

DESCRIPTION

Title of the Invention: SULFONAMIDE DERIVATIVE OR
PHARMACEUTICALLY ACCEPTABLE ACID ADDITION SALT THEREOF

Technical Field

5 [0001]

The present invention aims to provide a novel compound useful as a superior orexin agonist.

Background Art

[0002]

10 Narcolepsy is a sleeping disorder caused by the inability of the brain to control the sleep-wake cycle. The major symptoms of narcolepsy includes, for example, excessive daytime sleepiness during the day, cataplexy induced by emotion (particularly strong joy and surprise), hypnagogic
15 hallucination, and hypnagogic paralysis, and narcolepsy patients are under serious influence in general social life. The prevalence of narcolepsy is assumed to be 0.05 - 0.2% (0.16 - 0.18% in Japan), and the prevalence indicates that the disease is not rare.

20 [0003]

The therapy of narcolepsy mainly includes a drug therapy and life guidance. For drug therapy, methylphenidate, modafinil and pemoline are used to suppress daytime sleepiness, and tricyclic antidepressant, selective serotonin reuptake
25 inhibitor (SSRI), and serotonin and noradrenaline reuptake inhibitor (SNRI) are used to control cataplexy. While these treatment methods are symptomatic therapy of narcolepsy, they are not basic treatment methods.

[0004]

30 In recent years, the relationship between narcolepsy and orexin system dysfunction is attracting attention. Orexins are neuropeptides present in the lateral hypothalamic area, which are two kinds of peptide of orexin-A and orexin-B (hypocretin 1, hypocretin 2 (non-patent document 1)). They bind to orexin
35 1 receptor (hereinafter to be also referred to as OX1R) and orexin 2 receptor (hereinafter to be also referred to as OX2R), which are G-protein coupled receptors (non-patent document 2). It was suggested from model experiments using mouse and dog that lack of orexin receptor (both OX1R and OX2R are
40 expressed), or lack of OX2R causes narcolepsy (non-patent document 3). Furthermore, it was suggested from model experiments using mouse that the function of OX2R is important for maintaining wakefulness (non-patent document 4, non-patent

document 5).

[0005]

On the other hand, many narcolepsy patients were confirmed to show disappearance of orexin neuron, and
5 decreased orexin concentration (non-patent document 6).
Therefore, it is strongly suggested that narcolepsy is highly possibly caused by the lack of orexin.

[0006]

The orexin receptor is widely expressed in the brain.
10 Orexins are peptides, and are not useful for pharmaceutical use since permeability through the blood-brain barrier is extremely low. Therefore, a low molecule orexin receptor agonist has been desired. In recent years, a compound with a cyclic guanidine skeleton is reported as a small- molecule
15 OX2R agonist (patent document 1).

[0007]

In addition, orexin system is considered to not only control the above-mentioned sleep-wake but also appropriately control feeding behavior with emotion and energy balance. A
20 mouse under fasting increases the amount of behavior for searching food by increasing the waking time and decreasing the sleep hours. On the other hand, it was clarified that the waking time and the amount of behavior do not increase in orexin receptor-deficient mouse (non-patent document 7).
25 Moreover, it was suggested that an increase of the leptin sensitivity by OX2R regulates the homeostasis of body weight (non-patent document 8). From these findings, an orexin receptor (particularly OX2R) agonist is a potential therapeutic drug for not only narcolepsy but also diabetes,
30 obesity and metabolic syndrome.

[Document List]

[patent document]

[0008]

patent document 1: US-B-8,258,163

35 [non-patent documents]

[0009]

non-patent document 1: Proc. Natl. Acad. Sci. USA, 95, 322-327 (1998)

non-patent document 2: Cell, 92, 573-585 (1998)

40 non-patent document 3: Cell, 98, 365-376 (1999)

non-patent document 4: Cell, 98, 437-451 (1999)

non-patent document 5: Neuron, 38, 715-730 (2003)

non-patent document 6: Arch. Neurol., 59, 1553-1562 (2002)

non-patent document 7: Neuron, 38, 701-713 (2003)
non-patent document 8: Cell Metab., 9, 64-76 (2009)

SUMMARY OF THE INVENTION

Problems to be Solved by the Invention

5 [0010]

The present invention aims to provide a novel low-molecular-weight compound showing an orexin agonist activity, which is expected to be useful as a superior therapeutic or prophylactic agent for narcolepsy.

10 Means of Solving the Problems

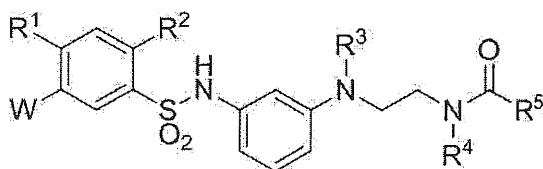
[0011]

The present inventors have conducted intensive studies in an attempt to solve the aforementioned problems and found a compound represented by the formula (I) mentioned below and
15 having a superior OX2R agonist activity, which resulted in the completion of the present invention.

That is, the present invention provides

[1] A compound represented by the formula (I)

[0012]



(I)

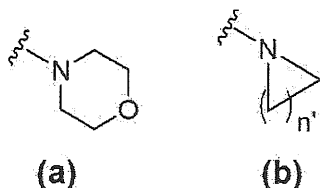
[0013]

wherein

R¹ is a hydrogen atom, alkyl having 1 to 4 carbon atoms or halogen,

25 R² is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, -OH, -NR^{2a}R^{2b} (wherein R^{2a} is a hydrogen atom, alkyl having 1 to 6 carbon atoms or cycloalkyl having 3 to 8 carbon atoms, R^{2b} is a hydrogen atom, alkyl having 1 to 6 carbon atoms or cycloalkyl having 3 to 8
30 carbon atoms), or the formula (a) or (b)

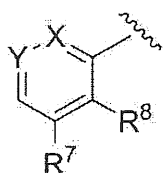
[0014]



[0015]

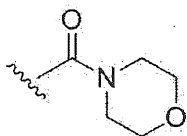
(wherein n' is an integer of 1 to 4)

- R^3 is a hydrogen atom or alkyl having 1 to 4 carbon atoms,
 R^4 is a hydrogen atom or alkyl having 1 to 4 carbon atoms,
 R^5 is aryl or heteroaryl (wherein aryl or heteroaryl is
optionally substituted by optionally selected 1 to 4 R^6),
5 R^6 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy
having 1 to 4 carbon atoms, halogen, -CN, -CF₃, -CH₂F, -CHF₂, -
OCF₃, -OH, -NO₂ or -NR^{6a}R^{6b} (wherein R^{6a} is a hydrogen atom,
alkyl having 1 to 4 carbon atoms, -CH₂CH₂OMe or -CH₂CH₂OEt, R^{6b}
is a hydrogen atom, alkyl having 1 to 4 carbon atoms, -
10 CH₂CH₂OMe or -CH₂CH₂OEt),
W is the formula (II):
[0016]

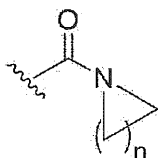


(II)

- [0017]
15 (wherein
 R^7 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy
having 1 to 4 carbon atoms or halogen,
 R^8 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy
having 1 to 4 carbon atoms or halogen,
20 X is -N= or -CR⁹=,
Y is -N= or -CR⁹=,
 R^9 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy
having 1 to 4 carbon atoms, halogen, -OH, -NH₂, -NR¹⁰R¹¹, -
CH₂OR¹⁰, -CF₃, -OCF₃, -CN, -C(O)OR¹⁰, -C(O)NR¹⁰R¹¹, the formula
25 (III) or (IV),
[0018]



(III)

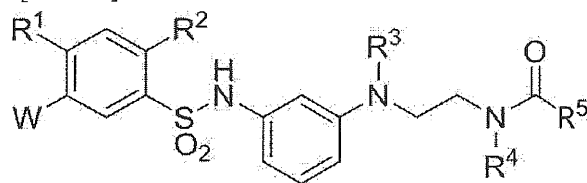


(IV)

- [0019]
(wherein n is an integer of 1 to 4)
30 X and Y are -CR⁹=, each R⁹ may be the same or different,
R¹⁰ is a hydrogen atom, alkyl having 1 to 4 carbon atoms, -
CH₂CF₃, -CH₂CH₂OH or -CH₂CH₂OMe,
R¹¹ is a hydrogen atom, alkyl having 1 to 4 carbon atoms, -
CH₂CF₃, -CH₂CH₂OH or -CH₂CH₂OMe,

or a pharmaceutically acceptable acid addition salt thereof,
 [1'] a compound represented by the formula (I)

[0020]



(I)

5 [0021]

wherein

R¹ is a hydrogen atom, alkyl having 1 to 4 carbon atoms or halogen,

R² is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy
 10 having 1 to 4 carbon atoms or halogen,

R³ is a hydrogen atom or alkyl having 1 to 4 carbon atoms,

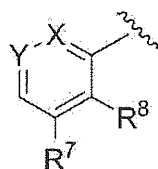
R⁴ is a hydrogen atom or alkyl having 1 to 4 carbon atoms,

R⁵ is aryl or heteroaryl (wherein aryl or heteroaryl is optionally substituted by optionally selected 1 to 4 R⁶),

