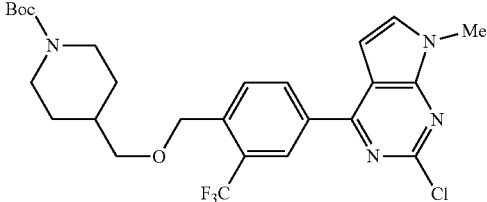
PEX	PSyn	Str	DAT
69	19		ESI+: 246, 248, 250
70	1		ESI+: 330, 332
71	28		ESI+: 453, 455
72	1		ESI+: 525, 527
73	33		ESI+: 540, 542
74	1		ESI+: 583, 585
75	33		ESI+: 539, 541

TABLE 13

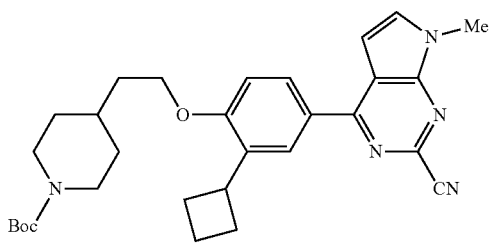
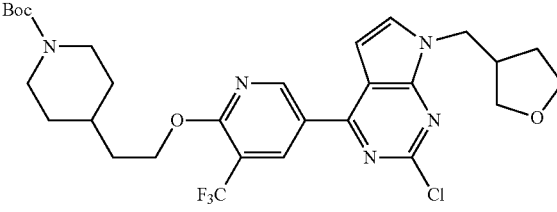
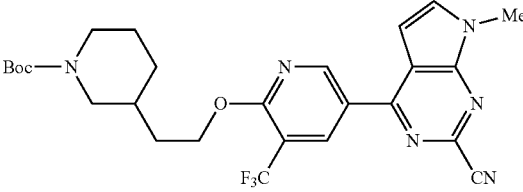
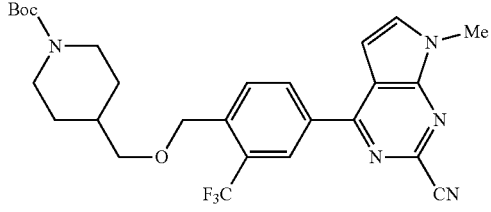
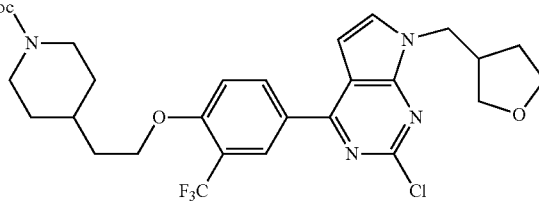
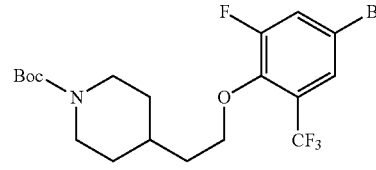
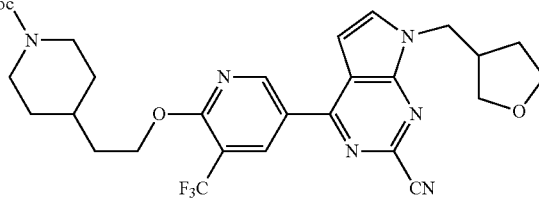
PEX	PSyn	Str	DAT
76	3		ESI+: 516
77	33		ESI+: 610, 612
78	3		ESI+: 531
79	3		ESI+: 530
80	1		ESI+: 609, 611
81	7		ESI+: 492, 494 [M + Na]
82	3		ESI+: 601

TABLE 14

PEX	PSyn	Str	DAT
83	33		ESI+: 631, 633 [M + Na]
84	3		ESI+: 600
85	3		ESI+: 622 [M + Na]
86	3		ESI+: 321
87	19		ESI+: 609, 611
88	3		ESI+: 600

TABLE 15

PEX	PSyn	Str	DAT
89	20		ESI+: 286, 288, 290
90	20		NMR1: 1.57-1.61 (1H, m), 1.88-1.99 (1H, m), 2.88 (1H, m), 3.47-3.53 (1H, m), 3.59-3.68 (2H, m), 3.75-3.82 (1H, m), 4.20-4.31 (2H, m), 8.79 (1H, s) ESI+: 273, 275, 277
91	1		ESI+: 679, 681
92	3		ESI+30: 562 [M + Na]
93	20		ESI+: 391
94	33		ESI+: 448, 450
95	1		ESI+: 610, 612

TABLE 16

PEX	PSyn	Str	DAT
96	33		ESI+: 611, 613
97	1		ESI+: 595, 597
98	10		ESI+: 226
99	3		ESI+: 670
100	3		ES+: 586
101	3		ES+: 601
102	25		ES+: 516

TABLE 17

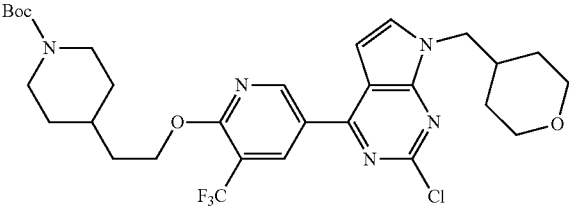
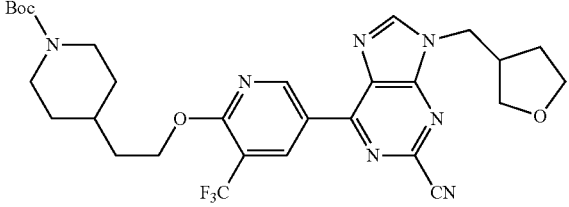
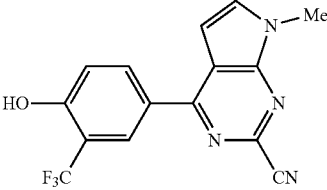
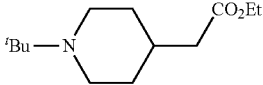
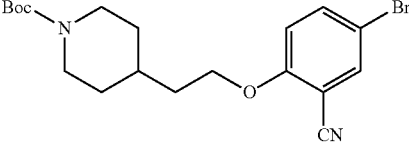
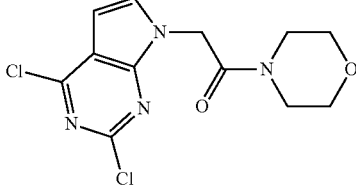
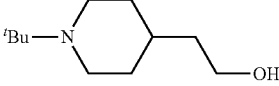
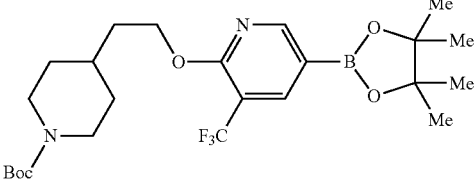
PEX	PSyn	Str	DAT
103	33		ESI+: 624, 626
104	3		ESI+: 602
105	5		ESI+: 319
106	9		ESI+: 228
107	2		ESI+: 409, 411
108	19		ESI+: 315, 317, 319
109	30		ESI+: 186
110	110		ESI+: 501

TABLE 18

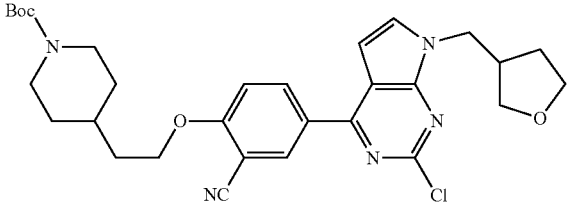
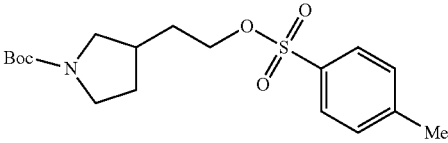
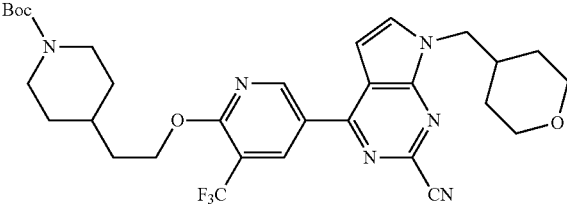
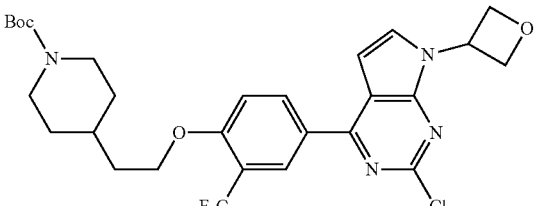
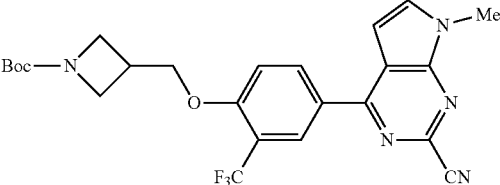
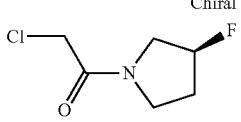
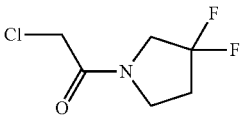
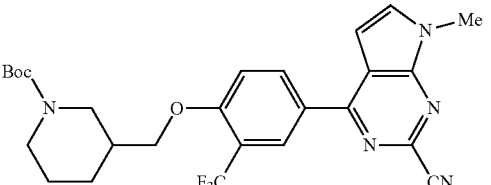
PEX	PSyn	Str	DAT
111	33		ESI+: 566
112	15		ESI+: 370
113	3		ESI+: 637 [M + Na]
114	32		ESI+: 581, 583
115	16		ESI+: 488
116	31		ESI+: 166, 168
117	31		ESI+: 184, 186
18	16		ESI+: 538 [M + Na]

TABLE 19

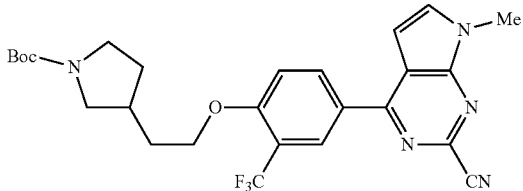
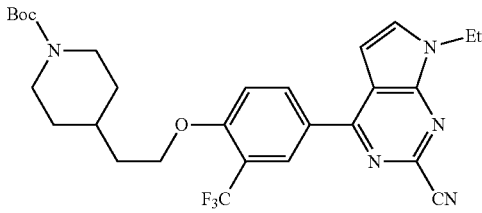
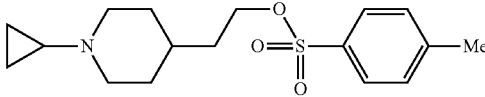
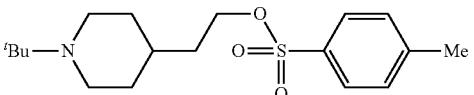
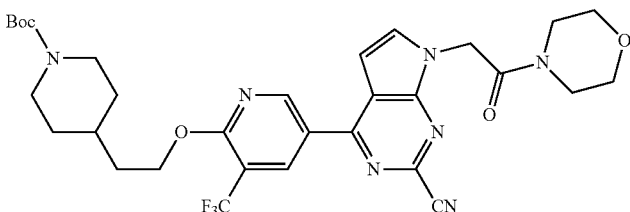
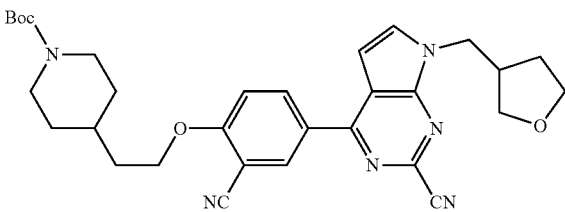
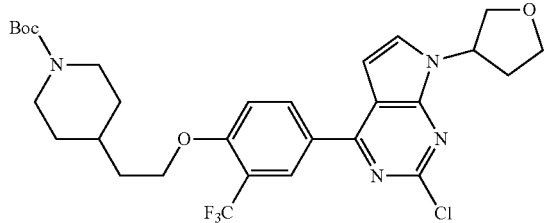
PEX	PSyn	Str	DAT
119	16		ESI+: 516
120	19		ESI+: 544
121	15		ESI+: 324
122	15		ESI+: 340
123	33		ESI+: 653, 655
124	3		ESI+: 557
125	32		ESI+: 595, 597

TABLE 20

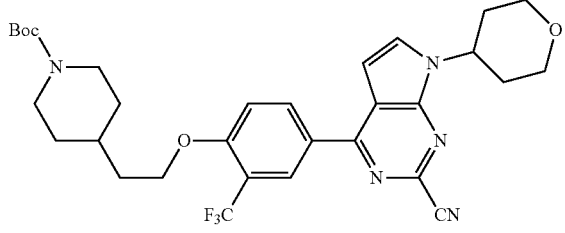
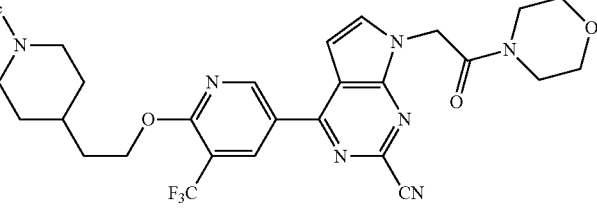
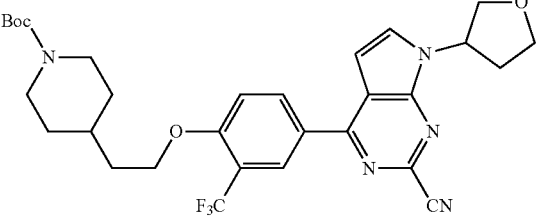
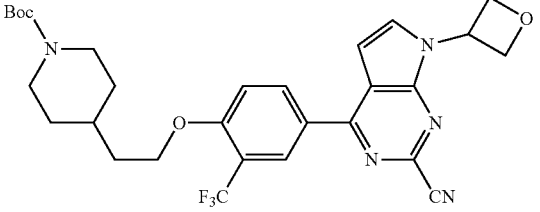
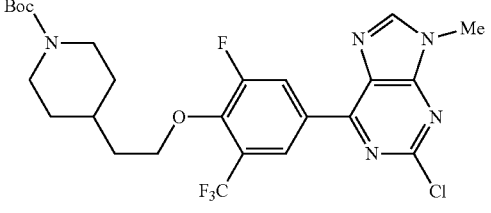
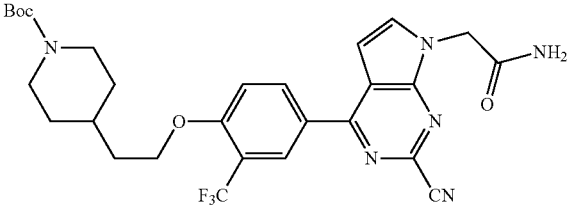
PEX	PSyn	Str	DAT
126	3		ESI+: 600
127	3		ESI+: 644
128	3		ESI+: 586
129	3		ESI+: 572
130	33		ESI+: 558, 560
131	19		ESI+: 573

TABLE 21

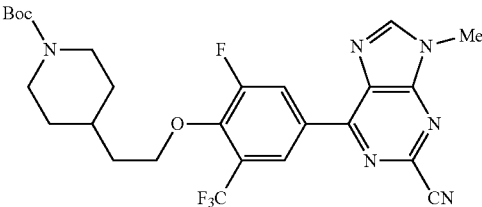
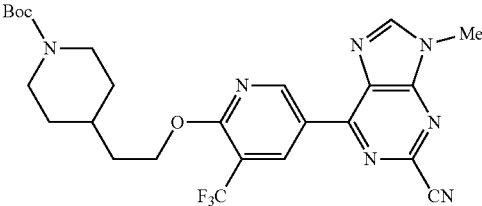
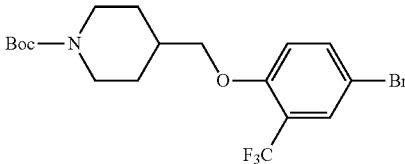
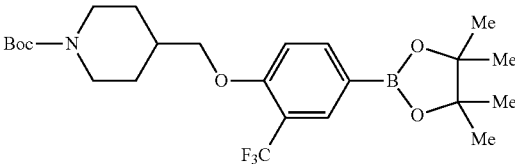
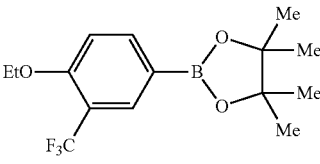
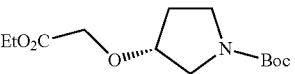
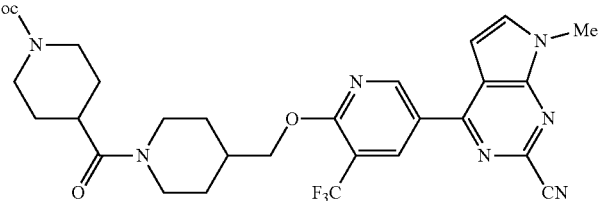
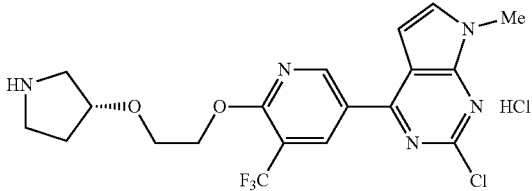
PEX	PSyn	Str	DAT
132	3		ESI+: 549
133	133		ESI+: 554 [M + Na]
134	4		ESI+: 460, 462 [M + Na]
135	27		ESI+: 508 [M + Na]
136	27		NMR2: 1.34 (12H, s), 1.44 (3H, t, J = 7.0 Hz), 4.14 (2H, q, J = 7.0 Hz), 6.96 (1H, d, J = 8.4 Hz), 7.90 (1H, dd, J = 1.3, 8.4 Hz), 7.98-8.01 (1H, m)
137	137	Chiral 	ESI+: 296 [M + Na]
138	138		ESI+: 628
139	139	Chiral 	ESI+: 433

TABLE 22

PEX	PSyn	Str	DAT
140	138		Chiral ESI+: 629
141	10		ESI+: 210
142	9		ESI+: 212
143	11		ESI+: 170
144	2		ESI+: 393, 395
145	33		ESI+: 481, 483
146	28		ESI+: 339, 341 [M - Boc + H]
147	2		ESI+: 409, 411

TABLE 23

PEX	PSyn	Str	DAT
148	33		ESI+: 497, 499
149	137		ESI+: 274
150	30		ESI+: 232
151	2		ESI+: 455, 457
152	2		ESI+: 469, 471
153	33		ESI+: 556, 558
154	33		ESI+: 542, 544
155	3		ESI+: 547