15 R⁶ is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, -CN, -CF₃, -CH₂F, -CHF₂, -OCF₃, -OH, -NO₂ or -NR^aR^b (wherein R^a is a hydrogen atom, alkyl having 1 to 4 carbon atoms, -CH₂CH₂OMe or -CH₂CH₂OEt, R^b is a hydrogen atom, alkyl having 1 to 4 carbon atoms, -CH₂CH₂OMe or
 20 -CH₂CH₂OEt,

W is the formula (II):

[0022]



(II)

[0023]

25 (wherein

R⁷ is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms or halogen,

R⁸ is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms or halogen,

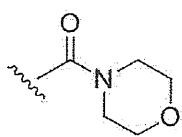
30 X is -N= or -CR⁹=,

Y is -N= or -CR⁹=,

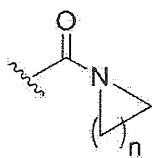
R⁹ is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, -OH, -NH₂, -NR¹⁰R¹¹, -

$\text{CH}_2\text{OR}^{10}$, $-\text{CF}_3$, $-\text{OCF}_3$, $-\text{CN}$, $-\text{C}(\text{O})\text{OR}^{10}$, $-\text{C}(\text{O})\text{NR}^{10}\text{R}^{11}$, the formula (III) or (IV),

[0024]



(III)



(IV)

5 [0025]

(wherein n is an integer of 1 to 4)

when X and Y are $-\text{CR}^9=$, each R^9 may be the same or different, R^{10} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-\text{CH}_2\text{CF}_3$, $-\text{CH}_2\text{CH}_2\text{OH}$ or $-\text{CH}_2\text{CH}_2\text{OMe}$, and

10 R^{11} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-\text{CH}_2\text{CF}_3$, $-\text{CH}_2\text{CH}_2\text{OH}$ or $-\text{CH}_2\text{CH}_2\text{OMe}$,

or a pharmaceutically acceptable acid addition salt thereof,

[2] the compound of the above-mentioned [1] or [1'], wherein

R^1 is a hydrogen atom, methyl, ethyl, a fluorine atom or a

15 chlorine atom,

R^2 is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a

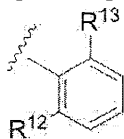
fluorine atom, a chlorine atom, $-\text{OH}$, $-\text{NH}_2$, $-\text{NH}(\text{Me})$ or $-\text{N}(\text{Me})_2$,

R^3 is a hydrogen atom or methyl,

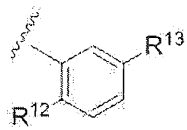
R^4 is a hydrogen atom or methyl,

20 R^5 is any group selected from the formulas (V) to (XV)

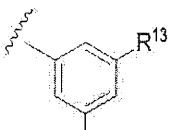
[0026]



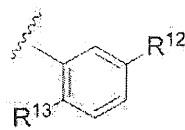
(V)



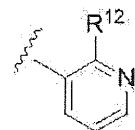
(VI)



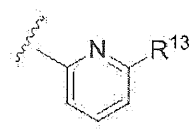
(VII)



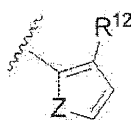
(VIII)



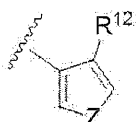
(IX)



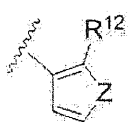
(X)



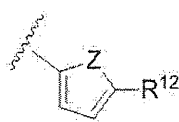
(XI)



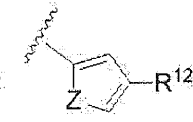
(XII)



(XIII)



(XIV)



(XV)

[0027]

(wherein

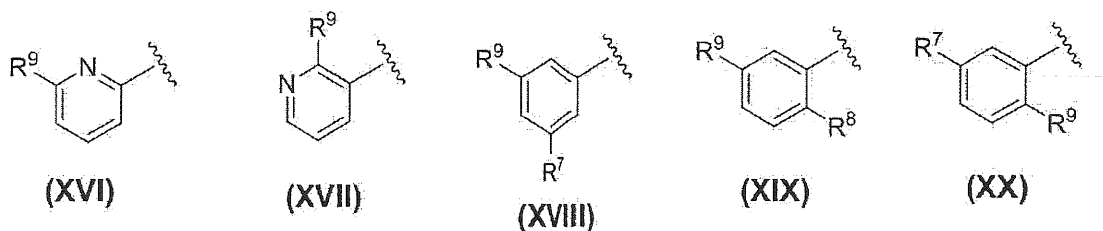
25 R^{12} is methyl, ethyl, n -propyl, isopropyl, $tert$ -butyl, methoxy, ethoxy, a fluorine atom, a chlorine atom, $-\text{CH}_2\text{F}$, $-\text{CHF}_2$, $-\text{CF}_3$ or $-\text{CN}$,

R^{13} is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a fluorine atom or a chlorine atom,

30 Z is an oxygen atom or a sulfur atom,

W is any group selected from the formulas (XVI) to (XX)

[0028]



[0029]

5 (wherein

R⁷ is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a fluorine atom or a chlorine atom,

R⁸ is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a fluorine atom or a chlorine atom,

10 R⁹ is a hydrogen atom, methyl, ethyl, n-propyl, isopropyl, methoxy, ethoxy, n-propoxy, isopropoxy, a fluorine atom, a chlorine atom, -OH, -NR¹⁰R¹¹, -CH₂OR¹⁰, -CF₃, -OCF₃, -CN, -C(O)NR¹⁰R¹¹, the formula (III) or (IV),

R¹⁰ is a hydrogen atom, methyl, ethyl, n-propyl, isopropyl, 15 tert-butyl, isobutyl or -CH₂CH₂OMe,

R¹¹ is a hydrogen atom, methyl, ethyl, n-propyl, isopropyl, tert-butyl, isobutyl or -CH₂CH₂OMe,

n is an integer of 2 to 4,

or a pharmaceutically acceptable acid addition salt thereof,

20 [2'] the compound of the above-mentioned [1] or [1'], wherein

R¹ is a hydrogen atom or alkyl having 1 to 4 carbon atoms,

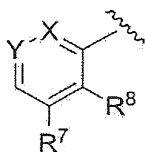
R² is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms or halogen,

R⁶ is alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4

25 carbon atoms, halogen, -CN, -CF₃, -OH or -NR^{6a}R^{6b} (wherein R^{6a} is alkyl having 1 to 4 carbon atoms, R^{6b} is alkyl having 1 to 4 carbon atoms),

W is a compound of the formula (II):

[0030]



30 (II)

[0031]

(wherein

R⁷ is a hydrogen atom, alkoxy having 1 to 4 carbon atoms or halogen,

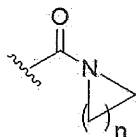
R^8 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms or halogen,

X is $-CR^9=$ or $-N=$,

Y is $-CR^9=$ or $-N=$,

- 5 R^9 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, $-OH$, $-NR^{10}R^{11}$, $-CH_2OH$, $-C(O)OR^{10}$, $-C(O)NR^{10}R^{11}$ or the formula (IV):

[0032]



(IV)

- 10 [0033]

(wherein n is an integer of 1 to 4),

R^{10} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-CH_2CF_3$, $-CH_2CH_2OH$ or $-CH_2CH_2OMe$, and

R^{11} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-$

- 15 CH_2CF_3 , $-CH_2CH_2OH$ or $-CH_2CH_2OMe$,

or a pharmaceutically acceptable acid addition salt thereof,

[3] the compound of the above-mentioned [2], wherein

R^1 is a hydrogen atom or methyl,

R^2 is methyl, methoxy, ethoxy or a fluorine atom,

- 20 R^3 is a hydrogen atom,

R^4 is a hydrogen atom,

W is the formula (XVI)

(wherein

R^9 is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a

- 25 fluorine atom or a chlorine atom),

R^{12} is methyl, ethyl, methoxy, a fluorine atom or a chlorine atom,

R^{13} is a hydrogen atom, methyl, methoxy, a fluorine atom or a chlorine atom,

- 30 or a pharmaceutically acceptable acid addition salt thereof,

[4] the compound of the above-mentioned [3], wherein

R^2 is methoxy, and

R^5 is the formula (V), (VIII), (IX) or (X)

(wherein

- 35 R^{13} is a hydrogen atom or a fluorine atom),

or a pharmaceutically acceptable acid addition salt thereof,

[5] the compound of the above-mentioned [2], wherein

R^1 is a hydrogen atom,

R^2 is methoxy or ethoxy,

R^3 is a hydrogen atom,
 R^4 is a hydrogen atom,
W is the formula (XVI)
(wherein
5 R^9 is $-C(O)NR^{10}R^{11}$ or the formula (IV),
 R^{10} is a hydrogen atom, methyl or ethyl,
 R^{11} is a hydrogen atom, methyl or ethyl,
n is an integer of 2 to 3),
 R^{12} is methyl, ethyl, methoxy, a fluorine atom or a chlorine
10 atom,
 R^{13} is a hydrogen atom, methyl, methoxy, a fluorine atom or a
chlorine atom,
or a pharmaceutically acceptable acid addition salt thereof,
[6] the compound of the above-mentioned [5], wherein
15 R^2 is methoxy,
 R^5 is the formula (V), (VIII), (IX) or (X)
(wherein
 R^{13} is a hydrogen atom),
or a pharmaceutically acceptable acid addition salt thereof,
20 [7] the compound of the above-mentioned [2], wherein
 R^1 is a hydrogen atom or methyl,
 R^2 is methyl, methoxy, ethoxy or a fluorine atom,
 R^3 is a hydrogen atom,
 R^4 is a hydrogen atom,
25 W is the formula (XVII),
(wherein
 R^9 is a hydrogen atom, methyl, methoxy, a fluorine atom or a
chlorine atom)
 R^{12} is methyl, ethyl, methoxy, a fluorine atom or a chlorine
30 atom,
 R^{13} is a hydrogen atom, methyl, methoxy, a fluorine atom or a
chlorine atom,
or a pharmaceutically acceptable acid addition salt thereof,
[8] the compound of the above-mentioned [7], wherein
35 R^1 is a hydrogen atom,
 R^2 is methoxy,
 R^5 is the formula (V), (VIII), (IX), (X) or (XI),
(wherein
 R^{12} is methyl, methoxy, a fluorine atom or a chlorine atom,
40 R^{13} is a hydrogen atom or a fluorine atom)
 R^9 is a hydrogen atom or a fluorine atom,
or a pharmaceutically acceptable acid addition salt thereof,
[9] the compound of the above-mentioned [2], wherein