TABLE 24

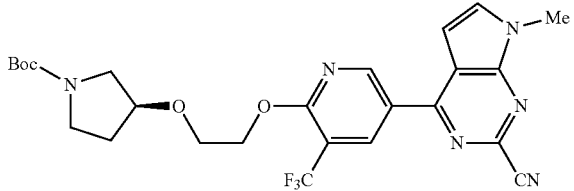
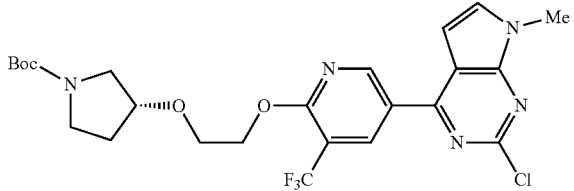
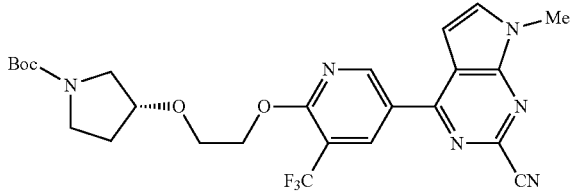
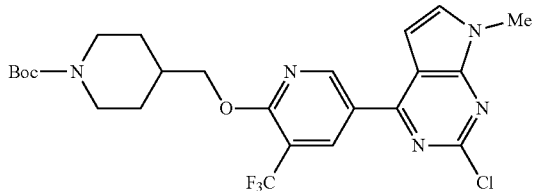
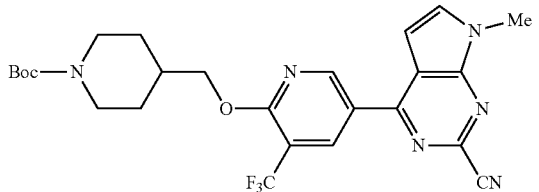
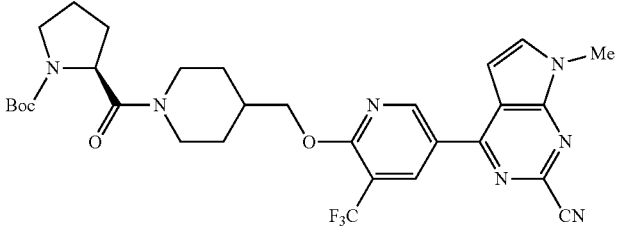
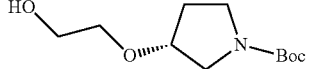
PEX	PSyn	Str	DAT
156	3		Chiral ESI+: 533
157	33		Chiral ESI+: 542, 544
158	3		Chiral ESI+: 533
159	33		ESI+: 526, 528
160	3		ESI+: 517
161	138		Chiral ESI+: 614
162	30		Chiral ESI+: 232

TABLE 24-continued

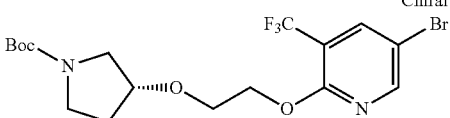
PEX	PSyn	Str	DAT
163	2		Chiral ESI+: 477, 479 [M + Na]

TABLE 25

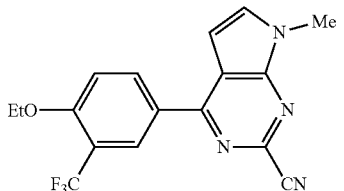
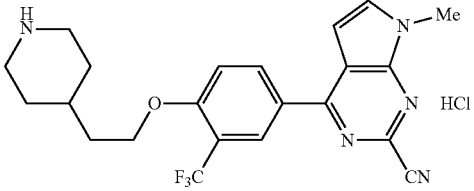
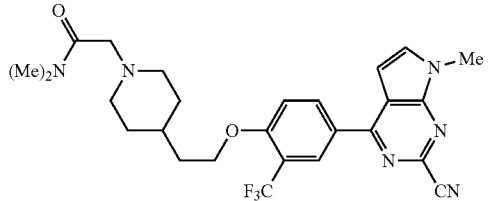
Ex	Str
1	
2	
3	

TABLE 25-continued

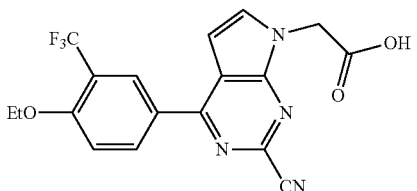
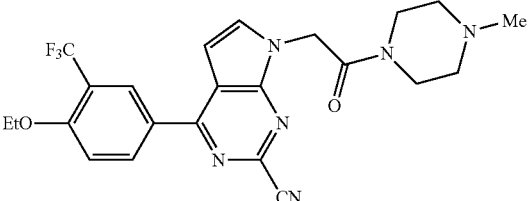
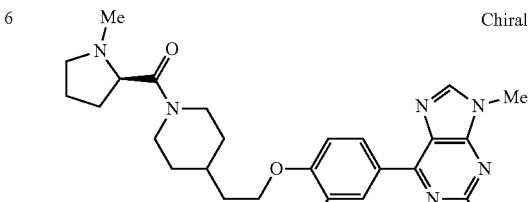
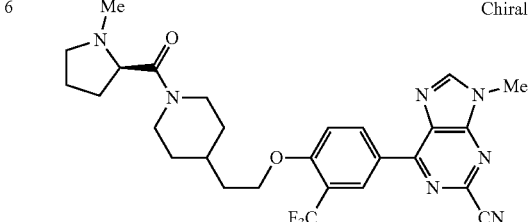
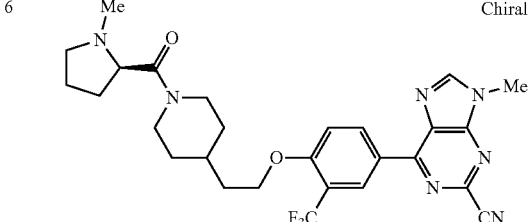
Ex	Str
15	
20	
25	
30	
35	

TABLE 26

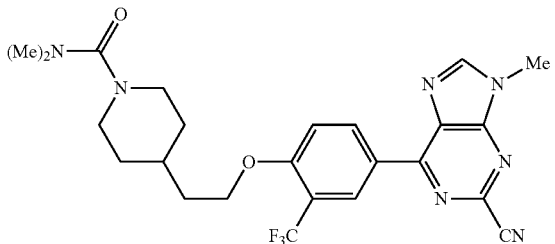
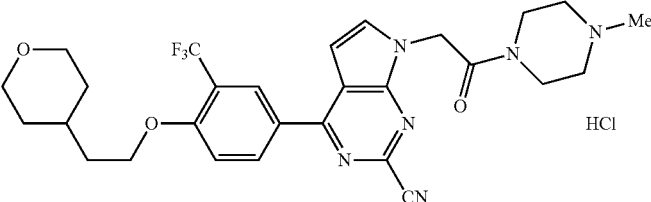
Ex	Str
7	
8	

TABLE 26-continued

Ex	Str
9#	<p>Chemical structure 9# consists of a 4-methylpiperazine ring connected via a propyl chain to a secondary amine group (-NH-). This secondary amine is attached to a benzene ring at the para position. The benzene ring also has a trifluoromethyl group (-F₃C) at the meta position and is connected at the other para position to a pyrazolo[1,5-a]pyrimidine ring system. The pyrazolo ring has an NH group, and the pyrimidine ring has a cyano group (-CN) at the 4-position.</p>
10	<p>Chemical structure 10 consists of a piperazine ring connected via a propyl chain to an ether oxygen atom (-O-). This ether oxygen is attached to a benzene ring at the para position. The benzene ring also has a trifluoromethyl group (-F₃C) at the meta position and is connected at the other para position to a pyrazolo[1,5-a]pyrimidine ring system. The pyrazolo ring has a methyl group (-Me) at the 2-position, and the pyrimidine ring has a cyano group (-CN) at the 4-position. The structure is shown as a hydrochloride salt (HCl).</p>
11	<p>Chemical structure 11 consists of a 4-ethylpiperazine ring connected via a propyl chain to an ether oxygen atom (-O-). This ether oxygen is attached to a benzene ring at the para position. The benzene ring also has a trifluoromethyl group (-F₃C) at the meta position and is connected at the other para position to a pyrazolo[1,5-a]pyrimidine ring system. The pyrazolo ring has a methyl group (-Me) at the 2-position, and the pyrimidine ring has a cyano group (-CN) at the 4-position.</p>
12	<p>Chemical structure 12 consists of a 4-ethylpiperazine ring connected via a propyl chain to an ether oxygen atom (-O-). This ether oxygen is attached to a pyridine ring at the 2-position. The pyridine ring also has a trifluoromethyl group (-F₃C) at the 3-position and is connected at the 4-position to a pyrazolo[1,5-a]pyrimidine ring system. The pyrazolo ring has a methyl group (-Me) at the 2-position, and the pyrimidine ring has a cyano group (-CN) at the 4-position. The structure is shown as a hydrochloride salt (HCl).</p>

TABLE 27

Ex	Str
13	<p>Chemical structure 13 consists of a 4-ethylpiperazine ring connected via a propyl chain to an ether oxygen atom (-O-). This ether oxygen is attached to a benzene ring at the para position. The benzene ring also has a trifluoromethyl group (-F₃C) at the meta position and is connected at the other para position to a pyrazolo[1,5-a]pyrimidine ring system. The pyrazolo ring has a propyl methoxy group (-CH₂CH₂CH₂OMe) at the 2-position, and the pyrimidine ring has a cyano group (-CN) at the 4-position. The structure is shown as a hydrochloride salt (HCl).</p>
14	<p>Chemical structure 14 consists of a piperazine ring with a cyclopropyl group attached to the nitrogen atom. The piperazine ring is connected via a propyl chain to an ether oxygen atom (-O-). This ether oxygen is attached to a benzene ring at the para position. The benzene ring also has a trifluoromethyl group (-F₃C) at the meta position and is connected at the other para position to a pyrazolo[1,5-a]pyrimidine ring system. The pyrazolo ring has a methyl group (-Me) at the 2-position, and the pyrimidine ring has a cyano group (-CN) at the 4-position.</p>

TABLE 27-continued

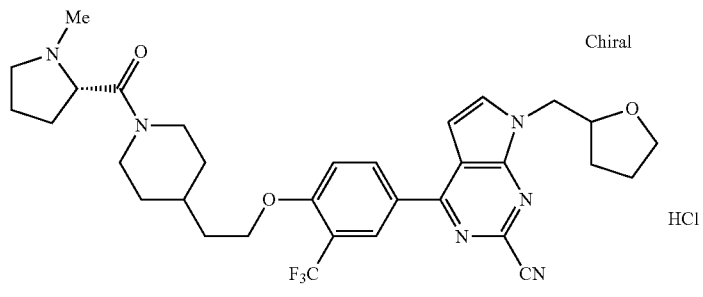
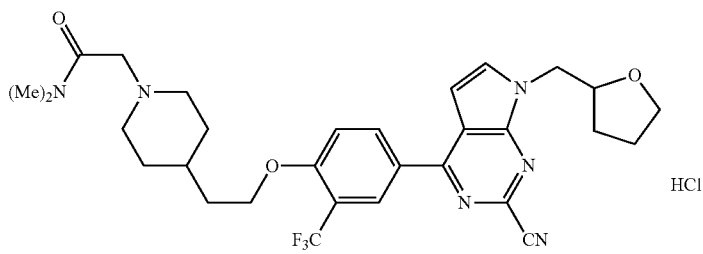
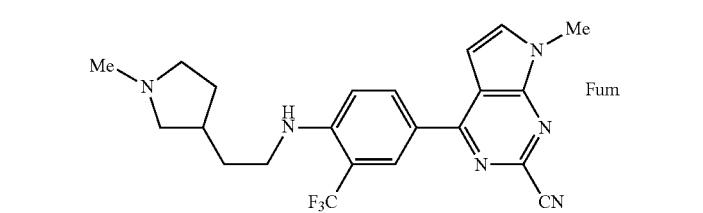
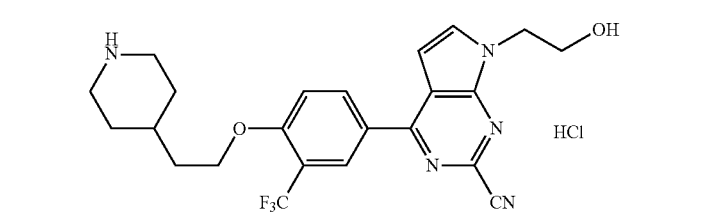
Ex	Str
15	 <p>Chiral HCl</p>
16	 <p>HCl</p>
17#	 <p>Fum</p>
18	 <p>HCl</p>

TABLE 28

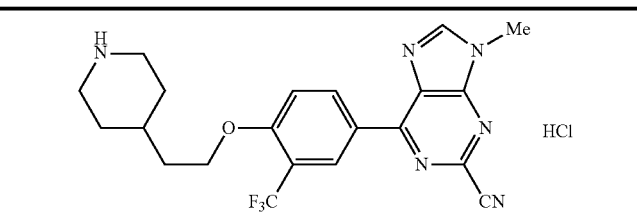
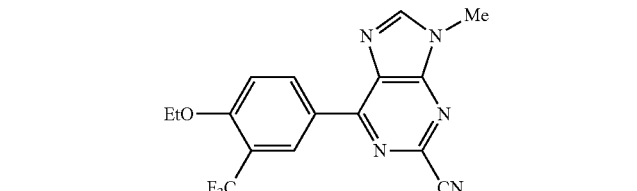
Ex	Str
19	 <p>HCl</p>
20	

TABLE 28-continued

Ex	Str
21	
22#	
23	
24	

TABLE 29

Ex	Str
25	
26	

TABLE 29-continued

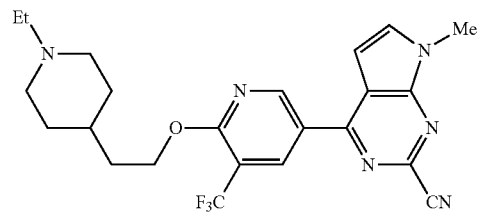
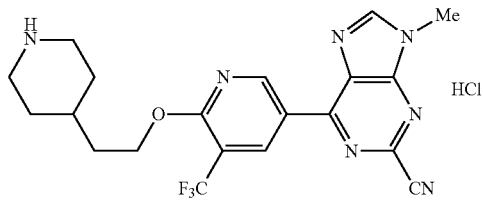
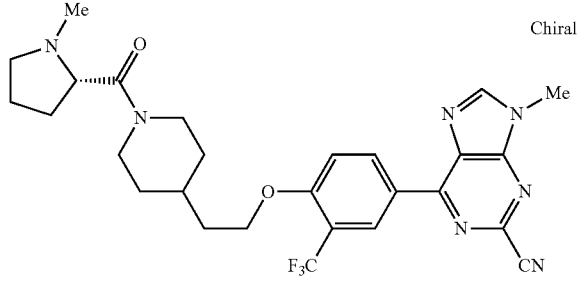
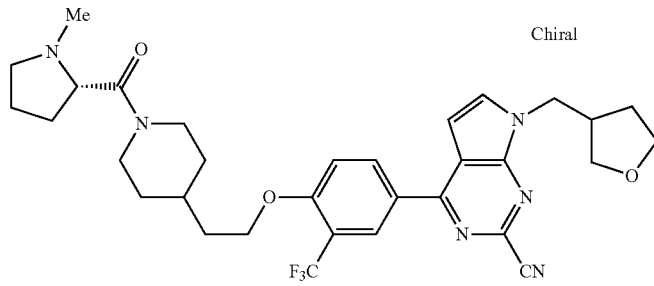
Ex	Str
27	
28	
29	
30	

TABLE 30

45

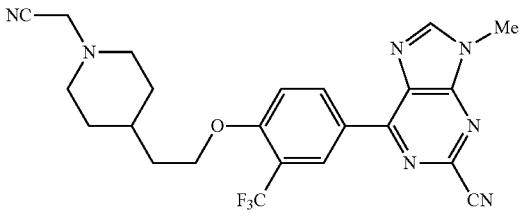
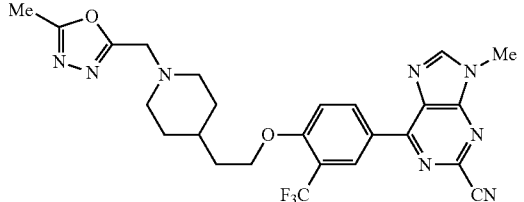
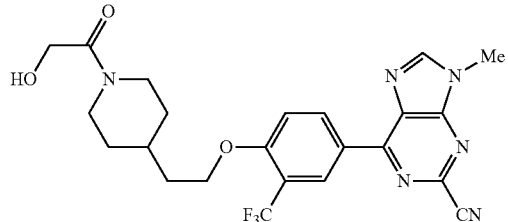
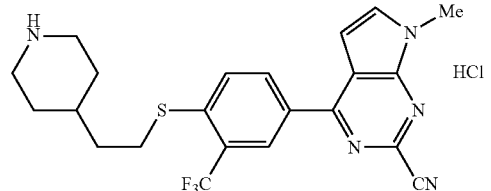
Ex	Str
31	
32	

TABLE 30-continued

50

Ex	Str
33	
34	
50	
55	
60	
65	

105

TABLE 30-continued

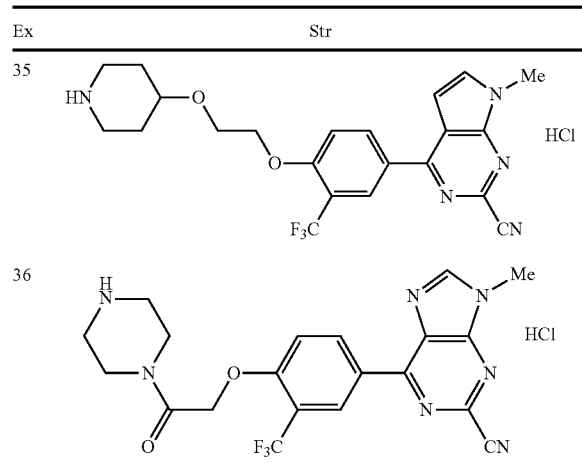
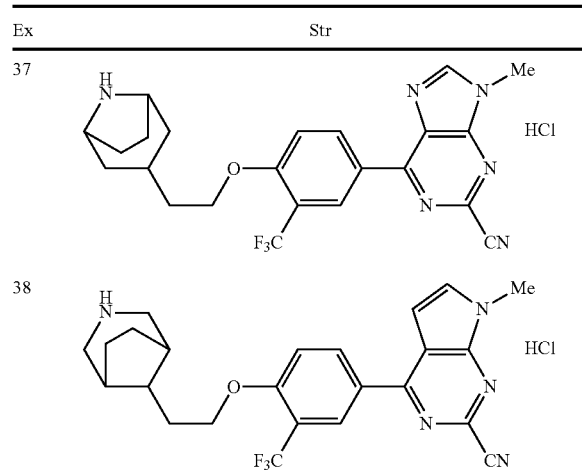