R^1 is a hydrogen atom or methyl,
 R^2 is methyl, methoxy, ethoxy or a fluorine atom,
 R^3 is a hydrogen atom,
 R^4 is a hydrogen atom,
5 W is the formula (XVIII),
 (wherein
 R^7 is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a
 fluorine atom or a chlorine atom,
 R^9 is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a
 10 fluorine atom, a chlorine atom, $-CF_3$ or $-NR^{10}R^{11}$,
 R^{10} is a hydrogen atom, methyl, ethyl, n-propyl, isopropyl or
 isobutyl,
 R^{11} is a hydrogen atom or methyl)
 R^{12} is methyl, ethyl, methoxy, a fluorine atom or a chlorine
 15 atom,
 R^{13} is a hydrogen atom, methyl, methoxy, a fluorine atom or a
 chlorine atom,
 or a pharmaceutically acceptable acid addition salt thereof,
 [10] the compound of the above-mentioned [9], wherein
 20 R^2 is methoxy,
 R^5 is the formula (V), (VIII), (IX) or (X),
 (wherein
 R^{12} is methyl, methoxy, a fluorine atom or a chlorine atom,
 R^{13} is a hydrogen atom or a fluorine atom)
 25 R^9 is methoxy, ethoxy, a fluorine atom or a chlorine atom,
 or a pharmaceutically acceptable acid addition salt thereof,
 [11] the compound of the above-mentioned [2], wherein
 R^1 is a hydrogen atom or methyl,
 R^2 is methyl, methoxy, ethoxy or a fluorine atom,
 30 R^3 is a hydrogen atom,
 R^4 is a hydrogen atom,
 W is the formula (XVIII),
 (wherein
 R^7 is a hydrogen atom or a fluorine atom,
 35 R^9 is $-C(O)NR^{10}R^{11}$ or the formula (IV),
 R^{10} is a hydrogen atom, methyl, ethyl, n-propyl, isopropyl or
 tert-butyl,
 R^{11} is a hydrogen atom, methyl or ethyl,
 n is an integer of 2 to 4)
 40 R^{12} is methyl, ethyl, methoxy, a fluorine atom, a chlorine atom,
 $-CF_3$ or $-CN$,
 R^{13} is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a
 fluorine atom or a chlorine atom,

or a pharmaceutically acceptable acid addition salt thereof,
 [12] the compound of the above-mentioned [11], wherein
 R^2 is methoxy,
 R^5 is the formula (V), (VIII), (IX), (X) or (XI),
 5 (wherein
 R^{12} is methyl, methoxy, a fluorine atom or a chlorine atom,
 R^{13} is a hydrogen atom or a fluorine atom,
 Z is a sulfur atom), and
 R^7 is a hydrogen atom,
 10 or a pharmaceutically acceptable acid addition salt thereof,
 [13] the compound of the above-mentioned [2], wherein
 R^1 is a hydrogen atom or methyl,
 R^2 is methyl, methoxy, ethoxy or a fluorine atom,
 R^3 is a hydrogen atom,
 15 R^4 is a hydrogen atom,
 W is the formula (XIX),
 (wherein
 R^8 is methyl, ethyl, methoxy, ethoxy, a fluorine atom or a
 chlorine atom,
 20 R^9 is methyl, ethyl, methoxy, ethoxy, a fluorine atom, a
 chlorine atom or $-NR^{10}R^{11}$,
 R^{10} is a hydrogen atom, methyl, ethyl, n-propyl, isopropyl or
 isobutyl,
 R^{11} is a hydrogen atom or methyl)
 25 R^{12} is methyl, ethyl, methoxy, a fluorine atom or a chlorine
 atom,
 R^{13} is a hydrogen atom, methyl, methoxy, a fluorine atom or a
 chlorine atom,
 or a pharmaceutically acceptable acid addition salt thereof,
 30 [14] the compound of the above-mentioned [13], wherein
 R^2 is methoxy,
 R^5 is the formula (V), (VIII), (IX) or (X)
 (wherein
 R^{12} is methyl, methoxy, a fluorine atom or a chlorine atom,
 35 R^{13} is a hydrogen atom, a fluorine atom or a chlorine atom),
 or a pharmaceutically acceptable acid addition salt thereof,
 [15] the compound of the above-mentioned [2], wherein
 R^1 is a hydrogen atom or methyl,
 R^2 is methyl, methoxy, ethoxy or a fluorine atom,
 40 R^3 is a hydrogen atom,
 R^4 is a hydrogen atom,
 W is the formula (XIX),
 (wherein

R^8 is a fluorine atom,
 R^9 is $-C(O)NR^{10}R^{11}$ or the formula (IV),
 R^{10} is a hydrogen atom, methyl, ethyl, n-propyl, isopropyl or tert-butyl,
5 R^{11} is a hydrogen atom, methyl or ethyl,
n is an integer of 2 to 4)
 R^{12} is methyl, ethyl, methoxy, a fluorine atom or a chlorine atom,
 R^{13} is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a
10 fluorine atom or a chlorine atom,
or a pharmaceutically acceptable acid addition salt thereof,
[16] the compound of the above-mentioned [15], wherein
 R^1 is a hydrogen atom,
 R^2 is methoxy,
15 R^5 is the formula (V), (VIII), (IX) or (X),
(wherein
 R^{13} is a hydrogen atom or a fluorine atom),
or a pharmaceutically acceptable acid addition salt thereof,
[17] the compound of the above-mentioned [2], wherein
20 R^1 is a hydrogen atom or methyl,
 R^2 is methyl, methoxy, ethoxy or a fluorine atom,
 R^3 is a hydrogen atom,
 R^4 is a hydrogen atom,
W is the formula (XX),
25 (wherein
 R^7 is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a fluorine atom or a chlorine atom,
 R^9 is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a fluorine atom, a chlorine atom or $-CF_3$)
30 R^{12} is methyl, ethyl, methoxy, a fluorine atom or a chlorine atom,
 R^{13} is a hydrogen atom, methyl, methoxy, a fluorine atom or a chlorine atom,
or a pharmaceutically acceptable acid addition salt thereof,
35 [18] the compound of the above-mentioned [17], wherein
 R^2 is methoxy,
 R^5 is the formula (V), (VIII), (IX) or (X),
(wherein
 R^{12} is methyl, methoxy, a fluorine atom or a chlorine atom,
40 R^{13} is a hydrogen atom or a fluorine atom)
 R^7 is a hydrogen atom,
or a pharmaceutically acceptable acid addition salt thereof,
[19] the compound of the above-mentioned [2], wherein

R^1 is a hydrogen atom or methyl,
 R^2 is methyl, methoxy, ethoxy or a fluorine atom,
 R^3 is a hydrogen atom,
 R^4 is a hydrogen atom,
5 W is the formula (XX),
 (wherein
 R^7 is a hydrogen atom or a fluorine atom,
 R^9 is $-C(O)NR^{10}R^{11}$,
 R^{10} is a hydrogen atom, methyl, ethyl, n-propyl, isopropyl or
10 tert-butyl,
 R^{11} is a hydrogen atom, methyl or ethyl)
 R^{12} is methyl, ethyl, methoxy, a fluorine atom or a chlorine
 atom, and
 R^{13} is a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a
15 fluorine atom or a chlorine atom,
 or a pharmaceutically acceptable acid addition salt thereof,
 [20] the compound of the above-mentioned [19], wherein
 R^2 is methoxy,
 R^5 is the formula (V), (VIII), (IX) or (X)
20 (wherein
 R^{12} is methyl, methoxy, a fluorine atom or a chlorine atom,
 R^{13} is a hydrogen atom or a fluorine atom), and
 R^{10} is methyl or ethyl,
 or a pharmaceutically acceptable acid addition salt thereof,
25 [21] a medicament comprising the compound of any of the above-
 mentioned [1] to [20] or a pharmaceutically acceptable acid
 addition salt thereof,
 [22] an orexin receptor agonist comprising the compound of any
 of the above-mentioned [1] to [20] or a pharmaceutically
30 acceptable acid addition salt thereof,
 [23] an anti-narcolepsy agent comprising the compound of any
 of the above-mentioned [1] to [20] or a pharmaceutically
 acceptable acid addition salt thereof,
 [24] an agent for the treatment or prophylaxis of narcolepsy,
35 comprising the compound of any of the above-mentioned [1] to
 [20] or a pharmaceutically acceptable acid addition salt
 thereof,
 [25] a method for the treatment or prophylaxis of narcolepsy,
 comprising administering an effective amount of the compound
40 of any of the above-mentioned [1] to [20] or a
 pharmaceutically acceptable acid addition salt thereof, and
 [26] the compound of any of the above-mentioned [1] to [20] or
 a pharmaceutically acceptable acid addition salt thereof, for

the treatment or prophylaxis of narcolepsy.