TABLE 31



106

TABLE 31-continued

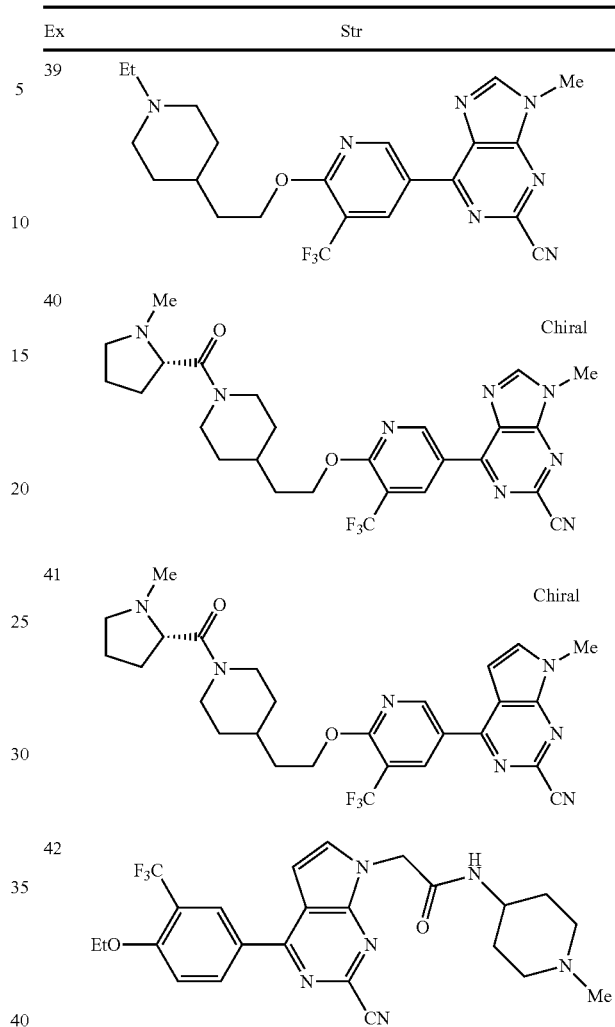


TABLE 32

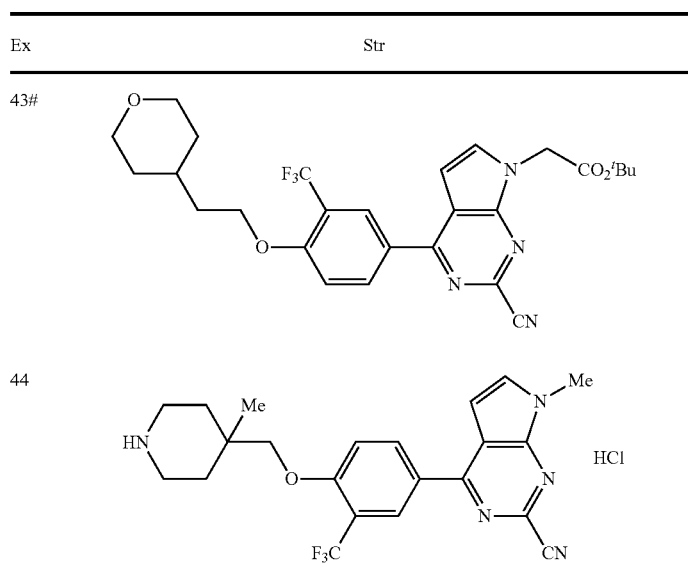


TABLE 32-continued

Ex	Str
45	
46#	
47	
48#	
49#	

TABLE 33

Ex	Str
50	
51	

TABLE 33-continued

Ex	Str
52	
55	
53#	
60	
65	

109

TABLE 33-continued

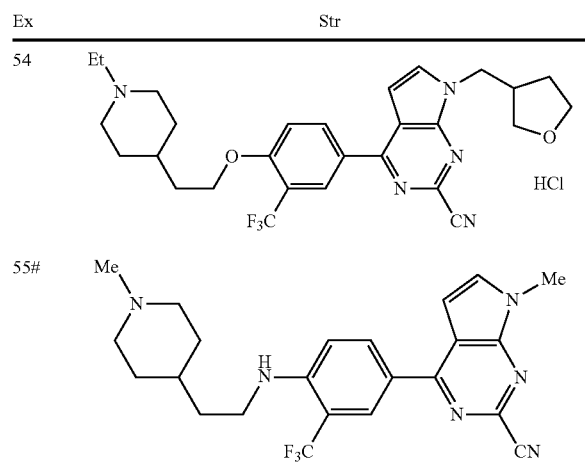
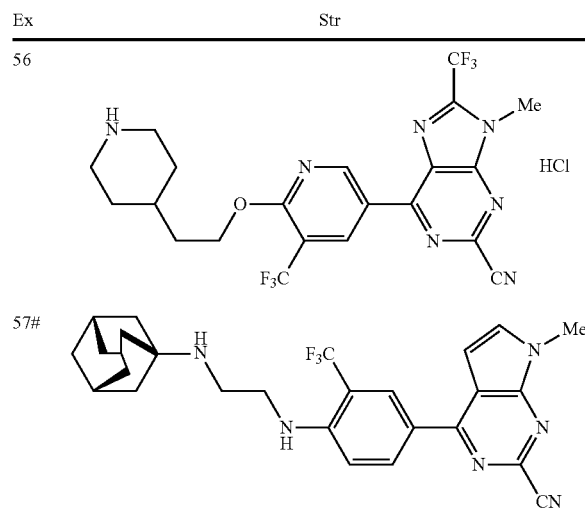


TABLE 34



110

TABLE 34-continued

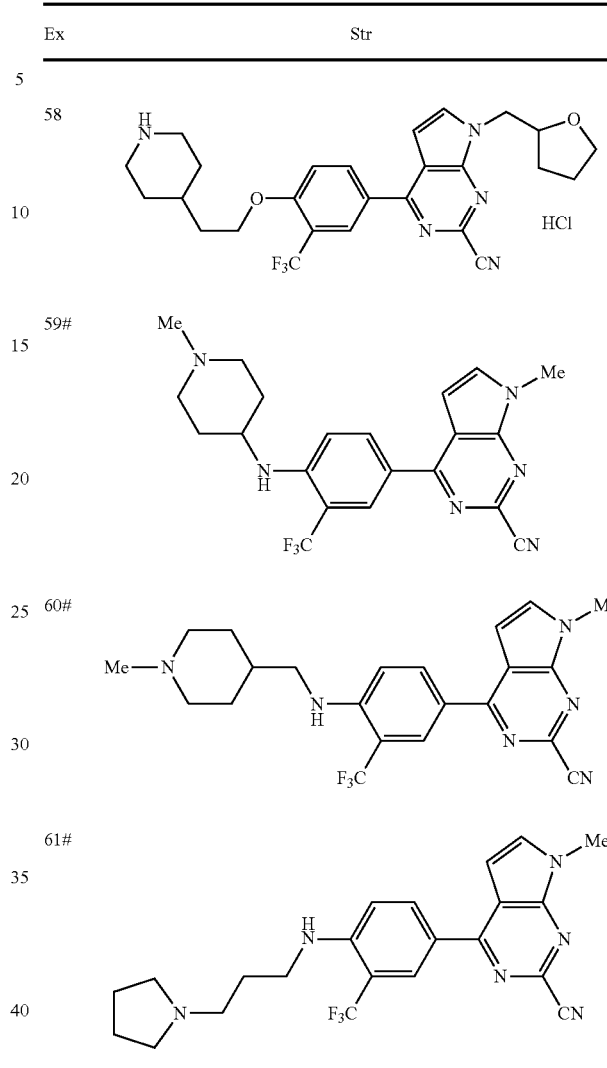
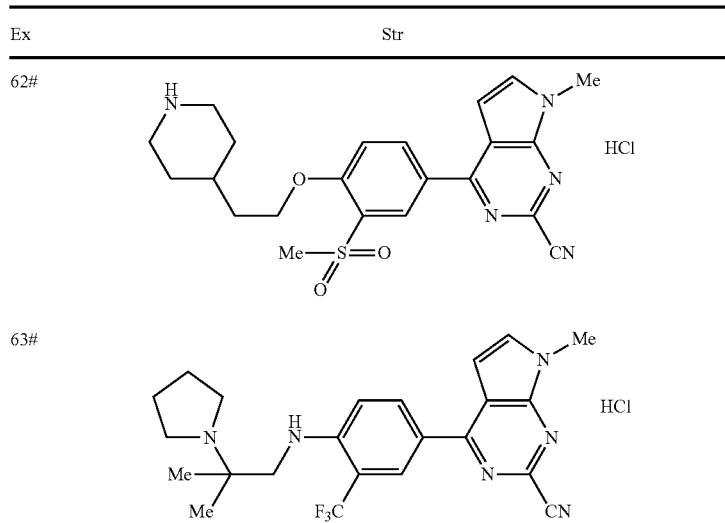


TABLE 35



Ex	Str
64#	<p>Chemical structure 64# shows a piperidine ring with a methyl group on the nitrogen, connected via a propyl chain to the nitrogen of a secondary amine. This secondary amine is attached to a benzene ring, which is also substituted with a trifluoromethyl group (F₃C) and a cyano group (CN). The benzene ring is further connected to a pyrazolo[1,5-a]pyrimidine ring system, which has a methyl group on the nitrogen and a cyano group (CN) at the 4-position.</p>
65	<p>Chemical structure 65 shows a piperidine ring with a hydroxymethyl group (-CH₂OH) on the nitrogen, connected via a propyl chain to the oxygen of an ether linkage. This ether is attached to a benzene ring, which is also substituted with a trifluoromethyl group (F₃C) and a cyano group (CN). The benzene ring is further connected to a pyrazolo[1,5-a]pyrimidine ring system, which has a methyl group on the nitrogen and a cyano group (CN) at the 4-position. The pyrazolo[1,5-a]pyrimidine ring is also connected to a tetrahydrofuran ring via a propyl chain. The structure is shown as a hydrochloride salt (HCl).</p>
66	<p>Chemical structure 66 shows a piperidine ring with a dimethylamino group (-NMe₂) on the nitrogen, connected via a propyl chain to the oxygen of an ether linkage. This ether is attached to a benzene ring, which is also substituted with a trifluoromethyl group (F₃C) and a cyano group (CN). The benzene ring is further connected to a pyrazolo[1,5-a]pyrimidine ring system, which has a methyl group on the nitrogen and a cyano group (CN) at the 4-position. The pyrazolo[1,5-a]pyrimidine ring is also connected to a tetrahydrofuran ring via a propyl chain. The structure is shown as a hydrochloride salt (HCl).</p>
67#	<p>Chemical structure 67# shows a piperidine ring with a methyl group on the nitrogen, connected via a propyl chain to the nitrogen of a secondary amine. This secondary amine is attached to a benzene ring, which is also substituted with a trifluoromethyl group (F₃C) and a cyano group (CN). The benzene ring is further connected to a pyrazolo[1,5-a]pyrimidine ring system, which has a methyl group on the nitrogen and a cyano group (CN) at the 4-position. The pyrazolo[1,5-a]pyrimidine ring is also connected to a tetrahydrofuran ring via a propyl chain.</p>

TABLE 36

Ex	Str
68	<p>Chemical structure 68 shows a piperidine ring connected via a propyl chain to the oxygen of an ether linkage. This ether is attached to a benzene ring, which is also substituted with a trifluoromethyl group (F₃C) and a cyano group (CN). The benzene ring is further connected to a pyrazolo[1,5-a]pyrimidine ring system, which has a hydrogen atom on the nitrogen and a cyano group (CN) at the 4-position. The structure is shown as a hydrochloride salt (HCl).</p>
69#	<p>Chemical structure 69# shows a piperidine ring with a methyl group on the nitrogen, connected via a propyl chain to the nitrogen of a secondary amine. This secondary amine is attached to a benzene ring, which is also substituted with a trifluoromethyl group (F₃C) and a cyano group (CN). The benzene ring is further connected to a pyrazolo[1,5-a]pyrimidine ring system, which has a methyl group on the nitrogen and a cyano group (CN) at the 4-position. The pyrazolo[1,5-a]pyrimidine ring is also connected to a tetrahydrofuran ring via a propyl chain. The structure is labeled as Chiral.</p>

TABLE 36-continued

Ex	Str
70	
71	
72	
73	
74	

TABLE 37

Ex	Str
75	

TABLE 37-continued

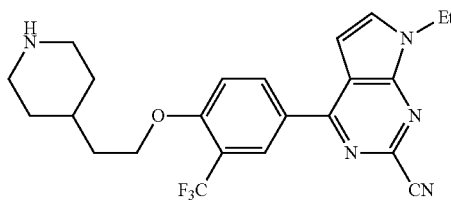
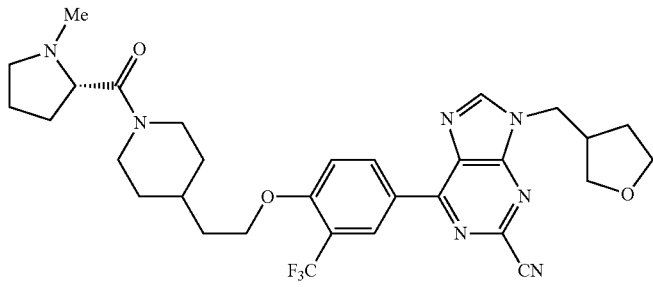
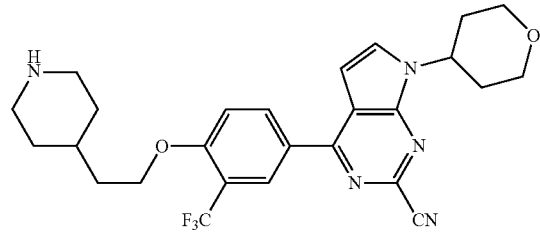
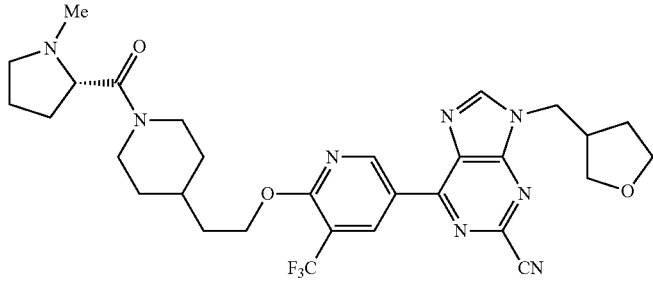
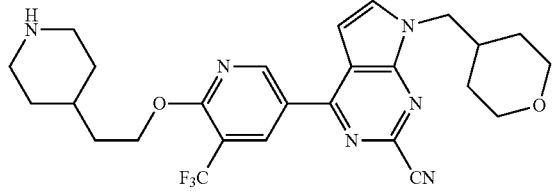
Ex	Str
76	 <p>HCl</p>
77	 <p>Chiral</p>
78	 <p>HCl</p>
79	 <p>Chiral</p>
80	 <p>HCl</p>

TABLE 38

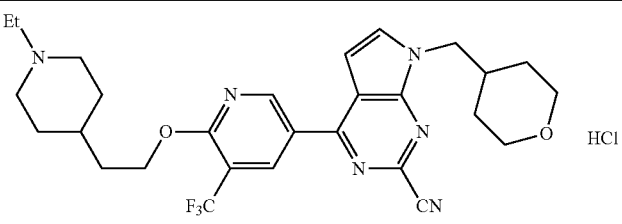
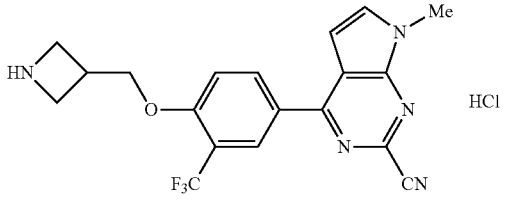
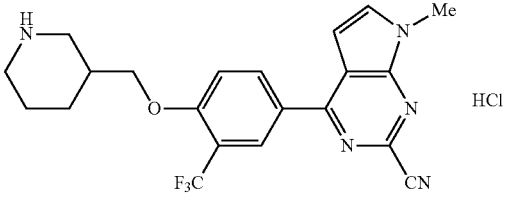
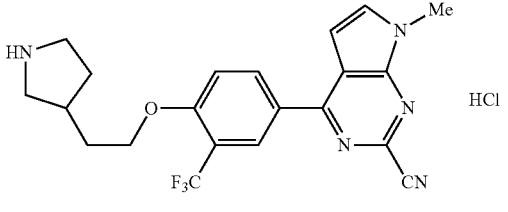
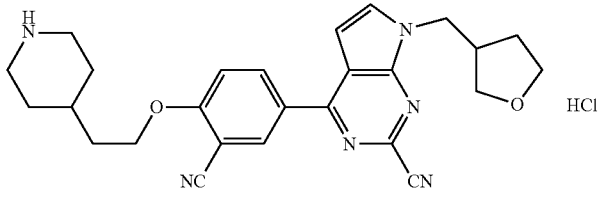
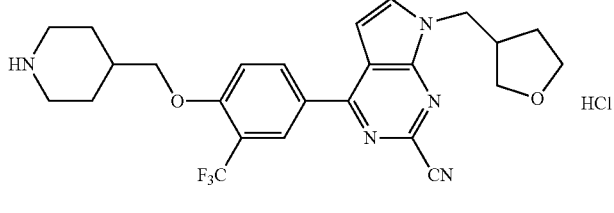
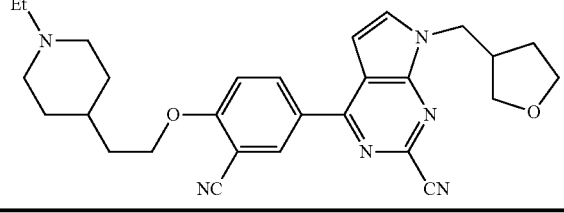
Ex	Str
81	 HCl
82	 HCl
83	 HCl
84	 HCl
85	 HCl
86	 HCl
87	