Effect of the Invention

[0034]

The compound represented by the formula (I) or a
5 pharmaceutically acceptable acid addition salt thereof of the
present invention has a superior OX2R agonist activity.

Description of Embodiments

[0035]

The following terms used in the present specification are
10 as defined below unless otherwise specified.

The "alkyl" in the present specification means a
monovalent straight chain or branched saturated hydrocarbon
group composed of a carbon atom and a hydrogen atom. For
example, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl,
15 sec-butyl, tert-butyl, pentyl, hexyl and the like can be
mentioned.

The "alkoxy" in the present specification means an oxy
group to which the above-mentioned "alkyl" is bonded. For
example, methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy,
20 isobutoxy, tert-butoxy and the like can be mentioned.

The "halogen" in the present specification means a
fluorine atom, a chlorine atom, a bromine atom or an iodine
atom.

The "cycloalkyl" in the present specification means a
25 monocyclic aliphatic carbon cyclic group having 3 to 8 carbon
atoms. For example, cyclopropyl, cyclobutyl, cyclopentyl,
cyclohexyl, cycloheptyl, cyclooctyl and the like can be
mentioned.

The "aryl" in the present specification means a
30 monocyclic or fused aromatic carbon cyclic group having 6 to
10 carbon atoms. For example, phenyl, 1-naphthyl, 2-naphthyl
and the like can be mentioned.

The "heteroaryl" in the present specification means a 5-
to 10-membered monocyclic or bicyclic aromatic heterocyclic
35 group containing, as a ring-constituting atom besides carbon
atom, one or two kinds of 1 to 4 hetero atoms selected from
oxygen atom, sulfur atom and nitrogen atom. For example,
thienyl, furyl, pyrrolyl, imidazolyl, pyrazolyl, triazolyl,
tetrazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl,
40 pyridyl, pyridazinyl, pyrimidinyl, furazanyl, pyrazinyl,
thiadiazolyl, oxadiazolyl, benzofuryl, benzothienyl,
benzimidazolyl, benzothiazolyl, quinolyl, isoquinolyl,
cinnolyl, quinazolyl, quinoxalyl, phthalazinyl, 1H-indazolyl

and the like can be mentioned.

The "anti-narcolepsy agent" in the present specification means an agent for the treatment or prophylaxis of narcolepsy.
[0036]

5 The definition of each symbol in the formula (I) is explained in detail below.

Examples of the "alkyl having 1 to 4 carbon atoms" for R^1 , R^2 , R^3 , R^4 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} , R^a or R^b include methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl,
10 tert-butyl and the like; however, these are not limitative.

Examples of the "halogen" for R^1 , R^2 , R^6 , R^7 , R^8 or R^9 include a fluorine atom, a chlorine atom, a bromine atom, an iodine atom and the like; however, these are not limitative.

Examples of the "alkoxy having 1 to 4 carbon atoms" for
15 R^2 , R^6 , R^7 , R^8 or R^9 include methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, tert-butoxy and the like can be mentioned; however, these are not limitative.

[0037]

Preferable embodiment of each symbol in the formula (I)
20 is explained below.

As R^1 , a hydrogen atom, methyl, ethyl, a fluorine atom or a chlorine atom is preferable, and a hydrogen atom or methyl is particularly preferable.

As R^2 , a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a
25 fluorine atom or a chlorine atom is preferable, methyl, methoxy, ethoxy or a fluorine atom is further preferable, and methoxy is particularly preferable.

As R^{2a} and R^{2b} , each is independently preferably a hydrogen atom, methyl, ethyl, n-propyl, isopropyl, n-butyl,
30 isobutyl, sec-butyl or tert-butyl, and a hydrogen atom or methyl is particularly preferable.

As R^3 , a hydrogen atom, methyl or ethyl is preferable, a hydrogen atom or methyl is further preferable, and a hydrogen atom is particularly preferable.

35 As R^4 , a hydrogen atom, methyl or ethyl is preferable, a hydrogen atom or methyl is further preferable, and a hydrogen atom is particularly preferable.

As R^5 , phenyl, naphthyl, pyridyl, thienyl, furyl or pyrazolyl is preferable, phenyl, pyridyl or thienyl is further
40 preferable, and phenyl or pyridyl is particularly preferable.

As R^6 , methyl, ethyl, methoxy, ethoxy, a fluorine atom or a chlorine atom is preferable, methyl, methoxy, a fluorine atom or a chlorine atom is further preferable, and methoxy or

a chlorine atom is particularly preferable.

As R^{6a} and R^{6b} , each is independently preferably a hydrogen atom, methyl or ethyl, and a hydrogen atom or methyl is particularly preferable.

5 In the structure of W shown by the above-mentioned formula (II), R^7 is preferably a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a fluorine atom or a chlorine atom, further preferably a hydrogen atom or a fluorine atom, particularly preferably a hydrogen atom.

10 As R^8 , a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a fluorine atom or a chlorine atom is preferable, and a hydrogen atom or a fluorine atom is particularly preferable.

As R^9 , methyl, ethyl, methoxy, ethoxy, a fluorine atom, a chlorine atom, $-CF_3$, $-CN$, $-NR^{10}R^{11}$, $-C(O)NR^{10}R^{11}$ or the formula
15 (IV) is preferable, methoxy, a fluorine atom, a chlorine atom, $-C(O)NR^{10}R^{11}$ or the formula (IV) is further preferable, and methoxy or $-C(O)NR^{10}R^{11}$ is particularly preferable.

As R^{10} , a hydrogen atom, methyl, ethyl, n-propyl, isopropyl, isobutyl or tert-butyl is preferable, and methyl,
20 ethyl or isopropyl is particularly preferable.

As R^{11} , a hydrogen atom, methyl, ethyl, n-propyl, isopropyl, isobutyl or tert-butyl is preferable, a hydrogen atom, methyl or ethyl is further preferable, and a hydrogen atom or methyl is particularly preferable.

25 As R^{12} , methyl, ethyl, n-propyl, isopropyl, tert-butyl, methoxy, ethoxy, a fluorine atom, a chlorine atom, trifluoromethyl or cyano is preferable, methyl, ethyl, methoxy, ethoxy, fluoro or chloro is further preferable, and methyl, methoxy, fluoro or chloro is particularly preferable.

30 As R^{13} , a hydrogen atom, methyl, ethyl, methoxy, ethoxy, a fluorine atom or a chlorine atom is preferable, a hydrogen atom, methyl, methoxy, a fluorine atom or a chlorine atom is further preferable, and a hydrogen atom or a fluorine atom is particularly preferable.

35 In the structure of R^9 shown by the above-mentioned formula (IV), as n, 2, 3 or 4 is preferable, and 2 or 3 is particularly preferable.

[0038]

As a preferable example of the compound represented by
40 the above-mentioned formula (I) of the present invention, the following compound can be mentioned.

[Compound A]

Compound (I) wherein

R¹ is a hydrogen atom or alkyl having 1 to 4 carbon atoms (e.g., methyl),

R² is a hydrogen atom, alkyl having 1 to 4 carbon atoms (e.g., methyl), alkoxy having 1 to 4 carbon atoms (e.g., methoxy or
5 ethoxy) or halogen (e.g., fluorine atom),

R³ is a hydrogen atom or alkyl having 1 to 4 carbon atoms (e.g., methyl),

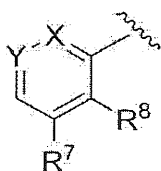
R⁴ is a hydrogen atom or alkyl having 1 to 4 carbon atoms (e.g., methyl),

10 R⁵ is aryl (e.g., phenyl) or heteroaryl (e.g., pyridyl, pyrazolyl, thienyl, imidazolyl, furyl) (wherein aryl or heteroaryl is optionally substituted by optionally selected 1 to 4 R⁶),

R⁶ is alkyl having 1 to 4 carbon atoms (e.g., methyl), alkoxy
15 having 1 to 4 carbon atoms (e.g., methoxy), halogen (e.g., a fluorine atom, a chlorine atom, a bromine atom), -CN, -CF₃, -OH or -NR^{6a}R^{6b} (wherein R^{6a} is alkyl having 1 to 4 carbon atoms (e.g., methyl), R^{6b} is alkyl having 1 to 4 carbon atoms (e.g., methyl)),

20 W is the formula (II):

[0039]



(II)

[0040]

(wherein

25 R⁷ is a hydrogen atom, alkoxy having 1 to 4 carbon atoms (e.g., methoxy) or halogen (e.g., a fluorine atom, a chlorine atom), R⁸ is a hydrogen atom, alkyl having 1 to 4 carbon atoms (e.g., methyl), alkoxy having 1 to 4 carbon atoms (e.g., methoxy) or halogen (e.g., a fluorine atom, a chlorine atom),

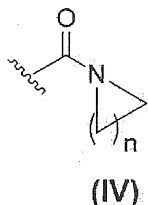
30 X is -CR⁹= or -N=,

Y is -CR⁹= or -N=,

R⁹ is a hydrogen atom, alkyl having 1 to 4 carbon atoms (e.g., methyl), alkoxy having 1 to 4 carbon atoms (e.g., methoxy), halogen (e.g., a fluorine atom, a chlorine atom), -OH, -NR¹⁰R¹¹,

35 -CH₂OH, -C(O)OR¹⁰, -C(O)NR¹⁰R¹¹ or the formula (IV):

[0041]



[0042]

(wherein n is an integer of 1 to 4) (e.g., azetidine-1-carbonyl, pyrrolidine-1-carbonyl, piperidine-1-carbonyl),

5 R^{10} is a hydrogen atom, alkyl having 1 to 4 carbon atoms (e.g., methyl, isobutyl, tert-butyl), $-CH_2CF_3$, $-CH_2CH_2OH$ or $-CH_2CH_2OMe$, and

R^{11} is a hydrogen atom, alkyl having 1 to 4 carbon atoms (e.g., methyl, isobutyl, tert-butyl), $-CH_2CF_3$, $-CH_2CH_2OH$ or $-CH_2CH_2OMe$),

10 or a pharmaceutically acceptable acid addition salt thereof.

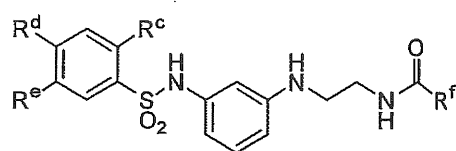
[0043]

As a pharmaceutically acceptable acid addition salt of the compound of the formula (I) of the present invention, inorganic acid salts such as hydrochloride, sulfate, nitrate, hydrobromide, hydroiodide, phosphate and the like, organic carbonates such as acetate, lactate, citrate, oxalate, glutarate, malate, tartrate, fumarate, mandelate, maleate, benzoate, phthalate and the like, organic sulfonates such as methanesulfonate, ethanesulfonate, benzenesulfonate, p-toluenesulfonate, camphorsulfonate and the like, and the like can be mentioned; however, these are not limitative. Of these, hydrochloride, hydrobromide, phosphate, tartrate, methanesulfonate or camphorsulfonate is preferable, and hydrochloride, tartrate or methanesulfonate is further preferably used; however, these are also not limitative.

[0044]

Of the compounds of the formula (I) of the present invention, specific preferable examples thereof are shown in Table 1 to Table 8; however, they do not limit the present invention.

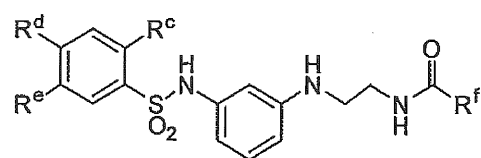
[0045]
Table 1



(Ia)

R ^c	R ^d	R ^e	R ^f
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		

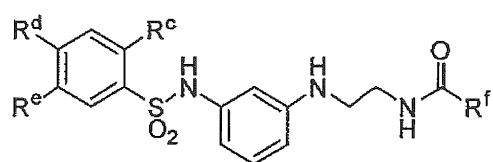
[0046]
Table 2



(Ia)

R ^a	R ^d	R ^e	R ^f
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		

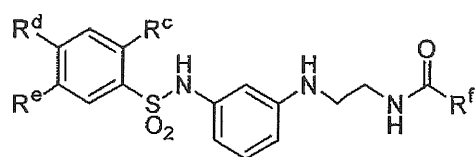
[0047]
Table 3



(Ia)

R ^c	R ^d	R ^e	R ^f
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		

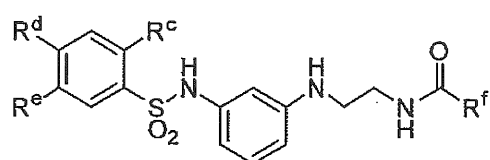
[0048]
Table 4



(1a)

R ^c	R ^d	R ^e	R ^f
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		

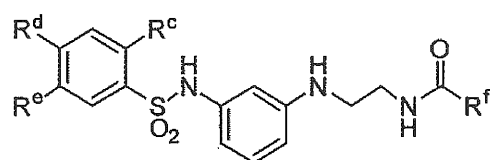
[0049]
Table 5



(1a)

R ^c	R ^d	R ^e	R ^f
OMe	Me		
OMe	H		
OMe	Me		
OMe	H		
OMe	H		
OMe	H		
OMe	Me		
OMe	Me		

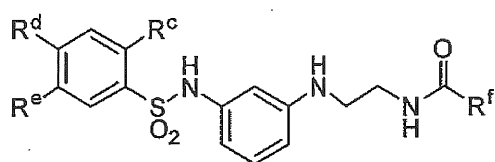
[0050]
Table 6



(1a)

R ^e	R ^d	R ^a	R ^f
OMe	H		
OMe	Me		
H	H		
Me	H		
F	H		
OEt	H		
OMe	H		
OMe	H		

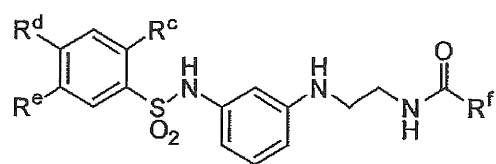
[0051]
Table 7



(Ia)

R ^c	R ^d	R ^e	R ^f
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		

[0052]
Table 8



(Ia)

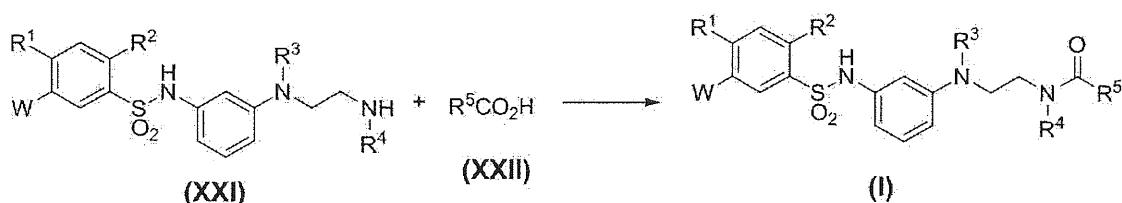
R ^c	R ^d	R ^e	R ^f
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		
OMe	H		

[0053]

The compound of the above-mentioned formula (I) of the present invention can be produced by an appropriate method based on the characteristics derived from the basic skeleton and substituents thereof. While the starting materials and reagents to be used for the production of these compounds are generally available or can be synthesized by a method known to those of ordinary skill in the art, which follows the procedures described in reference documents such as Organic Reactions (Wiley&Sons), Fieser and Fieser's Reagent for Organic Synthesis (Wiley&Sons) and the like.

As a specific production method of the compound of the above-mentioned formula (I) of the present invention, for example, the methods shown in Schemes 1 to 4 can be mentioned.

[0054]



Scheme 1

[0055]

wherein R¹, R², R³, R⁴, R⁵ and W are as defined above.

Specifically, the compound of the formula (I) can be obtained by, for example, amidating an amine derivative represented by the formula (XXI) with a carboxylic acid represented by the formula (XXII).

As a solvent, halogenated solvents such as dichloromethane, chloroform, 1,2-dichloroethane and the like, ether solvents such as diethyl ether, tetrahydrofuran (THF), 1,2-dimethoxyethane (DME), dioxane and the like, aprotic polar solvents such as N,N-dimethylformamide (DMF), dimethyl sulfoxide (DMSO) and the like, alcohol solvents such as methanol, ethanol, propanol and the like or a mixed solvent thereof can be used. In general, dichloromethane or THF is preferably used. Carboxylic acid (XXII) is used in an amount of 0.5 - 20 equivalents, preferably 0.5 - 10 equivalents, relative to amine derivative (XXI).

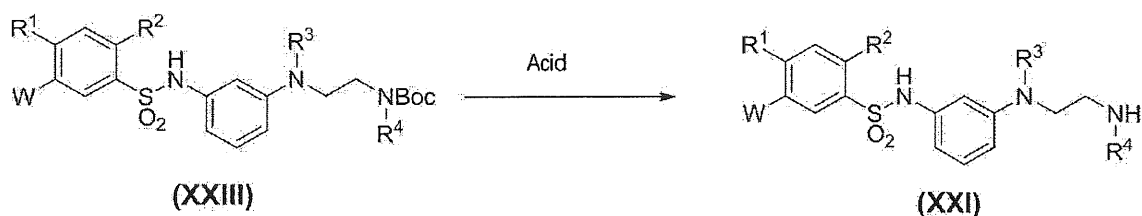
As the condensing agent, dicyclohexylcarbodiimide, 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (EDCI), benzotriazol-1-yloxytris(dimethylamino)phosphonium hexafluorophosphate (BOP) reagent and the like can be used,

and a BOP reagent is particularly preferably used. The condensing agent is used in an amount of 1.0 - 100 equivalents, preferably 1.0 - 10 equivalents, relative to amine derivative (XXI).

5 When a base is used, triethylamine, diisopropylethylamine, pyridine, N-methylmorpholine and the like can be used, and triethylamine or diisopropylethylamine is preferably used. The base is used in an amount of 3.0 - 100 equivalents, preferably 3.0 - 10 equivalents, relative to amine derivative (XXI).

10 The reaction temperature is generally -40 - 150°C, preferably 0 - 60°C. While the reaction time is appropriately selected according to the conditions such as reaction temperature and the like, it is generally about 20 min - 48 hr. While the concentration of substrate (XXI) in the reaction
15 system is not particularly limited, it is generally preferably 0.001 mmol/L - 1 mol/L.

[0056]



Scheme 2

[0057]

20 wherein R¹, R², R³, R⁴ and W are as defined above.