TABLE 39

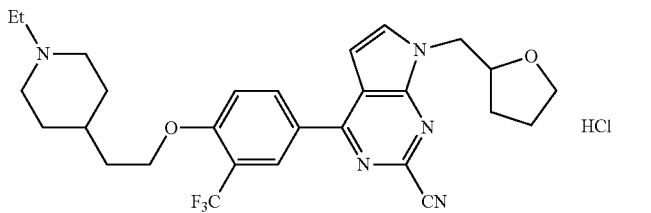
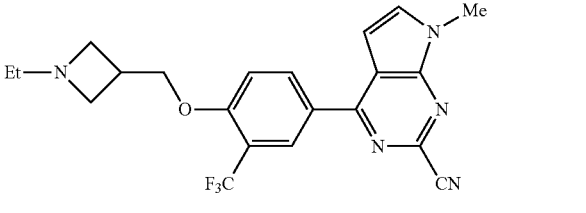
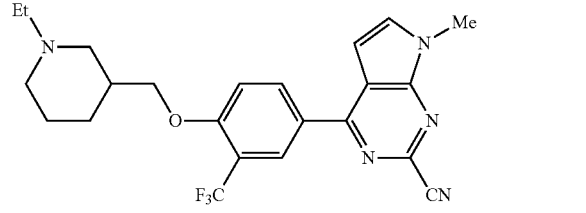
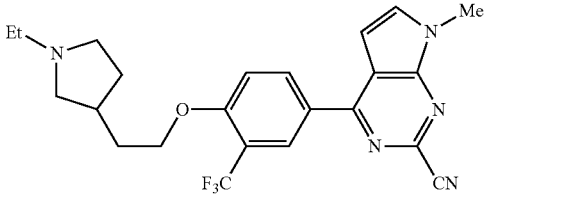
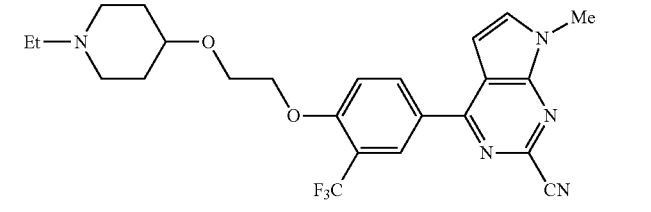
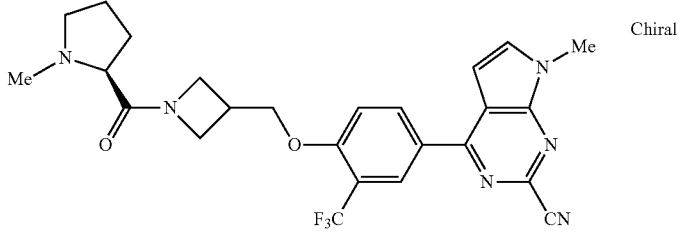
Ex	Str
88	 <chem>CCN1CCOCC1COc2ccc(C(F)(F)F)cc2-c3nc4c(ncn3COC5OCCO5)C#N.Cl</chem>
89	 <chem>CCN1CC1COc2ccc(C(F)(F)F)cc2-c3nc4c(ncn3C5=CN(C)C=C5)C#N</chem>
90	 <chem>CCN1CCOCC1COc2ccc(C(F)(F)F)cc2-c3nc4c(ncn3C5=CN(C)C=C5)C#N</chem>
91	 <chem>CCN1CCOCC1COc2ccc(C(F)(F)F)cc2-c3nc4c(ncn3C5=CN(C)C=C5)C#N</chem>
92	 <chem>CCN1CCOCC1COCCOc2ccc(C(F)(F)F)cc2-c3nc4c(ncn3C5=CN(C)C=C5)C#N</chem>
93	 <chem>C[C@@H](C1CCN1)C(=O)N2CCN2COc3ccc(C(F)(F)F)cc3-c4nc5c(ncn4C6=CN(C)C=C6)C#N</chem>

TABLE 40

Ex	Str
94	<p>Chiral</p>
95	<p>Chiral</p>
96	<p></p>
97	<p>HCl</p>
98#	<p></p>
99#	<p></p>

TABLE 41

Ex	Str
100#	<p>Chemical structure 100# consists of a central benzene ring substituted at the 1 and 4 positions with a trifluoromethyl group (F₃C) and a 4-methyl-1H-imidazole-2-yl group, respectively. At the 3 position of the benzene ring, there is a 2-(2-(pyrrolidin-2-yl)ethyl)amino group.</p>
101	<p>Chemical structure 101 consists of a central benzene ring substituted at the 1 and 4 positions with a trifluoromethyl group (F₃C) and a 4-(2-methyl-1H-imidazol-5-yl) group, respectively. At the 3 position of the benzene ring, there is a 2-(2-(piperidin-4-yl)ethoxy) group. The imidazole ring is substituted at the 2 position with a tetrahydrofuran-2-yl group. The structure is shown as a hydrochloride salt (HCl).</p>
102	<p>Chemical structure 102 consists of a central benzene ring substituted at the 1 and 4 positions with a trifluoromethyl group (F₃C) and a 4-(2-methyl-1H-imidazol-5-yl) group, respectively. At the 3 position of the benzene ring, there is a 2-(2-(1-ethylpiperidin-4-yl)ethoxy) group. The imidazole ring is substituted at the 2 position with a 2-aminoethyl group.</p>
103	<p>Chemical structure 103 consists of a central benzene ring substituted at the 1 and 4 positions with a trifluoromethyl group (F₃C) and a 4-(2-methyl-1H-imidazol-5-yl) group, respectively. At the 3 position of the benzene ring, there is a 2-(2-(1-ethylpiperidin-4-yl)ethoxy) group. The imidazole ring is substituted at the 2 position with an ethyl group. The structure is shown as a fumarate salt (Fum).</p>
104	<p>Chemical structure 104 consists of a central benzene ring substituted at the 1 and 4 positions with a trifluoromethyl group (F₃C) and a 4-(2-methyl-1H-imidazol-5-yl) group, respectively. At the 3 position of the benzene ring, there is a 2-(2-(1-methylpyrrolidin-2-yl)ethoxy) group. The imidazole ring is substituted at the 2 position with a tetrahydrofuran-2-yl group. The structure is labeled as Chiral.</p>
105	<p>Chemical structure 105 consists of a central benzene ring substituted at the 1 and 4 positions with a trifluoromethyl group (F₃C) and a 4-(2-methyl-1H-imidazol-5-yl) group, respectively. At the 3 position of the benzene ring, there is a 2-(2-(1-methylpiperidin-4-yl)ethoxy) group. The imidazole ring is substituted at the 2 position with a tetrahydrofuran-2-yl group. The structure is labeled as Chiral.</p>

TABLE 42

Ex	Str
106	 <chem>CCN1CCCCC1CCOC2=CC=C(C=C2)C3=C(C=C3)N4C=NC(=N4)C5=CC=CN5CC6OCCO6</chem> Fum
107	 <chem>C1CCNCC1CCOC2=CC=NC(=C2)C3=C(C=C3)N4C=NC(=N4)C5=CC=CN5CC(=O)N6CCOCC6</chem> HCl
108	 <chem>OCC1CCN(C1)CCOC2=CC=C(C=C2)C3=C(C=C3)N4C=NC(=N4)C5=CC=CN5CC6OCCO6</chem> HCl
109	 <chem>CCN1CCCCC1CCOC2=CC=NC(=C2)C3=C(C=C3)N4C=NC(=N4)C5=CC=CN5CC(=O)N6CCOCC6</chem> HCl
110	 <chem>CCN1CCCCC1CCOC2=CC=C(C=C2)C3=C(C=C3)N4C=NC(=N4)C5=CC=CN5CCCO</chem> HCl
111	 <chem>CC(C)N1CCCCC1CCOC2=CC=C(C=C2)C3=C(C=C3)N4C=NC(=N4)C5=CC=CN5CC6OCCO6</chem>

TABLE 43

Ex	Str
112	 <chem>CC1(CCN1)CCOC2=CC=C(C=C2)C(=N3C=NC(=N3)C#N)C(=N4C=NC(=N4)C#N</chem>
113	 <chem>CC1(CCN1)CCOC2=CC=C(C=C2)C(=N3C=NC(=N3)C#N)C(=N4C=NC(=N4)C#N)C(F)=C5C=C(C=C5)C(F)(F)F</chem> HCl
114	 <chem>CCN1CC(C)CC1CCOC2=CC=C(C=C2)C(=N3C=NC(=N3)C#N)C(=N4C=NC(=N4)C#N)C(F)=C5C=C(C=C5)C(F)(F)F</chem> Fum
115	 <chem>CC1(CCN1)CCOC2=CC=C(C=C2)C(=N3C=NC(=N3)C#N)C(=N4C=NC(=N4)C#N)C(F)=C5C=C(C=C5)C(F)(F)F</chem> Chiral Fum
116	 <chem>CC1(CCN1)CCOC2=CC=NC=C2C(=N3C=NC(=N3)C#N)C(=N4C=NC(=N4)C#N)C(F)(F)F</chem> HCl
117	 <chem>CC1(CCN1)CCOC2=CC=C(C=C2)C(=N3C=NC(=N3)C#N)C(=N4C=NC(=N4)C#N)C(F)(F)F</chem> Chiral

TABLE 44

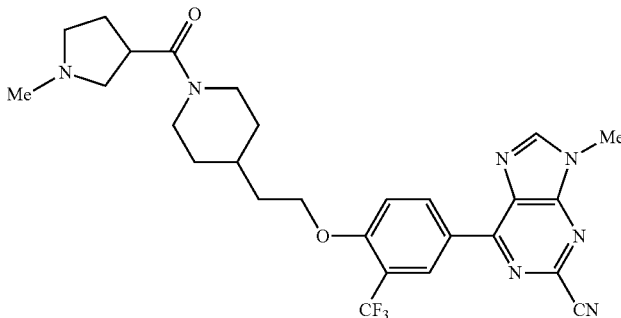
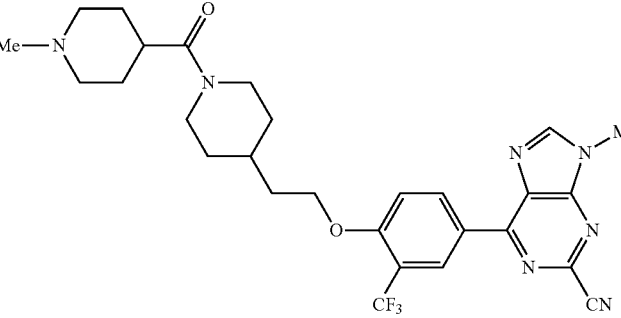
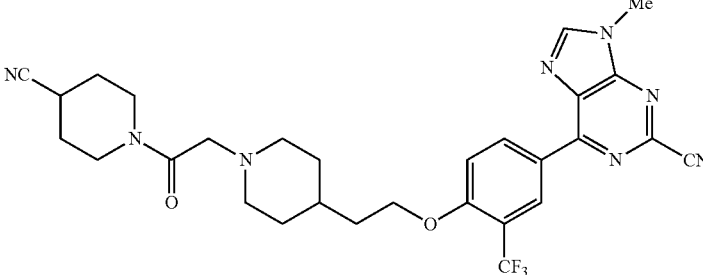
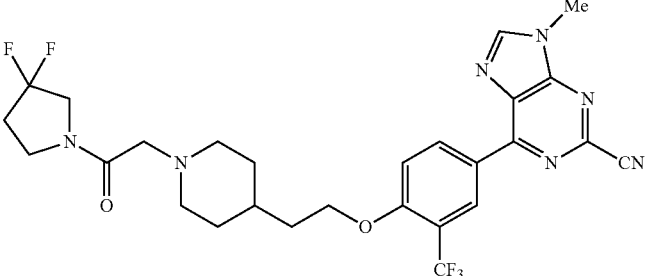
Ex	Str
118	
119	
120	
121	

TABLE 45

Ex	Str
122	<p>Chiral</p>
123	
124	
125	
126	
127	

TABLE 46

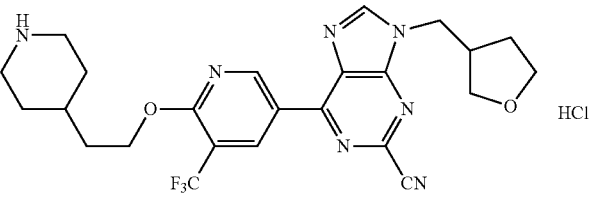
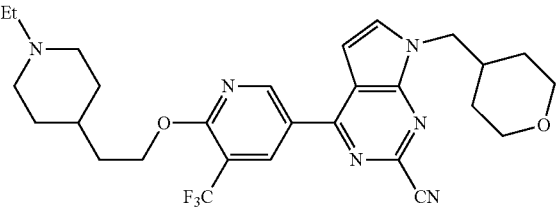
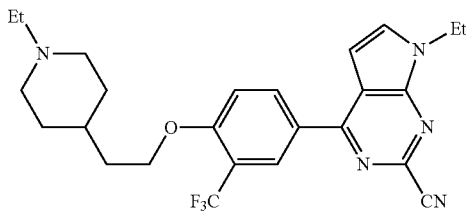
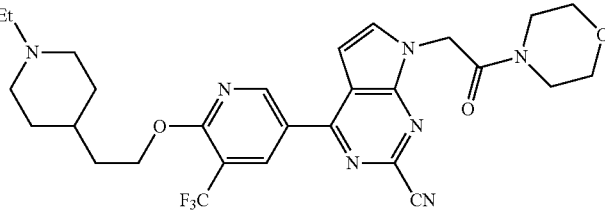
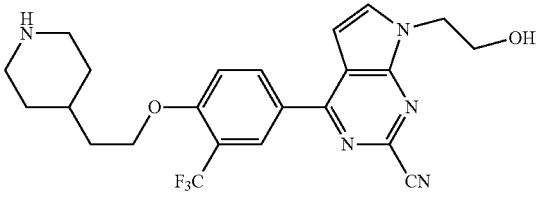
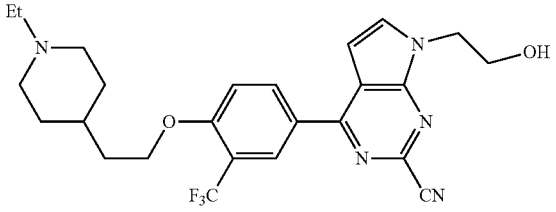
Ex	Str
128	 <chem>C1CCN(C1)CCCOc2nc(C(F)(F)F)c(C3=NC(=NC4=CN(C4)C5CCOC5)N3)nc6nccn6</chem> HCl
129	 <chem>CCN1CCOCC1CCCOc2nc(C(F)(F)F)c(C3=NC(=NC4=CN(C4)C5CCOC5)N3)nc6nccn6</chem>
130	 <chem>CCN1CCOCC1CCCOc2cc(C(F)(F)F)c(C3=NC(=NC4=CN(C4)C5CCOC5)N3)nc6nccn6C</chem>
131	 <chem>CCN1CCOCC1CCCOc2nc(C(F)(F)F)c(C3=NC(=NC4=CN(C4)C5CCOC(=O)N6CCOCC6)N3)nc6nccn6</chem>
132	 <chem>C1CCN(C1)CCCOc2nc(C(F)(F)F)c(C3=NC(=NC4=CN(C4)C5CCO)N3)nc6nccn6</chem>
133	 <chem>CCN1CCOCC1CCCOc2nc(C(F)(F)F)c(C3=NC(=NC4=CN(C4)C5CCO)N3)nc6nccn6</chem>

TABLE 47

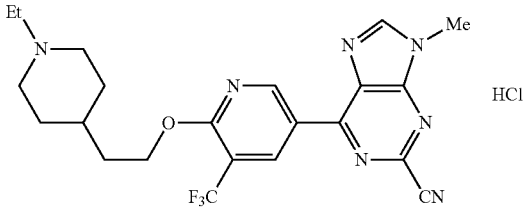
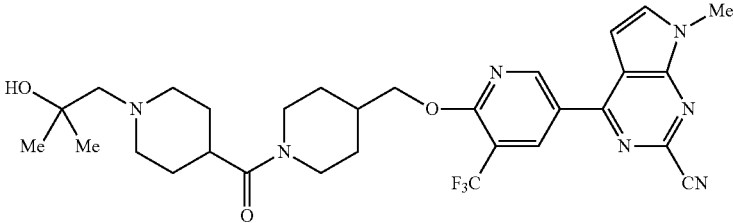
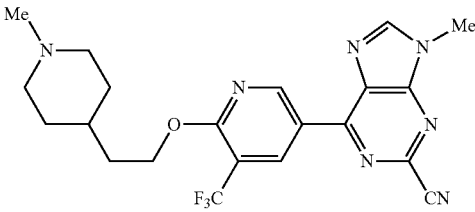
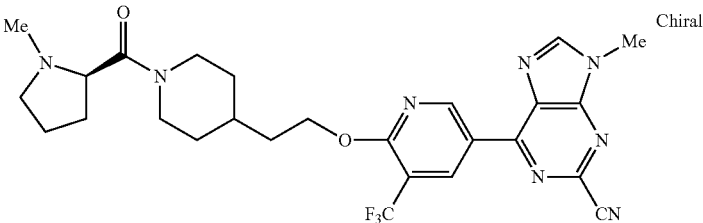
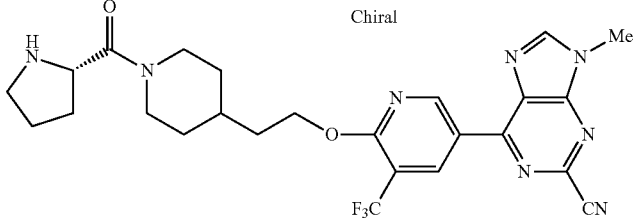
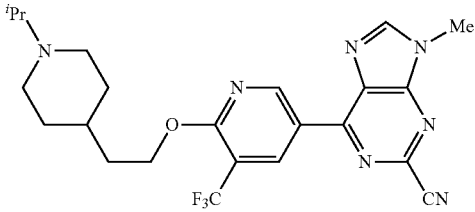
Ex	Str
134	 <p style="text-align: right; margin-right: 20px;">HCl</p>
135	
136	
137	 <p style="text-align: right; margin-right: 20px;">Chiral</p>
138	 <p style="text-align: right; margin-right: 20px;">Chiral</p>
139	

TABLE 48

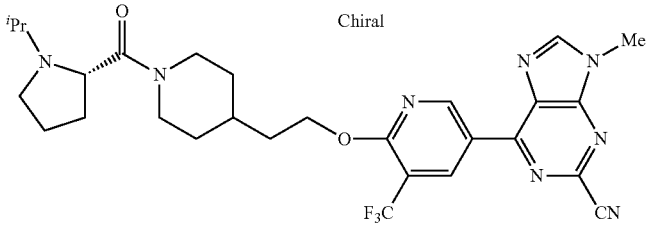
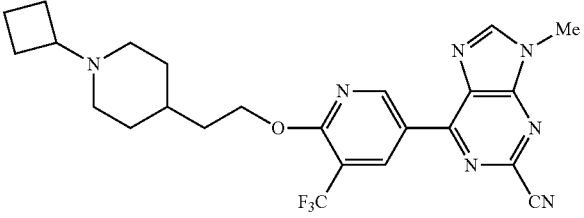
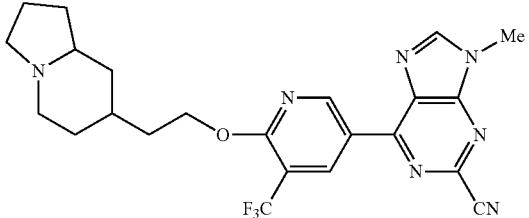
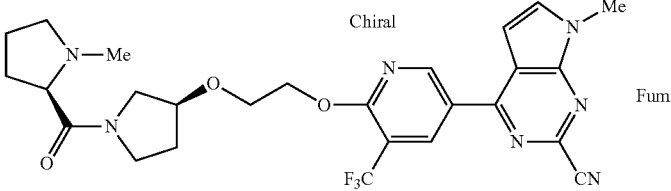
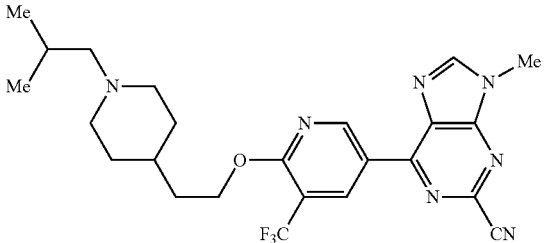
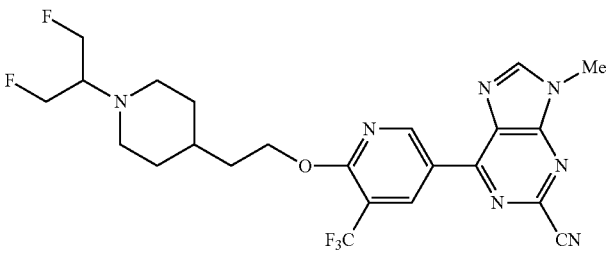
Ex	Str
140	<p>Chiral</p> 
141	
142	
143	<p>Chiral</p> <p>Fum</p> 
144	
145	

TABLE 49

Ex	Str
146	 <chem>CC1(C)CCN(C1)CCCC2=CN=C(C=C2C3=NC(=NC=C3N)C#N)C(F)(F)F</chem>
147	 <chem>CN1CCCCC1OCCCO=C2N=C(C=C2C3=NC(=NC=C3N)C#N)C(F)(F)F</chem> HCl
148	 <chem>CN1C[C@H](OCCCO=C2N=C(C=C2C3=NC(=NC=C3N)C#N)C(F)(F)F)C1</chem> Chiral HCl
149	 <chem>CN1C[C@H](OCCCO=C2N=C(C=C2C3=NC(=NC=C3N)C#N)C(F)(F)F)C(=O)N1C2CCCC2</chem> Chiral
150	 <chem>CN1C[C@H](OCCCO=C2N=C(C=C2C3=NC(=NC=C3N)C#N)C(F)(F)F)C(=O)N1</chem> Chiral
151	 <chem>CN(C)C(=O)CN1CCCCC1OCCCO=C2N=C(C=C2C3=NC(=NC=C3N)C#N)C(F)(F)F</chem>

TABLE 50

Ex	Str
152	<p>HCl</p>
153	<p>HCl</p>
154	<p>Chiral HCl</p>
155	<p>HCl</p>
156	<p>Chiral</p>
157	<p>Chiral</p>

TABLE 52-continued

Ex	Syn	DAT
		4.26-4.35 (2H, m), 7.55 (1H, d, J = 8.9 Hz), 8.37 (1H, br), 8.64 (1H, br), 9.08 (1H, dd, J = 2.2, 8.9 Hz), 9.11 (1H, d, J = 2.2 Hz); ESI+: 445
11	11	NMR2: 1.09 (3H, t, J = 7.2 Hz), 1.29-1.42 (2H, m), 1.55-1.68 (1H, m), 1.72-1.96 (6H, m), 2.40 (2H, q, J = 7.2 Hz), 2.75 (3H, s), 2.92-3.00 (2H, m), 3.86 (3H, s), 4.21 (2H, t, J = 6.3 Hz), 7.14 (1H, d, J = 8.8 Hz), 9.04 (1H, dd, J = 2.2, 8.8 Hz), 9.16 (1H, d, J = 2.2 Hz); ESI+: 473
12	12	NMR1: 1.13-1.27 (4H, m), 1.64-1.92 (5H, m), 1.97-2.10 (1H, m), 2.57-2.69 (1H, m), 2.70-2.83 (1H, m), 2.99-3.13 (2H, m), 3.37-3.53 (2H, m), 3.94 (3H, s), 4.53-4.71 (2H, m), 7.23 (1H, d, J = 3.6 Hz), 8.09 (1H, d, J = 3.6 Hz), 8.73 (1H, d, J = 2.2 Hz), 9.25 (1H, d, J = 2.2 Hz), 9.58 (1H, br); ESI+: 459

TABLE 53

Ex	Syn	DAT
13	13	NMR1: 1.23 (3H, t, J = 7.3 Hz), 1.44-1.59 (2H, m), 1.70-1.84 (2H, m), 1.88-2.01 (3H, m), 2.77-2.90 (2H, m), 3.02-3.11 (2H, m), 3.25 (3H, s), 3.43-3.52 (2H, m), 3.76 (2H, t, J = 5.2 Hz), 4.29-4.36 (2H, m), 4.53 (2H, t, J = 5.2 Hz), 7.12 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.8 Hz), 8.07 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 3.6 Hz), 8.48 (1H, dd, J = 2.1, 8.8 Hz), 9.68 (1H, br); ESI+: 502
14	14	NMR1: 0.22-0.30 (2H, m), 0.35-0.42 (2H, m), 1.06-1.22 (2H, m), 1.44-1.60 (2H, m), 1.62-1.76 (4H, m), 2.04-2.16 (2H, m), 2.87-2.96 (2H, m), 3.92 (3H, s), 4.28 (2H, t, J = 6.4 Hz), 7.11 (1H, d, J = 3.6 Hz), 7.50 (1H, d, J = 8.9 Hz), 8.04 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.2 Hz), 8.46 (1H, dd, J = 2.2, 8.8 Hz); ESI+: 470
15	15	NMR1: 1.06-1.34 (4H, m), 1.54-1.67 (1H, m), 1.72-2.16 (5H, m), 2.45-2.56 (4H, m), 2.65-2.83 (4H, m), 3.01-3.15 (2H, m), 3.54-3.73 (3H, m), 3.75-3.84 (1H, m), 4.20-4.66 (8H, m), 7.12 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 8.8 Hz), 8.06 (1H, d, J = 3.7 Hz), 8.39 (1H, d, J = 2.2 Hz), 8.48 (1H, dd, J = 2.2, 8.8 Hz), 9.43-9.60 (1H, br); ESI+: 611
16	16	NMR1: 1.50-1.68 (3H, m), 1.71-2.05 (8H, m), 2.86-3.05 (8H, m), 3.43-3.54 (2H, m), 3.60-3.69 (1H, m), 3.75-3.83 (1H, m), 4.16-4.49 (7H, m), 7.12 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.9 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.2 Hz), 8.49 (1H, dd, J = 2.1, 8.7 Hz), 9.31-9.48 (1H, br); ESI+: 585
17#	17	NMR1: 1.48-1.61 (1H, m), 1.63-1.78 (2H, m), 2.03-2.17 (1H, m), 2.23-2.37 (1H, m), 2.55 (3H, s), 2.85-3.02 (2H, m), 3.10-3.20 (1H, m), 3.27-3.38 (3H, m), 3.90 (3H, s), 6.26-6.33 (1H, m), 6.54 (2H, s), 7.01-7.06 (1H, m), 7.07 (1H, d, J = 3.6 Hz), 7.96 (1H, d, J = 3.6 Hz), 8.28-8.34 (2H, m); ESI+: 429
18	18	ESI+: 460
19	2	NMR1: 1.33-1.48 (2H, m), 1.71-1.95 (5H, m), 2.76-2.91 (2H, m), 3.20-3.38 (2H, m), 3.93 (3H, s), 4.27-4.38 (2H, m), 7.56 (1H, d, J = 9.0 Hz), 8.49 (1H, br), 8.73 (1H, br), 8.92 (1H, s), 9.06 (1H, dd, J = 2.2, 9.0 Hz), 9.17 (1H, d, J = 2.2 Hz); ESI+: 431
20	1	NMR2: 1.51 (3H, t, J = 7.0 Hz), 3.98 (3H, s), 4.25 (2H, q, J = 7.0 Hz), 7.16 (1H, d, J = 8.8 Hz), 8.25 (1H, s), 9.06 (1H, dd, J = 2.2, 8.8 Hz), 9.21 (1H, d, J = 2.2 Hz); ESI+: 348
21	3	NMR2: 1.32-1.44 (2H, m), 1.52-1.67 (1H, m), 1.69-1.86 (4H, m), 2.04-2.14 (2H, m), 2.88-2.92 (2H, m), 2.94 (3H, s), 3.08 (3H, s), 3.15 (2H, s), 4.07 (2H, t, J = 5.0 Hz), 4.20 (2H, t, J = 6.4 Hz), 4.51 (2H, t, J = 5.0 Hz), 6.91 (1H, d, J = 3.6 Hz), 7.16 (1H, d, J = 8.8 Hz), 7.63 (1H, d, J = 3.6 Hz), 8.35 (1H, dd, J = 2.2, 8.8 Hz), 8.40 (1H, d, J = 2.2 Hz); ESI+: 545
22#	1	NMR2: 1.50 (9H, s), 1.51 (3H, t, J = 7.0 Hz), 4.25 (2H, q, J = 7.0 Hz), 5.01 (2H, s), 6.96 (1H, d, J = 3.6 Hz), 7.17 (1H, d, J = 8.4 Hz), 7.53 (1H, d, J = 3.6 Hz), 8.35 (1H, dd, J = 2.4, 8.4 Hz), 8.41 (1H, d, J = 2.4 Hz); ESI+: 447
23	10	NMR1: 1.34-1.49 (2H, m), 1.59-1.69 (1H, m), 1.73-1.97 (6H, m), 2.78-2.93 (3H, m), 3.21-3.32 (2H, m), 3.49-3.55 (1H, m), 3.61-3.69 (2H, m), 3.79-3.86 (1H, m), 4.27-4.38 (4H, m), 7.14 (1H, d, J = 3.6 Hz), 7.50 (1H, d, J = 8.8 Hz), 8.16 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.2 Hz), 8.40-8.56 (2H, m), 8.73 (1H, br); ESI+: 500
24	3	NMR1: 1.16-1.30 (2H, m), 1.44-1.57 (1H, m), 1.64-1.76 (4H, m), 1.86-2.01 (2H, m), 2.31-2.42 (2H, m), 2.78-2.95 (2H, m), 3.44-3.52 (2H, m), 3.92 (3H, s), 4.24-4.39 (3H, m), 7.56 (1H, d, J = 9.0 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.2, 8.9 Hz), 9.17 (1H, d, J = 2.1 Hz); ESI+: 475

TABLE 54

Ex	Syn	DAT
25	3	NMR1: 1.14-1.31 (2H, m), 1.42-1.56 (1H, m), 1.64-1.79 (6H, m), 1.80-1.88 (2H, m), 1.96-2.06 (2H, m), 2.79-2.89 (2H, m), 3.04 (2H, s), 3.23-3.33 (2H, m), 3.46 (2H, t, J = 6.7 Hz), 3.92 (3H, s), 4.30 (2H, t, J = 6.5 Hz), 7.56 (1H, d, J = 9.0 Hz), 8.92 (1H, s), 9.05 (1H, dd, J = 2.2, 8.9 Hz), 9.17 (1H, d, J = 2.2 Hz); ESI+: 542
26	11	NMR1: 0.98 (3H, t, J = 7.2 Hz), 1.12-1.29 (2H, m), 1.42-1.58 (1H, m), 1.64-1.88 (6H, m), 2.28 (2H, q, J = 7.2 Hz), 2.79-2.91 (2H, m), 3.92 (3H, s), 4.30 (2H, t, J = 6.4 Hz), 7.56 (1H, d, J = 9.0 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.2, 8.9 Hz), 9.16 (1H, d, J = 2.1 Hz); ESI+: 459
27	11	NMR1: 0.99 (3H, t, J = 7.2 Hz), 1.13-1.31 (3H, m), 1.41-1.55 (1H, m), 1.63-1.96 (5H, m), 2.25-2.39 (2H, m), 2.81-2.94 (2H, m), 3.93 (3H, s), 4.59 (2H, t, J = 6.5 Hz), 7.24 (1H, d, J = 3.6 Hz), 8.07 (1H, d, J = 3.6 Hz), 8.71 (1H, d, J = 2.0 Hz), 9.23 (1H, d, J = 2.0 Hz); ESI+: 459
28	28	NMR1: 1.33-1.49 (2H, m), 1.69-1.83 (3H, m), 1.84-1.93 (2H, m), 2.75-2.90 (2H, m), 3.20-3.32 (2H, m), 3.94 (3H, s), 4.61 (2H, t, J = 6.1 Hz), 8.53 (1H, br), 8.79 (1H, br), 8.97 (1H, s), 9.33 (1H, d, J = 2.2 Hz), 9.75 (1H, d, J = 2.2 Hz); ESI+: 432
29	6	NMR1: 0.95-1.21 (2H, m), 1.60-1.84 (8H, m), 1.98-2.10 (1H, m), 2.15-2.24 (4H, m), 2.45-2.58 (1H, m), 2.88-3.02 (2H, m), 3.04-3.18 (1H, m), 3.92 (3H, s), 4.09-4.45 (4H, m), 7.56 (1H, d, J = 9.0 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.2, 8.9 Hz), 9.16 (1H, d, J = 2.1 Hz); ESI+: 542
30	6	NMR2: 1.10-1.33 (2H, m), 1.66-2.15 (11H, m), 2.17-2.29 (1H, m), 2.31-2.39 (3H, m), 2.53-2.66 (1H, m), 2.88-3.09 (3H, m), 3.13-3.21 (1H, m), 3.61-3.67 (1H, m), 3.75-3.84 (2H, m), 3.95-4.02 (1H, m), 4.07-4.43 (5H, m), 4.62-4.73 (1H, m), 6.91 (1H, d, J = 3.6 Hz), 7.18 (1H, d, J = 8.7 Hz), 7.50 (1H, d, J = 3.6 Hz), 8.37 (1H, dd, J = 2.2, 8.7 Hz), 8.41 (1H, d, J = 2.2 Hz); ESI+: 611
31	3	NMR1: 1.20-1.32 (2H, m), 1.43-1.58 (1H, m), 1.68-1.80 (4H, m), 2.07-2.18 (2H, m), 2.74-2.83 (2H, m), 3.69 (2H, s), 3.92 (3H, s), 4.31 (2H, t, J = 6.4 Hz), 7.56 (1H, d, J = 9.0 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.2, 8.9 Hz), 9.17 (1H, d, J = 2.1 Hz); ESI+: 470
32	3	NMR1: 1.17-1.30 (2H, m), 1.42-1.55 (1H, m), 1.66-1.77 (4H, m), 2.02-2.12 (2H, m), 2.48 (3H, s), 2.79-2.86 (2H, m), 3.71 (2H, s), 3.92 (3H, s), 4.29 (2H, t, J = 6.4 Hz), 7.55 (1H, d, J = 9.0 Hz), 8.91 (1H, s), 9.04 (1H, dd, J = 2.2, 8.9 Hz), 9.16 (1H, d, J = 2.2 Hz); ESI+: 527
33	6	NMR1: 1.00-1.26 (2H, m), 1.70-1.83 (5H, m), 2.53-2.68 (1H, m), 2.85-2.98 (1H, m), 3.60-3.73 (1H, m), 3.92 (3H, s), 4.01-4.13 (2H, m), 4.26-4.39 (3H, m), 4.42 (1H, t, J = 5.4 Hz), 7.56 (1H, d, J = 9.0 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.2, 8.9 Hz), 9.16 (1H, d, J = 2.2 Hz); ESI+: 489
34	2	NMR1: 1.26-1.42 (2H, m), 1.57-1.81 (3H, m), 1.83-1.95 (2H, m), 2.76-2.94 (2H, m), 3.14-3.29 (4H, m), 3.94 (3H, s), 7.13 (1H, d, J = 4 Hz), 7.84 (1H, d, J = 9.0 Hz), 8.08 (1H, d, J = 4 Hz), 8.35-8.48 (3H, m), 8.62-8.8 (1H, br); ESI+: 446
35	2	NMR1: 1.66-1.77 (2H, m), 1.90-2.00 (2H, m), 2.91-3.03 (2H, m), 3.06-3.18 (2H, m), 3.65-3.73 (1H, m), 3.80-3.87 (2H, m), 3.93 (3H, s), 4.36-4.43 (2H, m), 7.12 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 8.8 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.40 (1H, d, J = 2.2 Hz), 8.49 (1H, dd, J = 2.2, 8.8 Hz), 8.57 (1H, br), 8.66 (1H, br); ESI+: 446

TABLE 55

Ex	Syn	DAT
36	2	NMR1: 3.02-3.26 (4H, m), 3.64-3.73 (4H, m), 3.93 (3H, s), 5.27 (2H, s), 7.44 (1H, d, J = 9.0 Hz), 8.93 (1H, s), 8.99 (1H, dd, J = 2.2, 9.0 Hz), 9.11 (2H, br), 9.18 (1H, d, J = 2.2 Hz); ESI+: 446
37	2	NMR1: 1.55-1.83 (4H, m), 1.89-2.25 (7H, m), 3.88-3.98 (5H, m), 4.24-4.36 (2H, m), 7.10-7.13 (1H, m), 7.47-7.52 (1H, m), 8.04-8.07 (1H, m), 8.37-8.40 (1H, m), 8.46-8.52 (1H, m), 8.55-8.97 (2H, m); ESI+: 456
38	2	NMR1: 1.63-2.04 (6H, m), 2.11-2.26 (3H, m), 2.81-2.97 (2H, m), 3.02-3.12 (1H, m), 3.13-3.24 (1H, m), 3.93 (3H, s), 4.25-4.41 (2H, m), 7.11 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 8.8 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.33-8.55 (3H, m), 8.92-9.03 (1H, m); ESI+: 456
39	39	NMR1: 0.97 (3H, t, J = 7.2 Hz), 1.13-1.26 (2H, m), 1.38-1.53 (1H, m), 1.64-1.85 (6H, m), 2.27 (2H, q, J = 7.2 Hz), 2.79-2.88 (2H, m), 3.94 (3H, s), 4.60 (2H, t, J = 6.6 Hz), 8.96 (1H, s), 9.33 (1H, d, J = 2.2 Hz), 9.76 (1H, d, J = 2.2 Hz); ESI+: 460
40	6	NMR1: 0.96-1.21 (2H, m), 1.58-1.82 (8H, m), 1.97-2.08 (1H, m), 2.11-2.23 (4H, m), 2.48-2.59 (1H, m), 2.88-3.00 (2H, m), 3.03-3.16 (1H, m), 3.94 (3H, s), 4.10-4.27 (1H, m), 4.31-4.47 (1H, m), 4.57-4.65 (2H, m), 8.96 (1H, s), 9.31 (1H, d, J = 2.1 Hz), 9.74 (1H, d, J = 2.1 Hz); ESI+: 543
41	6	NMR1: 0.96-1.23 (2H, m), 1.59-1.83 (8H, m), 1.97-2.08 (1H, m), 2.11-2.23 (4H, m), 2.45-2.59 (1H, m), 2.87-3.01 (2H, m), 3.02-3.15 (1H, m), 3.93 (3H, s), 4.11-4.26 (1H, m), 4.31-4.45 (1H, m), 4.60 (2H, t, J = 6.1 Hz), 7.23 (1H, d, J = 3.6 Hz), 8.07 (1H, d, J = 3.6 Hz), 8.71 (1H, d, J = 2.2 Hz), 9.23 (1H, d, J = 2.2 Hz); ESI+: 542