The amine derivative of the formula (XXI) can be obtained by, for example, deprotecting a tert-butoxycarbonyl (Boc) group of the derivative represented by the formula (XXIII) under acidic conditions.

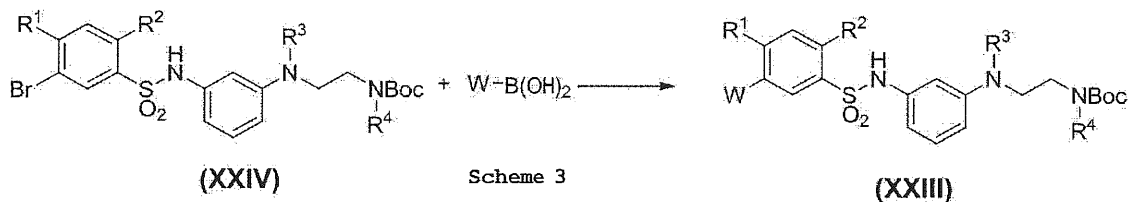
25 As a solvent, halogenated solvents such as dichloromethane, chloroform, 1,2-dichloroethane and the like, ether solvents such as THF, DME, dioxane and the like, alcohol solvents such as methanol, ethanol, propanol and the like or a mixed solvent thereof can be used. In general, dichloromethane
30 or dioxane is preferably used.

As the acid, organic acids such as trifluoroacetic acid, methanesulfonic acid, p-toluenesulfonic acid, trifluoromethanesulfonic acid and the like, or inorganic acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid,
35 sulfuric acid, nitric acid, phosphoric acid and the like can be used. In general, trifluoroacetic acid or hydrochloric acid is preferably used. In another method, a hydrogen chloride-

methanol solution, a hydrogen chloride-THF solution, a hydrogen chloride-ethyl acetate solution, or a hydrogen chloride-dioxane solution obtained by dissolving hydrogen chloride in an organic solvent is each independently used. In this case, particularly, preferable results are obtained by using a hydrogen chloride-methanol solution or a hydrogen chloride-THF solution. The acid is used in an amount of 1.0 - 100 equivalents, preferably 3.0 - 10 equivalents, relative to the formula (XXIII).

The reaction temperature is generally -40 - 150°C, preferably 0 - 80°C. While the reaction time is appropriately selected according to the conditions such as reaction temperature and the like, it is generally about 20 min - 48 hr. While the concentration of substrate (XXIII) in the reaction system is not particularly limited, it is generally preferably 0.001 mmol/L - 1 mol/L.

[0058]



[0059]

wherein R¹, R², R³, R⁴ and W are as defined above.

The derivative represented by the formula (XXIII) can be obtained by, for example, a Suzuki coupling of a derivative represented by the formula (XXIV) with a boronic acid derivative represented by the formula (XXV).

The Suzuki coupling reaction is performed in the presence of a palladium catalyst and a base, in the presence or absence of a phosphine ligand in a suitable solvent.

As a solvent, ether solvents such as THF, DME, dioxane and the like, aprotic polar solvents such as DMF, DMSO and the like, alcohol solvents such as methanol, ethanol, propanol and the like, aromatic solvents such as benzene, toluene, xylene and the like, water or a mixed solvent thereof can be used. In general, dioxane, DME, a mixed solvent of dioxane and water or a mixed solvent of DME and water is preferably used.

As boronic acid derivative (XXV), not only boronic acid but also boronates such as boronic acid pinacol ester, N-methyliminodiacetic acid (MIDA) boronate and the like, or

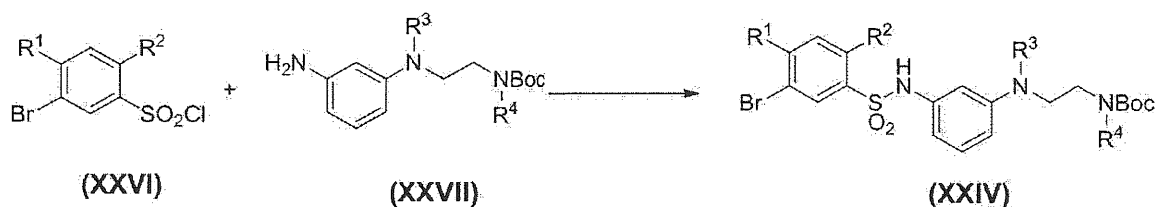
potassium trifluoroborate salt can be used. Particularly, boronic acid or boronic acid pinacol ester is preferably used. The boronic acid derivative (XXV) is used in an amount of 1.0 - 20 equivalents, preferably 1.0 - 10 equivalents, relative to the derivative of the formula (XXIV).

Examples of the palladium catalyst include tetrakis(triphenylphosphine)palladium, palladiumacetate, bis(triphenylphosphine)palladiumdichloride, bis(dibenzylideneacetone)palladium, bis(diphenylphosphino)ferrocenepalladiumdichloride and the like, and tetrakis(triphenylphosphine)palladium or bis(diphenylphosphino)ferrocene palladium dichloride is preferably used. The palladium catalyst is used in an amount of 0.001 - 1 equivalent, preferably 0.005 - 0.5 equivalent, relative to the derivative of the formula (XXIV).

Examples of the base include sodium carbonate, potassium carbonate, cesium carbonate, potassium phosphate, sodium hydroxide, potassium hydroxide, barium hydroxide, triethylamine, diisopropylethylamine and the like, and sodium carbonate or potassium carbonate is preferably used.

The reaction temperature is generally -40 - 150°C, preferably 20 - 110°C. While the reaction time is appropriately selected according to the conditions such as reaction temperature and the like, it is generally about 20 min - 48 hr. While the concentration of substrate (XXIV) in the reaction system is not particularly limited, it is generally preferably 0.001 mmol/L - 1 mol/L.

[0060]



Scheme 4

[0061]

wherein R¹, R², R³ and R⁴ are as defined above.

The derivative represented by the formula (XXIV) can be obtained by, for example, amidating the sulfonic acid chloride derivative of the formula (XXVI) with the amine derivative of the formula (XXVII).

As a solvent, halogenated solvents such as

dichloromethane, chloroform, 1,2-dichloroethane and the like, ether solvents such as diethyl ether, THF, DME, dioxane and the like, aprotic polar solvents such as DMF, DMSO and the like or a mixed solvent thereof can be used. In general, 5 dichloromethane or THF is preferably used. The sulfonic acid chloride derivative (XXVI) is used in an amount of 0.5 - 20 equivalents, preferably 1.0 - 10 equivalents, relative to amine derivative (XXVII).

Examples of the base include sodium carbonate, potassium 10 carbonate, cesium carbonate, potassium phosphate, sodium hydroxide, potassium hydroxide, barium hydroxide, triethylamine, diisopropylethylamine, pyridine, 4-dimethylaminopyridine and the like can be mentioned, triethylamine or pyridine is preferably used.

15 The reaction temperature is generally -40 - 150°C, preferably 0 - 80°C. While the reaction time is appropriately selected according to the conditions such as reaction temperature and the like, it is generally about 10 min - 48 hr. While the concentration of substrate (XXVII) in the reaction 20 system is not particularly limited, it is generally preferably 0.001 mmol/L - 1 mol/L.

[0062]

An orexin agonist containing the compound of the present invention is effective for not only human but also mammals 25 other than human, for example, mouse, rat, hamster, rabbit, cat, dog, bovine, sheep, monkey and the like.

Also, the compound of the present invention is used not only as an agent for the treatment or prophylaxis of narcolepsy as mentioned above but also can be used as a method 30 for the treatment or prophylaxis of narcolepsy, or for the production of a medicament for the treatment or prophylaxis of narcolepsy.

Furthermore, the compound of the present invention can also be used as an agent for the treatment or prophylaxis of 35 various neurological disorders related to an orexin receptor, such as sleeping sickness, sleeping disorder, waking disorder, jet lag, obesity, metabolic syndrome and the like.

When the compound of the present invention is clinically used as an agent for the treatment or prophylaxis of 40 narcolepsy, the medicament may be a free form of the compound

of the present invention or an acid addition salt thereof, or additives such as excipient, stabilizer, preservative, buffering agent, solubilizing agent, emulsifier, diluent, isotonicity agent and the like may be mixed as appropriate.

5 Examples of the administration form include oral preparations such as tablet, capsule, granule, powder, syrup and the like, parenteral agents such as injection, suppository, liquid and the like, topical administration of ointment, cream, adhesive preparation and the like, and the like.

10 The agent for the treatment or prophylaxis of narcolepsy of the present invention desirably contains 0.001 - 90 wt%, preferably 0.01 - 70 wt%, of the above-mentioned active ingredient. The amount thereof to be used is appropriately determined according to the symptom, age, body weight, and
15 administration method. In the case of injection for an adult, the amount of the active ingredient is 0.1 μ g - 1 g per day, 1 μ g - 1 g in the case of an oral preparation, and 1 μ g - 10 g in the case of an adhesive preparation, each of which can be administered in one to several portions.

20 In addition, the agent for the treatment or prophylaxis of narcolepsy of the present invention can also be used in combination with an agent for the treatment or prophylaxis of strong sleepiness and dozing during the day, an agent for the treatment or prophylaxis of deep sleep disorder, or an agent
25 for the treatment or prophylaxis of cataplexy.

As an agent for the treatment or prophylaxis of strong sleepiness and dozing during the day, central nervous system stimulants such as methylphenidate, pemoline, modafinil and the like, and the like can be mentioned.

30 As an agent for the treatment or prophylaxis of deep sleep disorder, tricyclic antidepressants such as triazolam, vegetamin B and the like, antianxiety drug and the like can be mentioned.

As an agent for the treatment or prophylaxis of cataplexy,
35 tricyclic antidepressants such as clomipramine hydrochloride, brotizolam, imipramine hydrochloride and the like, selective serotonin reuptake inhibitors (SSRI) such as fluvoxamine maleate, paroxetine, hydrochloride and the like, serotonin and noradrenaline reuptake inhibitors (SNRI) such as milnacipran
40 hydrochloride, duloxetine hydrochloride and the like, and the

like can be mentioned.

Examples

[0063]

The present invention is specifically explained in the following by referring to Examples. In the following Examples, the following abbreviations are used.

Boc: tert-butoxycarbonyl

Bn: benzyl

MIDA: N-methyliminodiacetic acid

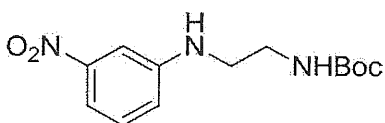
10 PMB: p-methoxybenzyl

[0064]

Reference Example 1

tert-butyl (2-((3-nitrophenyl)amino)ethyl)carbamate

[0065]



[0066]

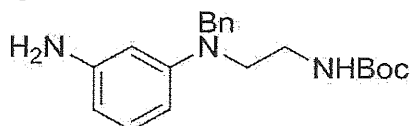
Under an argon atmosphere, to 3-fluoronitrobenzene (3.21 mL) was added ethylenediamine (25.0 mL), and the mixture was stirred at 100°C for 24 hr. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. To a solution of the obtained residue in dichloromethane (50 mL) were added triethylamine (4.60 mL), di-tert-butyl dicarbonate (6.55 g), and the mixture was stirred at room temperature for 2 hr. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/4→1/3) to give tert-butyl (2-((3-nitrophenyl)amino)ethyl)carbamate (5.31 g).

[0067]

35 Reference Example 2

tert-butyl (2-((3-aminophenyl)(benzyl)amino)ethyl)carbamate

[0068]



[0069]

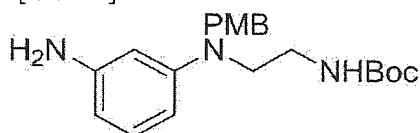
- (1) Under an argon atmosphere, to a solution of tert-butyl (2-((3-nitrophenyl)amino)ethyl)carbamate (4.46 g) in dichloromethane (30 mL) were added 50wt% aqueous sodium hydroxide solution (10 mL), benzyl bromide (2.84 mL) and tetrabutylammoniumiodide (587 mg), and the mixture was stirred at 20 - 30°C for 48 hr. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/6→1/4) to give tert-butyl (2-(benzyl(3-nitrophenyl)amino)ethyl)carbamate (4.62 g).
- (2) Under an argon atmosphere, to a mixed suspension of tert-butyl (2-(benzyl(3-nitrophenyl)amino)ethyl)carbamate in ethanol (50 mL) and water (20 mL) were added ammonium chloride (6.69 g) and iron powder (4.86 g), and the mixture was heated under reflux for 2 hr. The reaction mixture was filtered through celite, the filtrate was concentrated, saturated aqueous sodium hydrogen carbonate solution was added, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/ethyl acetate=2/1) to give tert-butyl (2-((3-aminophenyl)(benzyl)amino)ethyl)carbamate (4.58 g).

[0070]

Reference Example 3

- tert-butyl (2-((3-aminophenyl)(4-methoxybenzyl)amino)ethyl)carbamate

[0071]



[0072]

- (1) Under an argon atmosphere, to a solution of tert-butyl (2-((3-nitrophenyl)amino)ethyl)carbamate (4.46 g) in DMF (30 mL) were added potassium carbonate (1.21 g), sodium iodide (1.01 g) and 4-methoxybenzylchloride (1.00 mL), and the mixture was stirred at 60°C overnight. To this reaction mixture was added saturated brine, and the mixture was extracted with ethyl acetate. The organic layer was washed with water, dried over

sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/3→1/2→1/1) to give tert-butyl (2-(4-methoxybenzyl(3-

5 nitrophenyl)amino)ethyl)carbamate (2.78 g).

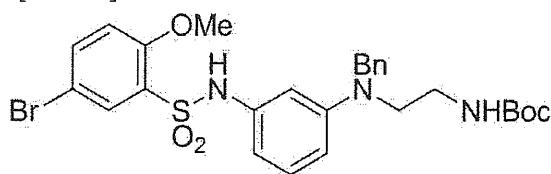
(2) To a solution of tert-butyl (2-(4-methoxybenzyl(3-nitrophenyl)amino)ethyl)carbamate (1.13 g) in methanol (20 mL) was added 10% palladium/carbon (339 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature for 1
10 hr. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-aminophenyl)(4-methoxybenzyl)amino)ethyl)carbamate (848 mg).

15 [0073]

Reference Example 4

tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate

[0074]



20

[0075]

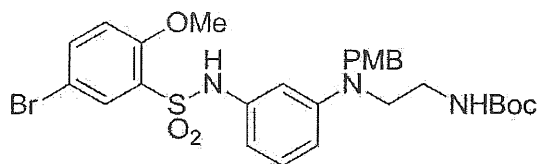
Under an argon atmosphere, to a solution of tert-butyl (2-((3-aminophenyl)(benzyl)amino)ethyl)carbamate (1.59 g) in dichloromethane (20 mL) were added pyridine (0.413 mL) and 5-
25 bromo-2-methoxybenzenesulfonyl chloride (1.33 g), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate,
30 filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2) to give tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (2.53 g).

35 [0076]

Reference Example 5

tert-butyl (2-((3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate

40 [0077]



[0078]

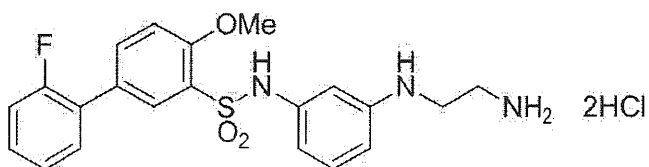
Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenylsulfonamido)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (848 mg) in dichloromethane (30 mL) were added pyridine (0.202 mL) and 5-bromo-2-methoxybenzenesulfonyl chloride (651 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=2/3→1/1) to give tert-butyl (2-((3-(5-bromo-2-methoxyphenylsulfonamido)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (1.15 g).

[0079]

Reference Example 6

N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride

[0080]



[0081]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenylsulfonamido)phenyl)amino)ethyl)carbamate (2.26 g) in DME (30 mL) were added 2-fluorophenylboronic acid (748 mg), sodium carbonate (811 mg), water (3.83 mL) and tetrakis(triphenylphosphine)palladium (221 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2→1/1) to give tert-butyl (2-((benzyl)(3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)carbamate (2.15 g).

(2) To a solution of tert-butyl (2-(benzyl(3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-

ylsulfonamide)phenyl)amino)ethyl)carbamate (2.15 g) in methanol (40 mL) was added 10% palladium/carbon (645 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature for 3 hr. The reaction mixture was filtered
5 through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) to give tert-butyl (2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (1.61 g).

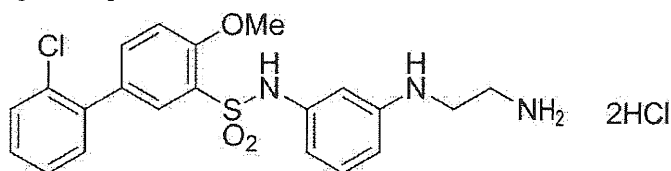
10 (3) To tert-butyl (2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (1.61 g) was added 10% hydrogen chloride-methanol solution (20 mL), and the mixture was stirred at 50°C for 3 hr. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-2'-
15 fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (1.55 g).

[0082]

Reference Example 7

N-(3-((2-aminoethyl)amino)phenyl)-2'-chloro-4-methoxy-[1,1'-
20 biphenyl]-3-sulfonamide dihydrochloride

[0083]



[0084]

(1) To a suspension of tert-butyl (2-((3-
25 nitrophenyl)amino)ethyl)carbamate (1.07 g) in methanol (15 mL) was added 10% palladium/carbon (321 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was
30 purified by amine silica gel column chromatography (eluate: ethyl acetate/hexane=2/1), methanol (1.5 mL) and 10% hydrogen chloride-methanol solution (1.5 mL) were further added, and the mixture was stirred at room temperature for 30 min. The reaction mixture was concentrated to give tert-butyl (2-((3-
35 aminophenyl)amino)ethyl)carbamate dihydrochloride (845 mg).

(2) Under an argon atmosphere, to a suspension of tert-butyl (2-((3-aminophenyl)amino)ethyl)carbamate dihydrochloride (143 mg) in dichloromethane (5.0 mL) were added pyridine (0.133 mL) and 5-bromo-2-methoxybenzenesulfonyl chloride (143 mg), and

the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) to give tert-butyl (2-((3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (133 mg).