TABLE 55-continued

Ex	Syn	DAT
42	5	NMR1: 1.40 (3H, t, J = 7.0 Hz), 1.46-1.52 (2H, m), 1.72-1.81 (2H, m), 1.91-2.26 (5H, m), 2.69-2.83 (2H, m), 3.48-3.61 (1H, m), 4.32 (2H, q, J = 7.0 Hz), 5.03 (2H, s), 7.13 (1H, d, J = 3.7 Hz), 7.50 (1H, d, J = 8.8 Hz), 8.02 (1H, d, J = 3.7 Hz), 8.34 (1H, d, J = 7.5 Hz), 8.39 (1H, d, J = 2.2 Hz), 8.48 (1H, dd, J = 2.2, 8.8 Hz); ESI+: 487
43#	1	NMR1: 1.20-1.31 (2H, m), 1.44 (9H, s), 1.61-1.67 (2H, m), 1.72-1.82 (3H, m), 3.24-3.33 (2H, m), 3.82-3.88 (2H, m), 4.31 (2H, t, J = 6 Hz), 5.18 (2H, s), 7.16 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 8.8 Hz), 8.05 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.2 Hz), 8.48 (1H, dd, J = 2.2, 8.8 Hz); ESI+: 531
44	2	NMR1: 1.16 (3H, s), 1.58-1.69 (2H, m), 1.74-1.85 (2H, m), 3.04-3.23 (4H, m), 3.93 (3H, s), 4.09 (2H, s), 7.11 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 8.8 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.2 Hz), 8.49 (1H, dd, J = 2.2, 8.8 Hz), 8.55-8.80 (2H, m); ESI+: 430
45	11	NMR1: 1.01 (3H, t, J = 7.2 Hz), 2.29-2.45 (6H, m), 3.41-3.51 (4H, m), 3.92 (3H, s), 5.20 (2H, s), 7.39 (1H, d, J = 9.0 Hz), 8.92 (1H, s), 8.99 (1H, dd, J = 2.1, 9.0 Hz), 9.18 (1H, d, J = 2.1 Hz); ESI+: 474
46#	4	NMR1: 1.20-1.31 (2H, m), 1.61-1.68 (2H, m), 1.71-1.82 (3H, m), 3.12-3.49 (2H, m), 3.80-3.89 (2H, m), 4.31 (2H, t, J = 6.0 Hz), 5.16 (2H, s), 7.15 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 8.8 Hz), 8.05 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.4 Hz), 8.48 (1H, dd, J = 2.4, 8.8 Hz), 13.37 (1H, br); ESI+: 475

TABLE 56

Ex	Syn	DAT
47	5	NMR1: 1.20-1.31 (2H, m), 1.40-1.52 (2H, m), 1.60-1.67 (2H, m), 1.70-1.83 (5H, m), 1.89-1.98 (2H, m), 2.14 (3H, s), 2.66-2.73 (2H, m), 3.28 (2H, dt, J = 1.6, 11.6 Hz), 3.46-3.57 (1H, m), 3.82-3.88 (2H, m), 4.31 (2H, t, J = 6.0 Hz), 5.03 (2H, s), 7.12 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.8 Hz), 8.02 (1H, d, J = 3.6 Hz), 8.32 (1H, d, J = 7.6 Hz), 8.38 (1H, d, J = 2.2 Hz), 8.47 (1H, dd, J = 2.2, 8.8 Hz); ESI+: 571
48#	2	NMR1: 1.36-1.50 (2H, m), 1.81-2.01 (3H, m), 2.80-2.94 (2H, m), 3.22-3.31 (2H, m), 3.45 (2H, d, J = 6.2 Hz), 3.95 (3H, s), 4.76 (2H, s), 7.15 (1H, d, J = 3.6 Hz), 7.95 (1H, d, J = 8.1 Hz), 8.10 (1H, d, J = 3.6 Hz), 8.40-8.54 (3H, m), 8.80 (1H, br); ESI+: 430
49#	11	NMR1: 0.98 (3H, t, J = 7.2 Hz), 1.16-1.29 (2H, m), 1.54-1.74 (3H, m), 1.78-1.88 (2H, m), 2.29 (2H, q, J = 7.2 Hz), 2.80-2.89 (2H, m), 3.41 (2H, d, J = 6.3 Hz), 3.94 (3H, s), 4.73 (2H, s), 7.17 (1H, d, J = 3.6 Hz), 7.96 (1H, d, J = 8.1 Hz), 8.09 (1H, d, J = 3.6 Hz), 8.43 (1H, d, J = 1.5 Hz), 8.51 (1H, dd, J = 1.5, 8.1 Hz); ESI+: 458
50	2	NMR1: 1.18-1.31 (1H, m), 1.54-2.01 (6H, m), 2.58-2.70 (1H, m), 2.70-2.83 (1H, m), 3.14-3.41 (2H, m), 3.94 (3H, s), 4.54-4.69 (2H, m), 7.23 (1H, d, J = 3.6 Hz), 8.09 (1H, d, J = 3.6 Hz), 8.60 (1H, br), 8.73 (1H, d, J = 2.2 Hz), 8.88 (1H, br), 9.24 (1H, d, J = 2.2 Hz); ESI+: 431
51	2	NMR1: 1.34-1.49 (2H, m), 1.59-1.69 (1H, m), 1.71-1.97 (6H, m), 2.76-2.93 (3H, m), 3.21-3.30 (2H, m), 3.49-3.55 (1H, m), 3.60-3.69 (2H, m), 3.79-3.86 (1H, m), 4.34-4.40 (2H, m), 4.57-4.64 (2H, m), 7.26 (1H, d, J = 3.6 Hz), 8.20 (1H, d, J = 3.6 Hz), 8.57 (1H, br), 8.72 (1H, d, J = 2.2 Hz), 8.80 (1H, br), 9.24 (1H, d, J = 2.2 Hz); ESI+: 501
52	12	NMR1: 1.22 (3H, t, J = 7.3 Hz), 1.41-1.55 (2H, m), 1.58-1.83 (3H, m), 1.87-2.00 (4H, m), 2.77-2.93 (3H, m), 3.01-3.12 (2H, m), 3.42-3.55 (3H, m), 3.60-3.69 (2H, m), 3.78-3.86 (1H, m), 4.33-4.40 (2H, m), 4.62 (2H, t, J = 6.0 Hz), 7.26 (1H, d, J = 3.6 Hz), 8.20 (1H, d, J = 3.6 Hz), 8.73 (1H, d, J = 2.2 Hz), 9.25 (1H, d, J = 2.2 Hz), 9.43 (1H, br); ESI+: 529
53#	2	NMR1: 1.32-1.49 (2H, m), 1.70-1.94 (6H, m), 1.97-2.23 (3H, m), 2.29-2.41 (2H, m), 2.78-2.94 (2H, m), 3.20-3.34 (2H, m), 3.56-3.80 (1H, m), 3.91 (3H, s), 4.13 (2H, t, J = 6.1 Hz), 7.06 (1H, d, J = 3.5 Hz), 7.15 (1H, d, J = 8.7 Hz), 7.96-8.01 (2H, m), 8.05 (1H, dd, J = 2.3, 8.6 Hz), 8.39-8.54 (1H, br), 8.66-8.83 (1H, br); ESI+: 416
54	13	NMR1: 1.24 (3H, t, J = 7.3 Hz), 1.46-1.69 (3H, m), 1.72-1.82 (3H, m), 1.88-2.01 (3H, m), 2.77-2.93 (3H, m), 3.01-3.10 (2H, m), 3.42-3.55 (3H, m), 3.57-3.68 (2H, m), 3.78-3.86 (1H, m), 4.28-4.40 (4H, m), 7.14 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.8 Hz), 8.16 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.2 Hz), 8.49 (1H, dd, J = 2.2, 8.8 Hz), 9.79 (1H, br); ESI+: 528
55#	9	NMR1: 1.12-1.36 (3H, m), 1.45-1.60 (2H, m), 1.63-1.72 (2H, m), 1.74-1.86 (2H, m), 2.13 (3H, s), 2.69-2.77 (2H, m), 3.23-3.38 (2H, m), 3.89 (3H, s), 6.14-6.21 (1H, m), 7.00 (1H, d, J = 8.8 Hz), 7.09 (1H, d, J = 3.6 Hz), 7.96 (1H, d, J = 3.6 Hz), 8.27-8.35 (2H, m); ESI+: 443
56	2	NMR1: 1.3-1.45 (2H, m), 1.75-1.94 (5H, m), 2.78-2.92 (2H, m), 3.20-3.40 (2H, m), 3.90 (3H, s), 4.6-4.67 (2H, m), 7.88-8.6 (2H, br), 9.23 (1H, d, J = 2.4 Hz), 9.77 (1H, d, J = 2.4 Hz); ESI+: 500

TABLE 56-continued

Ex	Syn	DAT
57#	9	NMR1: 1.53-1.67 (12H, m), 1.99-2.04 (3H, m), 2.78 (2H, t, J = 6.0 Hz), 3.20-3.26 (2H, m), 3.89 (3H, s), 6.31 (1H, t, J = 4.8 Hz), 7.00 (1H, d, J = 8.8 Hz), 7.08 (1H, d, J = 3.6 Hz), 7.96 (1H, d, J = 3.6 Hz), 8.29-8.34 (3H, m); ESI+: 495

TABLE 57

Ex	Syn	DAT
58	2	NMR1: 1.34-1.51 (2H, m), 1.55-1.67 (1H, m), 1.73-1.94 (7H, m), 1.95-2.06 (1H, m), 2.76-2.91 (2H, m), 3.22-3.36 (2H, m), 3.60-3.68 (1H, m), 3.75-3.83 (1H, m), 4.20-4.49 (5H, m), 7.11 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 9.0 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.1 Hz), 8.44-8.64 (2H, m), 8.71-8.87 (1H, br); ESI+: 500
59#	9	NMR1: 1.59-1.71 (2H, m), 1.85-1.95 (2H, m), 2.03-2.12 (2H, m), 2.19 (3H, s), 2.66-2.77 (2H, m), 3.48-3.61 (1H, m), 3.89 (3H, s), 5.30 (1H, d, J = 7.7 Hz), 7.06 (1H, d, J = 3.6 Hz), 7.09-7.13 (1H, m), 7.96 (1H, d, J = 3.6 Hz), 8.27-8.32 (2H, m); ESI+: 415
60#	9	NMR1: 1.14-1.30 (2H, m), 1.56-1.70 (3H, m), 1.74-1.84 (2H, m), 2.13 (3H, s), 2.72-2.79 (2H, m), 3.16-3.25 (2H, m), 3.89 (3H, s), 6.23-6.28 (1H, m), 7.00-7.05 (1H, m), 7.07 (1H, d, J = 3.6 Hz), 7.95 (1H, d, J = 3.6 Hz), 8.25-8.31 (2H, m); ESI+: 429
61#	9	NMR1: 1.68-1.77 (4H, m), 1.77-1.86 (2H, m), 2.44-2.52 (4H, m), 2.56-2.63 (2H, m), 3.32-3.40 (2H, m), 3.89 (3H, s), 6.95-7.02 (1H, m), 7.07 (1H, d, J = 3.6 Hz), 7.15-7.22 (1H, m), 7.95 (1H, d, J = 3.6 Hz), 8.28-8.34 (2H, m); ESI+: 429
62#	2	NMR1: 1.30-1.48 (2H, m), 1.77-1.96 (5H, m), 2.78-2.90 (2H, m), 3.23-3.33 (3H, m), 3.36 (3H, s), 3.93 (3H, s), 4.32-4.42 (2H, m), 7.09 (1H, d, J = 3.6 Hz), 7.54 (1H, d, J = 8.9 Hz), 8.08 (1H, d, J = 3.6 Hz), 8.46-8.62 (2H, m), 8.67 (1H, d, J = 2.3 Hz); ESI+: 440
63#	9	NMR1: 1.12 (6H, s), 1.63-1.75 (4H, m), 2.47-2.61 (4H, m), 3.10-3.16 (2H, m), 3.90 (3H, s), 6.02-6.11 (1H, br), 6.92-7.00 (1H, m), 7.08 (1H, d, J = 3.5 Hz), 7.97 (1H, d, J = 3.5 Hz), 8.28-8.35 (2H, m); ESI+: 443
64#	9	NMR1: 1.55-1.73 (3H, m), 1.74-1.97 (3H, m), 2.05-2.15 (1H, m), 2.27 (3H, s), 2.92-3.00 (1H, m), 3.25-3.40 (3H, m), 3.89 (3H, s), 6.93-6.99 (1H, m), 7.07 (1H, d, J = 3.6 Hz), 7.27-7.37 (1H, m), 7.95 (1H, d, J = 3.6 Hz), 8.25-8.36 (2H, m); ESI+: 429
65	13	NMR1: 1.50-2.00 (8H, m), 2.81-3.00 (3H, m), 3.08-3.16 (2H, m), 3.19-3.26 (1H, m), 3.48-3.56 (3H, m), 3.61-3.69 (2H, m), 3.73-3.86 (3H, m), 4.26-4.44 (4H, m), 5.30 (1H, br), 7.14 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.8 Hz), 8.17 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.1 Hz), 8.49 (1H, dd, J = 2.1, 8.8 Hz), 9.70 (1H, br); ESI+: 544
66	13	NMR1: 1.52-2.00 (7H, m), 2.81-3.06 (10H, m), 3.20-3.28 (1H, m), 3.36-3.55 (3H, m), 3.61-3.69 (2H, m), 3.78-3.86 (1H, m), 4.21-4.41 (6H, m), 7.14 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.8 Hz), 8.17 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.1 Hz), 8.49 (1H, dd, J = 2.1, 8.8 Hz), 9.46 (1H, br); ESI+: 585
67#	9	NMR1: 1.13-1.35 (3H, m), 1.49-1.72 (5H, m), 1.75-1.97 (3H, m), 2.13 (3H, s), 2.66-2.77 (2H, m), 2.80-2.91 (1H, m), 3.27-3.39 (2H, m), 3.47-3.53 (1H, m), 3.59-3.68 (2H, m), 3.77-3.85 (1H, m), 4.25-4.37 (2H, m), 6.15-6.20 (1H, m), 7.00 (1H, d, J = 8.8 Hz), 7.11 (1H, d, J = 3.6 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.27-8.33 (2H, m); ESI+: 513
68	2	NMR1: 1.34-1.49 (2H, m), 1.73-1.93 (5H, m), 2.76-2.89 (2H, m), 3.22-3.30 (2H, m), 4.27-4.35 (2H, m), 7.08 (1H, dd, J = 1.6, 3.6 Hz), 7.51 (1H, d, J = 8.9 Hz), 8.03 (1H, dd, J = 2.4, 3.6 Hz), 8.39 (1H, d, J = 2.1 Hz), 8.49 (1H, dd, J = 2.2, 8.8 Hz), 8.57-8.71 (1H, m), 8.81-8.96 (1H, br), 13.03 (1H, br-s); ESI+: 416
69#	9	NMR1: 0.92-1.08 (1H, m), 1.35-1.51 (1H, m), 1.58-1.72 (2H, m), 1.72-1.83 (1H, m), 1.87-2.02 (2H, m), 2.13 (3H, s), 2.46-2.60 (1H, m), 2.61-2.72 (1H, m), 3.17-3.27 (2H, m), 3.89 (3H, s), 6.37-6.51 (1H, m), 7.00-7.05 (1H, m), 7.08 (1H, d, J = 3.6 Hz), 7.95 (1H, d, J = 3.6 Hz), 8.27-8.33 (2H, m); ESI+: 429

TABLE 58

Ex	Syn	DAT
70	11	NMR1: 0.98 (3H, t, J = 7.2 Hz), 1.13-1.29 (2H, m), 1.42-1.56 (1H, m), 1.61-1.85 (7H, m), 1.90-2.03 (1H, m), 2.28 (2H, q, J = 7.2 Hz), 2.80-2.96 (3H, m), 3.51-3.58 (1H, m), 3.59-3.72 (2H, m), 3.77-3.85 (1H, m), 4.24-4.39 (4H, m), 7.56 (1H, d, J = 9.1 Hz), 9.03 (1H, s), 9.04 (1H, dd, J = 2.0, 9.1 Hz), 9.18 (1H, d, J = 2.0 Hz); ESI+: 529
71	11	NMR1: 0.97 (3H, t, J = 7.2 Hz), 1.13-1.27 (2H, m), 1.37-1.53 (1H, m), 1.60-1.85 (7H, m), 1.91-2.02 (1H, m), 2.27 (2H, q, J = 7.1 Hz), 2.76-2.97 (3H, m),

TABLE 58-continued

Ex	Syn	DAT
72	3	3.51-3.58 (1H, m), 3.59-3.73 (2H, m), 3.77-3.86 (1H, m), 4.29-4.44 (2H, m), 4.54-4.64 (2H, m), 9.08 (1H, s), 9.32 (1H, d, J = 2.1 Hz), 9.74 (1H, d, J = 2.1 Hz); ESI+: 530 NMR1: 1.19-1.31 (2H, m), 1.46-1.56 (1H, m), 1.66-1.77 (4H, m), 1.96-2.09 (2H, m), 2.80 (3H, s), 2.81-2.88 (2H, m), 3.01 (3H, s), 3.08-3.18 (2H, m), 3.92 (3H, s), 4.27-4.31 (2H, m), 7.55 (1H, d, J = 8.8 Hz), 8.91 (1H, s), 9.04 (1H, dd, J = 2.0, 8.8 Hz), 9.15 (1H, d, J = 2.0 Hz); ESI+: 516
73	14 + 13	NMR1: 1.33 (9H, s), 1.48-1.62 (2H, m), 1.73-1.86 (3H, m), 1.94-2.04 (2H, m), 2.80-2.93 (2H, m), 3.48-3.56 (2H, m), 3.93 (3H, s), 4.29-4.36 (2H, m), 7.12 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 8.9 Hz), 8.04 (1H, d, J = 3.6 Hz), 8.40 (1H, d, J = 2.2 Hz), 8.50 (1H, dd, J = 2.2, 8.7 Hz), 8.90 (1H, br); ESI+: 486
74	8	NMR1: 1.20-1.32 (2H, m), 1.60-1.81 (5H, m), 2.72-2.90 (3H, m), 3.08-3.67 (13H, m), 3.82-3.88 (2H, m), 4.29-4.35 (2H, m), 5.38 (2H, s), 7.15 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 9.0 Hz), 7.96 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.0 Hz), 8.48 (1H, dd, J = 2.0, 8.9 Hz); ESI+: 571
75	8	NMR1: 1.00-1.40 (2H, m), 1.52-2.10 (6H, m), 2.65-2.76 (1H, m), 2.95-3.43 (15H, m), 3.54-3.66 (3H, m), 3.81-3.90 (2H, m), 4.00-4.11 (1H, m), 4.25-4.38 (3H, m), 5.30-5.50 (2H, m), 7.14 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 8.8 Hz), 7.96 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 1.6 Hz), 8.48 (1H, dd, J = 1.6, 8.8 Hz); ESI+: 625
76	10	NMR1: 1.35-1.45 (5H, m), 1.74-1.93 (5H, m), 2.78-2.90 (2H, m), 3.23-3.41 (2H, m), 4.31 (2H, t, J = 6 Hz), 4.39 (2H, q, J = 7.2 Hz), 7.12 (1H, d, J = 3.6 Hz), 7.50 (1H, d, J = 8.8 Hz), 8.14 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.4 Hz), 8.44-8.55 (2H, m), 8.69-8.80 (1H, br); ESI+: 444
77	6	NMR1: 0.81-0.89 (1H, m), 0.97-1.32 (4H, m), 1.59-1.83 (7H, m), 1.90-2.07 (2H, m), 2.12-2.23 (4H, m), 2.83-3.01 (3H, m), 3.02-3.16 (1H, m), 3.51-3.58 (1H, m), 3.59-3.73 (2H, m), 3.76-3.86 (1H, m), 4.11-4.46 (6H, m), 7.56 (1H, d, J = 9.0 Hz), 9.00-9.07 (2H, m), 9.15-9.19 (1H, m); ESI+: 612
78	2	NMR1: 1.32-1.47 (2H, m), 1.74-1.85 (3H, m), 1.85-1.97 (4H, m), 2.11-2.25 (2H, m), 2.78-2.92 (2H, m), 3.22-3.38 (2H, m), 3.56-3.67 (2H, m), 3.98-4.08 (2H, m), 4.27-4.35 (2H, m), 4.97-5.08 (1H, m), 7.16 (1H, d, J = 3.7 Hz), 7.50 (1H, d, J = 8.9 Hz), 8.24-8.42 (3H, m), 8.48 (1H, dd, J = 2.2, 8.8 Hz), 8.56-8.68 (1H, br); ESI+: 500
79	6	NMR1: 0.81-1.34 (5H, m), 1.58-1.86 (7H, m) 1.90-2.09 (2H, m), 2.11-2.25 (4H, m), 2.84-3.01 (3H, m), 3.03-3.18 (1H, m), 3.51-3.59 (1H, m), 3.59-3.73 (2H, m), 3.76-3.87 (1H, m), 4.10-4.26 (1H, m), 4.29-4.45 (3H, m), 4.55-4.67 (2H, m), 9.08 (1H, s), 9.32 (1H, d, J = 1.7 Hz), 9.74 (1H, d, J = 1.7 Hz); ESI+: 613
80	10	NMR1: 1.21-1.50 (6H, m), 1.70-1.93 (5H, m), 2.09-2.25 (1H, m), 2.76-2.91 (2H, m), 3.17-3.31 (4H, m), 3.78-3.88 (2H, m), 4.05-4.43 (2H, m), 4.56-4.65 (2H, m), 7.25 (1H, d, J = 3.6 Hz), 8.13 (1H, d, J = 3.6 Hz), 8.54 (1H, br), 8.71 (1H, d, J = 2.2 Hz), 8.76 (1H, br), 9.24 (1H, d, J = 2.2 Hz); ESI+: 515

TABLE 59

Ex	Syn	DAT
81	13	NMR1: 1.19-1.59 (9H, m), 1.66-2.00 (5H, m), 2.10-2.24 (1H, m), 2.76-2.90 (2H, m), 3.00-3.29 (4H, m), 3.40-3.51 (2H, m), 3.79-3.88 (2H, m), 4.28 (2H, d, J = 7.2 Hz), 4.61 (2H, t, J = 6.0 Hz), 7.25 (1H, d, J = 3.6 Hz), 8.13 (1H, d, J = 3.6 Hz), 8.72 (1H, d, J = 2.2 Hz), 9.24 (1H, d, J = 2.2 Hz), 9.67 (1H, br); ESI+: 543
82	10	NMR1: 3.25-3.41 (1H, m), 3.84-3.97 (5H, m), 3.99-4.12 (2H, m), 4.44 (2H, d, J = 6.2 Hz), 7.13 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.9 Hz), 8.07 (1H, d, J = 3.6 Hz), 8.41 (1H, d, J = 2.1 Hz), 8.52 (1H, dd, J = 2.1, 8.9 Hz), 8.77-9.11 (2H, m); ESI+: 388
83	10	NMR1: 1.33-1.47 (1H, m), 1.62-1.77 (1H, m), 1.82-1.95 (2H, m), 2.25-2.39 (1H, m), 2.72-2.86 (2H, m), 3.23-3.47 (2H, m), 3.93 (3H, s), 4.11-4.20 (1H, m), 4.22-4.30 (1H, m), 7.12 (1H, d, J = 3.6 Hz), 7.49 (1H, d, J = 8.8 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.40 (1H, d, J = 2.0 Hz), 8.50 (1H, dd, J = 2.0, 8.8 Hz), 8.57-8.86 (2H, m); ESI+: 416
84	10	NMR1: 1.57-1.70 (1H, m), 1.87-2.03 (2H, m), 2.07-2.19 (1H, m), 2.34-2.47 (1H, m), 2.77-2.88 (1H, m), 3.05-3.17 (1H, m), 3.19-3.52 (2H, m), 3.93 (3H, s), 4.26-4.36 (2H, m), 7.12 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.8 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.40 (1H, d, J = 2.0 Hz), 8.50 (1H, dd, J = 2.0, 8.8 Hz), 8.75-9.01 (2H, m); ESI+: 416
85	10	NMR1: 1.38-1.52 (2H, m), 1.58-1.69 (1H, m), 1.77-1.98 (6H, m), 2.80-2.94 (3H, m), 3.22-3.32 (2H, m), 3.51 (1H, dd, J = 5.3, 8.7 Hz), 3.60-3.68 (2H, m), 3.78-3.86 (1H, m), 4.28-4.41 (4H, m), 7.25 (1H, d, J = 3.6 Hz), 7.49 (1H, d, J = 8.8 Hz), 8.17 (1H, d, J = 3.6 Hz), 8.40-8.52 (2H, m), 8.52-8.90 (2H, m); ESI+: 457
86	10	NMR1: 1.45-1.69 (3H, m), 1.88-2.00 (3H, m), 2.10-2.23 (1H, m), 2.81-3.00 (3H, m), 3.31-3.38 (2H, m), 3.53 (1H, dd, J = 5.3, 8.7 Hz), 3.60-3.68 (2H, m), 3.78-3.86 (1H, m), 4.16 (2H, d, J = 6.0 Hz),

TABLE 59-continued

Ex	Syn	DAT
87	11	4.30-4.42 (2H, m), 7.13 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.8 Hz), 8.16 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.0 Hz), 8.46-8.51 (1H, m), 8.51-8.80 (2H, m); ESI+: 486 NMR1: 0.98 (3H, t, J = 7.2 Hz), 1.16-1.30 (2H, m), 1.44-1.97 (9H, m), 2.29 (2H, q, J = 7.2 Hz), 2.78-2.93 (3H, m), 3.51 (1H, dd, J = 5.3, 8.7 Hz), 3.60-3.69 (2H, m), 3.78-3.86 (1H, m), 4.28-4.39 (4H, m), 7.25 (1H, d, J = 3.6 Hz), 7.49 (1H, d, J = 9.3 Hz), 8.16 (1H, d, J = 3.6 Hz), 8.45-8.51 (2H, m); ESI+: 485
88	12	NMR1: 1.23 (3H, t, J = 7.3 Hz), 1.40-1.67 (3H, m), 1.68-2.06 (8H, m), 2.76-2.92 (2H, m), 3.02-3.12 (2H, m), 3.42-3.52 (2H, m), 3.60-3.68 (1H, m), 3.75-3.83 (1H, m), 4.20-4.49 (5H, m), 7.12 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.9 Hz), 8.06 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.1 Hz), 8.49 (1H, dd, J = 2.1, 8.7 Hz), 9.33-9.60 (1H, br); ESI+: 528
89	11	NMR1: 0.87 (3H, t, J = 7.2 Hz), 2.41 (2H, q, J = 7.2 Hz), 2.80-2.92 (1H, m), 2.95-3.05 (2H, m), 3.25-3.33 (2H, m), 3.93 (3H, s), 4.34 (2H, d, J = 6.1 Hz), 7.12 (1H, d, J = 3.6 Hz), 7.50 (1H, d, J = 8.9 Hz), 8.05 (1H, d, J = 3.6 Hz), 8.39 (1H, d, J = 2.1 Hz), 8.47 (1H, dd, J = 2.1, 8.9 Hz); ESI+: 416
90	11	NMR1: 1.00 (3H, t, J = 7.2 Hz), 1.09-1.29 (1H, m), 1.43-1.57 (1H, m), 1.62-1.72 (1H, m), 1.72-1.81 (1H, m), 1.81-2.00 (2H, m), 2.00-2.12 (1H, m), 2.23-2.41 (2H, m), 2.66-2.80 (1H, m), 2.86-2.99 (1H, m), 3.92 (3H, s), 4.05-4.21 (2H, m), 7.11 (1H, d, J = 3.6 Hz), 7.47 (1H, d, J = 8.8 Hz), 8.04 (1H, d, J = 3.6 Hz), 8.37 (1H, d, J = 2.0 Hz), 8.46 (1H, dd, J = 2.0, 8.8 Hz); ESI+: 444

TABLE 60

Ex	Syn	DAT
91	11	NMR1: 1.01 (3H, t, J = 7.2 Hz), 1.36-1.49 (1H, m), 1.77-2.01 (3H, m), 2.10-2.17 (1H, m), 2.21-2.45 (4H, m), 2.48-2.58 (1H, m), 2.67-2.76 (1H, m), 3.92 (3H, s), 4.21-4.29 (2H, m), 7.11 (1H, d, J = 3.6 Hz), 7.48 (1H, d, J = 8.8 Hz), 8.04 (1H, d, J = 3.6 Hz), 8.37 (1H, d, J = 2.0 Hz), 8.46 (1H, dd, J = 2.0, 8.8 Hz); ESI+: 444
92	11	NMR1: 0.96 (3H, t, J = 7.2 Hz), 1.37-1.49 (2H, m), 1.76-1.86 (2H, m), 1.94-2.06 (2H, m), 2.26 (2H, q, J = 7.2 Hz), 2.58-2.70 (2H, m), 3.34-3.45 (1H, m), 3.76-3.83 (2H, m), 3.92 (3H, s), 4.33-4.40 (2H, m), 7.11 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.8 Hz), 8.04 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.1 Hz), 8.46 (1H, dd, J = 2.1, 8.8 Hz); ESI+: 474
93	6	NMR1: 1.64-1.83 (3H, m), 1.91-2.07 (1H, m), 2.09-2.19 (1H, m), 2.21-2.25 (3H, m), 2.73-2.83 (1H, m), 2.91-3.00 (1H, m), 3.04-3.17 (1H, m), 3.73-3.85 (1H, m), 3.87-4.03 (4H, m), 4.09-4.18 (1H, m), 4.30-4.44 (3H, m), 7.12 (1H, d, J = 3.6 Hz), 7.51 (1H, d, J = 8.9 Hz), 8.05 (1H, d, J = 3.6 Hz), 8.29 (1H, d, J = 2.0 Hz), 8.49 (1H, dd, J = 2.0, 8.9 Hz); ESI+: 499
94	6	NMR1: 1.31-1.52 (2H, m), 1.58-2.26 (11H, m), 2.59-2.77 (1H, m), 2.89-3.19 (3H, m), 3.93 (3H, s), 4.00-4.31 (4H, m), 7.11 (1H, d, J = 3.5 Hz), 7.44-7.55 (1H, m), 8.05 (1H, d, J = 3.5 Hz), 8.37-8.43 (1H, m), 8.44-8.53 (1H, m); ESI+: 527
95	6	NMR1: 1.48-1.79 (4H, m), 1.86-2.42 (9H, m), 2.91-3.07 (2H, m), 3.11-3.85 (4H, m), 3.93 (3H, s), 4.27-4.36 (2H, m), 7.09-7.15 (1H, m), 7.47-7.55 (1H, m), 8.02-8.07 (1H, m), 8.36-8.41 (1H, m), 8.45-8.52 (1H, m); ESI+: 527
96	3	NMR1: 1.12-1.29 (1H, m), 1.42-1.57 (1H, m), 1.60-1.80 (2H, m), 1.98-2.19 (3H, m), 2.58-2.88 (5H, m), 3.01 (3H, s), 3.03-3.19 (2H, m), 3.92 (3H, s), 4.08-4.20 (2H, m), 7.11 (1H, d, J = 3.5 Hz), 7.47 (1H, d, J = 8.8 Hz), 8.04 (1H, d, J = 3.5 Hz), 8.37 (1H, d, J = 1.4 Hz), 8.46 (1H, dd, J = 1.4, 8.8 Hz); ESI+: 501
97	12	NMR1: 1.08-1.30 (6H, m), 1.64-1.97 (4H, m), 2.99-3.20 (4H, m), 3.31-3.48 (2H, m), 3.93 (3H, s), 3.98-4.01 (1H, m), 4.24-4.29 (1H, m), 7.11 (1H, d, J = 3.6 Hz), 7.47-7.56 (1H, m), 8.06 (1H, d, J = 3.6 Hz), 8.38-8.42 (1H, m), 8.46-8.56 (1H, m), 9.59-9.89 (1H, m); ESI+: 458
98#	9	NMR1: 2.14 (6H, s), 2.40 (2H, t, J = 5.8 Hz), 3.48 (2H, q, J = 6.0 Hz), 3.53 (2H, t, J = 5.8 Hz), 3.63 (2H, t, J = 6.0 Hz), 3.90 (3H, s), 5.93-5.99 (1H, m), 7.08 (1H, d, J = 3.6 Hz), 7.11 (1H, m), 7.97 (1H, d, J = 3.6 Hz), 8.28-8.34 (2H, m); ESI+: 433
99#	9	NMR1: 1.96-2.06 (2H, m), 2.25 (3H, s), 3.26-3.37 (2H, m), 3.90 (3H, s), 3.96 (2H, t, J = 7.1 Hz), 6.25-6.35 (1H, m), 6.74 (1H, d, J = 1.2 Hz), 6.96-7.01 (1H, m), 7.05-7.10 (2H, m), 7.96 (1H, d, J = 3.5 Hz), 8.26-8.32 (2H, m); ESI+: 440
100#	9	NMR1: 1.86-1.96 (2H, m), 2.17-2.25 (2H, m), 3.38-3.49 (6H, m), 3.90 (3H, s), 6.25-6.31 (1H, br), 7.05-7.11 (2H, m), 7.97 (1H, d, J = 3.6 Hz), 8.27-8.36 (2H, m); ESI+: 429
101	2	NMR1: 1.33-1.47 (2H, m), 1.74-1.94 (5H, m), 2.16-2.26 (1H, m), 2.53-2.64 (1H, m), 2.77-2.90 (2H, m), 3.22-3.32 (2H, m), 3.83-3.92 (1H, m), 3.92-4.04 (2H, m), 4.10-4.19 (1H, m), 4.26-4.36 (2H, m), 5.54-5.62 (1H, m), 7.16 (1H, d, J = 3.7 Hz), 7.51 (1H, d, J = 9.0 Hz), 8.06 (1H, d, J = 3.8 Hz), 8.38 (1H, d, J = 2.2 Hz), 8.40-8.52 (2H, m), 8.63-8.78 (1H, m); ESI+: 486

TABLE 61

Ex	Syn	DAT
102	PSyn24 + 11	NMR1: 0.98 (3H, t, J = 7.2 Hz), 1.14-1.27 (2H, m), 1.45-1.55 (1H, m), 1.65-1.85 (6H, m), 2.28 (2H, q, J = 7.2 Hz), 2.81-2.88 (2H, m), 4.29 (2H, t, J = 6.8 Hz), 5.03 (2H, s), 7.12 (1H, d, J = 3.6 Hz), 7.35 (1H, br- s), 7.51 (1H, d, J = 9.2 Hz), 7.76 (1H, br- s), 8.02 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.4 Hz), 8.47 (1H, dd, J = 2.4, 9.2 Hz); ESI+: 501
103	103	NMR1: 1.08 (3H, t, J = 7.2 Hz), 1.29-1.40 (2H, m), 1.44 (3H, t, J = 7.6 Hz), 1.58-1.68 (1H, m), 1.72-1.84 (4H, m), 2.22-2.36 (2H, m), 2.60-2.69 (2H, m), 3.07-3.16 (2H, m), 4.30 (2H, t, J = 6.4 Hz), 4.39 (2H, q, J = 7.2 Hz), 6.54 (2H, s), 7.13 (1H, d, J = 3.6 Hz), 7.50 (1H, d, J = 8.8 Hz), 8.13 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.4 Hz), 8.47 (1H, dd, J = 2.4, 8.8 Hz); ESI+: 472
104	6	NMR1: 1.00-1.25 (2H, m), 1.57-1.85 (9H, m), 1.86-1.97 (1H, m), 1.98-2.10 (1H, m), 2.11-2.24 (4H, m), 2.50-2.62 (1H, m), 2.80-3.18 (4H, m), 3.51 (1H, dd, J = 5.3, 8.7 Hz), 3.60-3.69 (2H, m), 3.78-3.86 (1H, m), 4.10-4.46 (6H, m), 7.25 (1H, d, J = 3.6 Hz), 7.50 (1H, d, J = 9.3 Hz), 8.16 (1H, d, J = 3.6 Hz), 8.45-8.51 (2H, m); ESI+: 568
105	6	NMR1: 1.11-1.40 (2H, m), 1.58-2.21 (10H, m), 2.21 (3H, s), 2.52-2.65 (1H, m), 2.80-3.21 (4H, m), 3.51 (1H, dd, J = 5.3, 8.7 Hz), 3.60-3.69 (2H, m), 3.78-3.86 (1H, m), 4.09-4.52 (6H, m), 7.14 (1H, d, J = 3.6 Hz), 7.48 (1H, d, J = 8.9 Hz), 8.15 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.2 Hz), 8.47 (1H, dd, J = 2.2, 8.9 Hz); ESI+: 597
106	PSyn26 + 103	NMR1: 1.08 (3H, t, J = 7.2 Hz), 1.35-1.55 (2H, m), 1.59-1.68 (1H, m), 1.76-1.99 (4H, m), 2.20-2.35 (2H, m), 2.55-2.64 (2H, m), 2.80-2.93 (1H, m), 3.08-3.16 (2H, m), 3.51 (1H, dd, J = 5.3, 8.7 Hz), 3.60-3.69 (2H, m), 3.78-3.86 (1H, m), 4.10-4.18 (2H, m), 4.30-4.41 (2H, m), 6.55 (2H, s), 7.13 (1H, d, J = 3.6 Hz), 7.49 (1H, d, J = 8.9 Hz), 8.15 (1H, d, J = 3.6 Hz), 8.38 (1H, d, J = 2.4 Hz), 8.47 (1H, dd, J = 2.4, 8.9 Hz); ESI+: 514
107	10	NMR1: 1.32-1.47 (2H, m), 1.71-1.94 (5H, m), 2.77-2.91 (2H, m), 3.22-3.76 (10H, m), 4.57-4.66 (2H, m), 5.41 (2H, s), 7.25 (1H, d, J = 3.6 Hz), 7.99 (1H, d, J = 3.6 Hz), 8.38 (1H, br), 8.63 (1H, br), 8.74 (1H, d, J = 2.2 Hz), 9.26 (1H, d, J = 2.2 Hz); ESI+: 544
108	16	NMR1: 1.46-1.67 (2H, m), 1.70-1.86 (4H, m), 1.86-2.05 (4H, m), 2.86-3.00 (2H, m), 3.08-3.17 (2H, m), 3.19-3.35 (1H, m), 3.46-3.56 (2H, m), 3.60-3.69 (1H, m), 3.71-3.83 (3H, m), 4.20-4.49 (5H, m), 5.20-5.40 (1H, br), 7.12 (1H, d, J = 3.6 Hz), 7.49-7.55 (1H, m), 8.06 (1H, d, J = 3.7 Hz), 8.39 (1H, d, J = 2.1 Hz), 8.49 (1H, dd, J = 2.2, 8.7 Hz), 9.31-9.54 (1H, br); ESI+: 544
109	13	NMR1: 1.23 (3H, t, J = 7.3 Hz), 1.43-1.59 (2H, m), 1.66-2.01 (5H, m), 2.77-2.91 (2H, m), 3.00-3.24 (2H, m), 3.41-3.52 (4H, m), 3.57-3.77 (6H, m), 4.62 (2H, t, J = 6.0 Hz), 5.42 (2H, s), 7.25 (1H, d, J = 3.7 Hz), 8.00 (1H, d, J = 3.7 Hz), 8.74 (1H, d, J = 2.2 Hz), 9.26 (1H, d, J = 2.2 Hz), 9.65 (1H, br); ESI+: 572
110	13	NMR1: 1.24 (3H, t, J = 7.3 Hz), 1.45-1.61 (2H, m), 1.66-2.01 (5H, m), 2.76-2.90 (2H, m), 3.01-3.18 (2H, m), 3.42-3.52 (2H, m), 3.74-3.85 (2H, m), 4.28-4.36 (2H, m), 4.41 (2H, t, J = 5.5 Hz), 5.00 (1H, br), 7.11 (1H, d, J = 3.7 Hz), 7.52 (1H, d, J = 8.8 Hz), 8.08 (1H, d, J = 3.7 Hz), 8.39 (1H, d, J = 2.2 Hz), 8.49 (1H, dd, J = 2.2, 8.8 Hz), 9.83 (1H, br); ESI+: 488
111	11	NMR1: 0.95 (6H, d, J = 6.6 Hz), 1.14-1.29 (2H, m), 1.42-1.56 (1H, m), 1.58-1.81 (5H, m), 1.86-1.97 (1H, m), 2.01-2.17 (2H, m), 2.59-2.93 (4H, m), 3.51 (1H, dd, J = 5.3, 8.7 Hz), 3.60-3.69 (2H, m), 3.78-3.86 (1H, m), 4.27-4.40 (4H, m), 7.25 (1H, d, J = 3.7 Hz), 7.49 (1H, d, J = 9.3 Hz), 8.16 (1H, d, J = 3.7 Hz), 8.44-8.52 (2H, m); ESI+: 499

TABLE 62

Ex	Syn	DAT
112	11	NMR1: 1.14-1.29 (2H, m), 1.45-1.80 (6H, m), 1.86-1.97 (1H, m), 2.37-2.46 (2H, m), 2.80-2.92 (3H, m), 2.93-3.11 (1H, m), 3.52 (1H, dd, J = 5.3, 8.7 Hz), 3.58-3.69 (2H, m), 3.78-3.86 (1H, m), 4.27-4.40 (4H, m), 4.50-4.55 (2H, m), 4.61-4.67 (2H, m), 7.25 (1H, d, J = 3.6 Hz), 7.49 (1H, d, J = 9.3 Hz), 8.16 (1H, d, J = 3.6 Hz), 8.44-8.50 (2H, m); ESI+: 535
113	10	NMR1: 1.32-1.49 (2H, m), 1.68-1.94 (5H, m), 2.75-2.92 (2H, m), 3.21-3.33 (2H, m), 3.94 (3H, s), 4.30-4.43 (2H, m), 8.40-8.62 (1H, br), 8.66-8.82 (1H, br), 8.86-8.94 (1H, m), 8.95-9.02 (2H, m); ESI+: 449
114	114	NMR1: 1.08 (3H, t, J = 7.2 Hz), 1.27-1.41 (2H, m), 1.56-1.68 (1H, m), 1.71-1.84 (4H, m), 2.18-2.34 (2H, m), 2.56-2.68 (2H, m), 2.80-3.90 (2H, m), 4.26-4.36 (2H, m), 4.99-5.10 (4H, m), 5.98-6.08 (1H, m), 6.54 (2H, s), 7.26 (1H, d, J = 3.8 Hz), 7.51 (1H, d, J = 9.0 Hz), 8.38 (1H, d, J = 2.2 Hz), 8.45-8.53 (2H, m); ESI+: 500
115	115	NMR1: 0.99-1.29 (2H, m), 1.63-1.88 (8H, m), 2.10-2.24 (1H, m), 2.30-2.52 (4H, m), 2.52-2.64 (1H, m), 2.77-3.86 (3H, m), 3.96-4.13 (1H, m), 4.26-4.44 (3H, m), 4.99-5.10 (4H, m),

TABLE 62-continued

Ex	Syn	DAT
116	10	5.99-6.08 (1H, m), 6.56 (2H, s), 7.26 (1H, d, J = 3.8 Hz), 7.52 (1H, d, J = 8.9 Hz), 8.38 (1H, d, J = 2.2 Hz), 8.45-8.53 (2H, m); ESI+: 583 NMR1: 1.34-1.50 (2H, m), 1.72-1.96 (5H, m), 2.78-2.91 (2H, m), 3.21-3.32 (2H, m), 3.94 (3H, s), 4.56-4.65 (2H, m), 7.23 (1H, d, J = 3.6 Hz), 8.08 (1H, d, J = 3.6 Hz), 8.48 (1H, br), 8.66-8.79 (2H, m), 9.22-9.25 (1H, m); ESI+: 431
117	6	NMR1: 0.96-1.26 (2H, m), 1.49-1.66 (1H, m), 1.66-1.84 (5H, m), 2.18 (3H, s), 2.21-2.40 (2H, m), 2.44-2.62 (1H, m), 2.78-3.19 (3H, m), 3.92 (3H, s), 4.10-4.22 (1H, m), 4.25-4.49 (4H, m), 4.75-4.87 (1H, m), 7.56 (1H, d, J = 8.9 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.0, 8.9 Hz), 9.17 (1H, d, J = 2.0 Hz); ESI+: 558
118	6	NMR1: 0.95-1.25 (2H, m), 1.67-1.83 (5H, m), 1.84-1.99 (2H, m), 2.18-2.23 (3H, m), 2.23-2.35 (1H, m), 2.36-2.45 (1H, m), 2.46-2.59 (2H, m), 2.64-2.78 (1H, m), 2.90-3.01 (1H, m), 3.14-3.25 (1H, m), 3.85-3.96 (4H, m), 4.24-4.42 (3H, m), 7.56 (1H, d, J = 8.8 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.4, 8.8 Hz), 9.16 (1H, d, J = 2.4 Hz); ESI+: 542
119	6	NMR1: 0.94-1.21 (2H, m), 1.46-1.65 (4H, m), 1.67-1.82 (5H, m), 1.82-1.95 (2H, m), 2.13 (3H, s), 2.42-2.58 (2H, m), 2.75 (2H, d, J = 11.6 Hz), 2.98 (1H, t, J = 11.6 Hz), 3.86-3.98 (4H, m), 4.27-4.35 (2H, m), 4.35-4.46 (1H, m), 7.56 (1H, d, J = 9.0 Hz), 8.92 (1H, s), 9.06 (1H, dd, J = 2.1, 9.0 Hz), 9.17 (1H, d, J = 2.1 Hz); ESI+: 556
120	3	NMR1: 1.13-1.29 (2H, m), 1.43-1.62 (2H, m), 1.63-1.77 (5H, m), 1.78-2.00 (4H, m), 2.73-2.84 (2H, m), 3.00-3.22 (4H, m), 3.32-3.42 (1H, m), 3.68-3.81 (2H, m), 3.92 (3H, s), 4.24-4.35 (2H, m), 7.55 (1H, d, J = 9.0 Hz), 8.91 (1H, s), 9.04 (1H, dd, J = 2.0, 9.0 Hz), 9.16 (1H, d, J = 2.0 Hz); ESI+: 581
121	3	NMR1: 1.16-1.30 (2H, m), 1.42-1.56 (1H, m), 1.64-1.79 (4H, m), 1.94-2.07 (2H, m), 2.28-2.53 (2H, m), 2.75-2.87 (2H, m), 3.03-3.08 (1H, m), 3.09-3.13 (1H, m), 3.51 (1H, t, J = 7.6 Hz), 3.69 (1H, t, J = 13.2 Hz), 3.79 (1H, t, J = 7.6 Hz), 3.92 (3H, s), 4.02 (1H, t, J = 13.2 Hz), 4.30 (2H, t, J = 6.4 Hz), 7.55 (1H, d, J = 9.0 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.1, 9.0 Hz), 9.16 (1H, d, J = 2.1 Hz); ESI+: 578

TABLE 63

Ex	Syn	DAT
122	3	NMR1: 1.14-1.30 (2H, m), 1.42-1.58 (1H, m), 1.62-1.79 (4H, m), 1.86-2.26 (4H, m), 2.77-2.91 (2H, m), 3.03-3.07 (1H, m), 3.07-3.12 (1H, m), 3.24-3.91 (4H, m), 3.92 (3H, s), 4.23-4.34 (2H, m), 5.17-5.49 (1H, m), 7.56 (1H, d, J = 8.8 Hz), 8.91 (1H, s), 9.05 (1H, dd, J = 2.0, 8.8 Hz), 9.17 (1H, d, J = 2.0 Hz); ESI+: 560
123	PSyn3	ESI+: 502
124	PSyn26	ESI+: 528
125	3	ESI+: 585
126	3	ESI+: 544
127	PSyn24	ESI+: 501
128	PSyn24	ESI+: 502
129	PSyn26	ESI+: 543
130	PSyn26	ESI+: 472
131	PSyn26	ESI+: 572
132	PSyn24	ESI+: 460
133	PSyn26	ESI+: 488
134	134	NMR1: 1.23 (3H, t, J = 7.3 Hz), 1.45-1.60 (2H, m), 1.65-1.84 (3H, m), 1.85-2.01 (2H, m), 2.75-2.90 (2H, m), 2.99-3.24 (2H, m), 3.40-3.50 (2H, m), 3.94 (3H, s), 4.62 (2H, t, J = 6.1 Hz), 8.97 (1H, s), 9.33 (1H, d, J = 2.0 Hz), 9.70-9.96 (2H, m); ESI+: 460
135	135	NMR2: 1.15 (6H, s), 1.24-1.41 (2H, m), 1.61-1.70 (2H, m), 1.80-2.03 (4H, m), 2.09-2.24 (1H, m), 2.31 (2H, s), 2.32-2.42 (2H, m), 2.44-2.54 (1H, m), 2.56-2.67 (1H, m), 2.92-3.03 (2H, m), 3.04-3.34 (1H, m), 3.93-4.06 (4H, m), 4.29-4.50 (2H, m), 4.66-4.78 (1H, m), 6.90 (1H, d, J = 3.6 Hz), 7.52 (1H, d, J = 3.6 Hz), 8.72 (1H, d, J = 2.1 Hz), 9.10 (1H, d, J = 2.1 Hz); ESI+: 600
136	136	NMR1: 1.19-1.30 (2H, m), 1.40-1.52 (1H, m), 1.67-1.77 (4H, m), 1.84-1.99 (2H, m), 2.20 (3H, s), 2.77-2.85 (2H, m), 3.94 (3H, s), 4.60 (2H, t, J = 6.6 Hz), 8.96 (1H, s), 9.32 (1H, d, J = 2.1 Hz), 9.75 (1H, d, J = 2.1 Hz); ESI+: 446
137	6	NMR1: 0.96-1.20 (2H, m), 1.58-1.82 (8H, m), 1.96-2.08 (1H, m), 2.10-2.23 (4H, m), 2.43-2.59 (1H, m), 2.87-3.01 (2H, m), 3.02-3.15 (1H, m), 3.94 (3H, s), 4.10-4.26 (1H, m), 4.30-4.45 (1H, m), 4.62 (2H, t, J = 6.1 Hz), 8.96 (1H, s), 9.33 (1H, d, J = 2.1 Hz), 9.76 (1H, d, J = 2.1 Hz); ESI+: 543
138	10	ESI+: 529
139	11	ESI+: 474

TABLE 63-continued

Ex	Syn	DAT
140	11	NMR1: 0.81-1.30 (8H, m), 1.53-1.84 (9H, m), 1.95-2.06 (1H, m), 2.41-2.54 (1H, m), 2.70-2.84 (1H, m), 2.87-3.00 (2H, m), 3.55-3.68 (1H, m), 3.94 (3H, s), 4.20-4.44 (2H, m), 4.62 (2H, t, J = 6.1 Hz), 8.96 (1H, s), 9.32 (1H, d, J = 2.1 Hz), 9.75 (1H, d, J = 2.1 Hz); ESI+: 571
141	11	NMR1: 1.10-1.25 (2H, m), 1.37-1.50 (1H, m), 1.53-1.80 (10H, m), 1.88-1.98 (2H, m), 2.56-2.66 (1H, m), 2.70-2.80 (2H, m), 3.94 (3H, s), 4.59 (2H, t, J = 6.5 Hz), 8.96 (1H, s), 9.31 (1H, d, J = 2.1 Hz), 9.74 (1H, d, J = 2.1 Hz); ESI+: 486
142	1	ESI+: 472

TABLE 64

Ex	Syn	DAT
143	143	ESI+: 544
144	3	ESI+: 488
145	11	ESI+: 510
146	1	ESI+: 488
147	10	ESI+: 447
148	10	ESI+: 433
149	6	ESI+: 558
150	6	ESI+: 544
151	3	ESI+: 516
152	13	NMR1: 1.53-2.00 (7H, m), 2.88-3.04 (8H, m), 3.19-3.54 (2H, m), 3.94 (3H, s), 4.20-4.31 (2H, m), 4.57-4.65 (2H, m), 7.23 (1H, d, J = 3.6 Hz), 8.09 (1H, d, J = 3.6 Hz), 8.72 (1H, d, J = 2.0 Hz), 9.24 (1H, d, J = 2.0 Hz), 9.44-9.55 (1H, br); ESI+: 516
153	10	ESI+: 417
154	10	ESI+: 514
155	135	ESI+: 489
156	11	ESI+: 556
157	6	ESI+: 544
158	6	NMR2: 1.71-2.29 (7H, m), 2.31-2.36 (3H, m), 2.91-3.02 (1H, m), 3.11-3.19 (1H, m), 3.51-3.78 (4H, m), 3.80-3.94 (2H, m), 3.98 (3H, s), 4.17-4.30 (1H, m), 4.62-4.73 (2H, m), 6.89-6.93 (1H, m), 7.50-7.54 (1H, m), 8.73 (1H, s), 9.09-9.14 (1H, m); ESI+: 544
159	10	ESI+: 528
160	11	ESI+: 612

INDUSTRIAL APPLICABILITY

The compound of the formula (I) or a salt thereof has a cathepsin S inhibitory effect, and can be used as an agent for preventing and/or treating autoimmune disease, allergic disease, graft rejection of an organ, bone marrow or tissue, systemic lupus erythematosus, lupus nephritis or the like.

The invention claimed is:

1. A method for treating systemic lupus erythematosus or lupus nephritis, comprising administering to a patient in need thereof an effective amount of a compound or a salt thereof, which is selected from the group consisting of:

6-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-9-methyl-9H-purine-2-carbonitrile or a salt thereof; and

- 40 4-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile or a salt thereof.
2. The method according to claim 1, comprising administering
- 45 6-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-9-methyl-9H-purine-2-carbonitrile or a salt thereof.
3. The method according to claim 1, comprising administering
- 50 4-{6-[2-(1-ethylpiperidin-4-yl)ethoxy]-5-(trifluoromethyl)pyridin-3-yl}-7-methyl-7H-pyrrolo[2,3-d]pyrimidine-2-carbonitrile or a salt thereof.

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