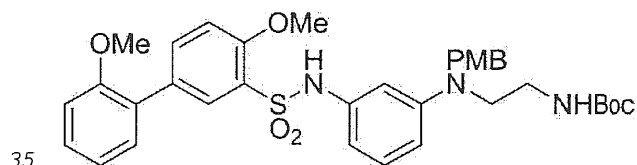
(3) Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (79.4 mg) in DME (2.0 mL) were added 2-chlorophenylboronic acid (34.2 mg), sodium carbonate (33.9 mg), water (0.160 mL) and tetrakis(triphenylphosphine)palladium (9.2 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: chloroform/methanol=40/1) to give tert-butyl (2-((3-(2'-chloro-4-methoxy-[1,1'-biphenyl]-3-yl)sulfonamide)phenyl)amino)ethyl)carbamate (53.6 mg).

(4) To tert-butyl (2-((3-(2'-chloro-4-methoxy-[1,1'-biphenyl]-3-yl)sulfonamide)phenyl)amino)ethyl)carbamate (53.6 mg) was added 10% hydrogen chloride-methanol solution (1.0 mL), and the mixture was stirred at 50°C for 3 hr. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-2'-chloro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (49.9 mg).

[0085]

Reference Example 8

tert-butyl (2-((3-(2',4-dimethoxy-[1,1'-biphenyl]-3-yl)sulfonamide)phenyl) (4-methoxybenzyl)amino)ethyl)carbamate [0086]



[0087]

Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl) (4-

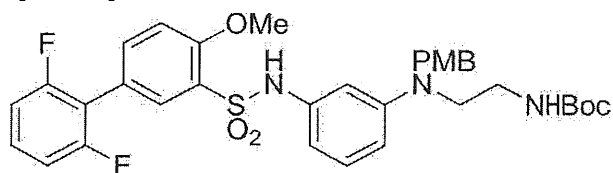
methoxybenzyl)amino)ethyl)carbamate (124 mg) in n-propanol (2.0 mL) were added 2-methoxyphenyltrifluoroborate potassium salt (51.4 mg), triethylamine (0.084 mL) and bis(diphenylphosphino)ferrocenepalladium dichloride dichloromethane complex (4.9 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) to give tert-butyl (2-((3-(2',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (116 mg).

[0088]

Reference Example 9

tert-butyl (2-((3-(2',6'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate

[0089]



[0090]

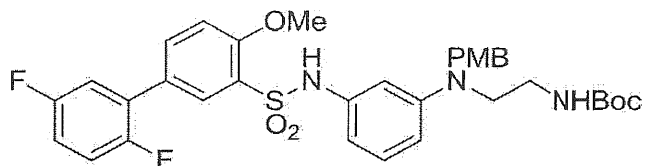
Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (100 mg) in DME (2.0 mL) were added 2,6-difluorophenyltrifluoroborate potassium salt (41.8 mg), triethylamine (0.067 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (3.9 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer column chromatography (eluate: ammonia-saturated chloroform) to give tert-butyl (2-((3-(2',6'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (69.6 mg).

[0091]

Reference Example 10

tert-butyl (2-((3-(2',5'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate

[0092]



[0093]

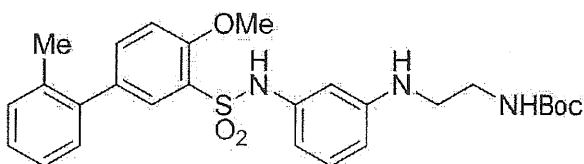
Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl carbamate (124 mg) in DME (4.0 mL) were added 2,5-difluorophenylboronic acid (344 mg), sodium carbonate (42.4 mg), water (0.200 mL) and tetrakis(triphenylphosphine)palladium (11.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: chloroform/methanol=40/1) to give tert-butyl (2-((3-(2',5'-difluoro-4-methoxy-[1,1'-biphenyl]-3-yl)sulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl carbamate (70.1 mg).

[0094]

Reference Example 11

tert-butyl (2-((3-(4-methoxy-2'-methyl-[1,1'-biphenyl]-3-yl)sulfonamide)phenyl)amino)ethyl carbamate

[0095]



[0096]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl carbamate (94.5 mg) in DME (4.0 mL) were added 2-trifluoromethylphenylboronic acid (41.8 mg), sodium carbonate (33.9 mg), water (0.160 mL) and tetrakis(triphenylphosphine)palladium (9.2 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) to give tert-butyl (2-((benzyl)(3-(4-methoxy-2'-(trifluoromethyl)-[1,1'-biphenyl]-3-yl)sulfonamide)phenyl)amino)ethyl carbamate (91.0 mg).

(2) To a solution of tert-butyl (2-((benzyl)(3-(4-methoxy-2'-(trifluoromethyl)-[1,1'-biphenyl]-3-

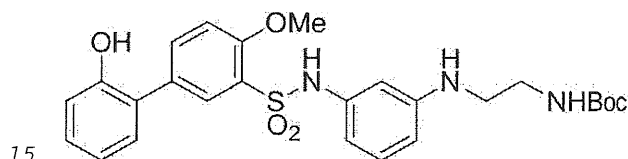
ylsulfonamide)phenyl)amino)ethyl)carbamate (91.0 mg) in methanol (3.0 mL) was added 10% palladium/carbon (27.3 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered
5 through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated chloroform) to give tert-butyl (2-((3-(4-methoxy-2'-methyl-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (11.7 mg).

10 [0097]

Reference Example 12

tert-butyl (2-((3-(2'-hydroxy-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate

[0098]



[0099]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl) (3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)amino)ethyl)carbamate (94.5
20 mg) in DME (4.0 mL) were added 2-benzyloxyphenylboronic acid (51.1 mg), sodium carbonate (33.9 mg), water (0.160 mL) and tetrakis(triphenylphosphine)palladium (9.2 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was
25 concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2) to give tert-butyl (2-((benzyl) (3-(2'-(benzyloxy)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (96.0 mg).

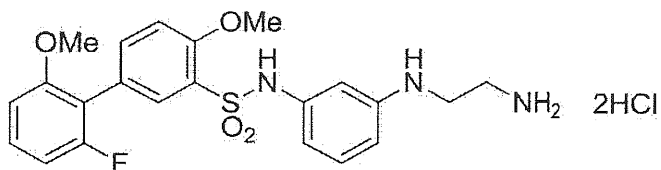
30 (2) To a solution of tert-butyl (2-((benzyl) (3-(2'-(benzyloxy)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (96.0 mg) in methanol (3.0 mL) was added 10% palladium/carbon (67.2 mg), and the mixture was stirred under a hydrogen atmosphere at
35 room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: chloroform/methanol=40/1) to give
40 tert-butyl (2-((3-(2'-hydroxy-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (7.2 mg).

[0100]

Reference Example 13

N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride

5 [0101]



[0102]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (118 mg) in DME (4.0 mL) were added 2-fluoro-6-methoxyphenylboronic acid (47.6 mg), sodium carbonate (42.4 mg), water (0.200 mL) and tetrakis(triphenylphosphine)palladium (11.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/1). To a solution of the obtained purified product in methanol (3.0 mL) was added 10% palladium/carbon (38.4 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated chloroform/ethyl acetate=6/1) to give tert-butyl (2-((3-(2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (34.6 mg).

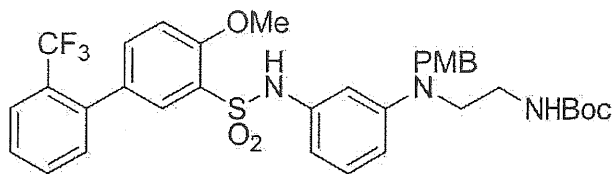
(2) To tert-butyl (2-((3-(2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (34.6 mg) was added 10% hydrogen chloride-methanol (1.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (36.6 mg).

[0103]

35 Reference Example 14

tert-butyl (2-((3-(4-methoxy-2'-(trifluoromethyl)-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate

[0104]



[0105]

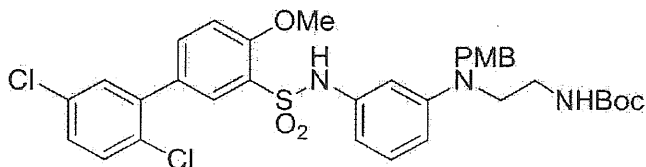
Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (160 mg) in DME (4.0 mL) were added 2-trifluoromethylphenylboronic acid (68.4 mg), sodium carbonate (55.1 mg), water (0.260 mL) and tetrakis(triphenylphosphine)palladium (15.0 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated chloroform) to give tert-butyl (2-((3-(4-methoxy-2'-(trifluoromethyl)-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (76.1 mg).

[0106]

Reference Example 15

tert-butyl (2-((3-(2',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate

[0107]



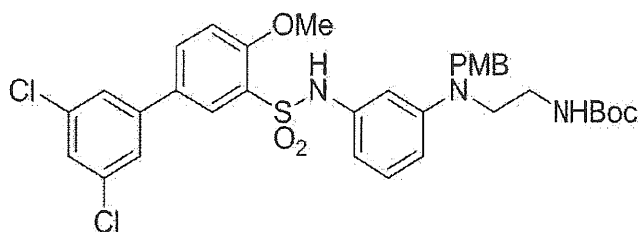
[0108]

Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (165 mg) in DME (4.0 mL) were added 2,5-dichlorophenylboronic acid (70.6 mg), sodium carbonate (56.2 mg), water (0.270 mL) and tetrakis(triphenylphosphine)palladium (15.0 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated chloroform) to give tert-butyl (2-((3-(2',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (177 mg).

[0109]

Reference Example 16

tert-butyl (2-((3-(3',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate
[0110]



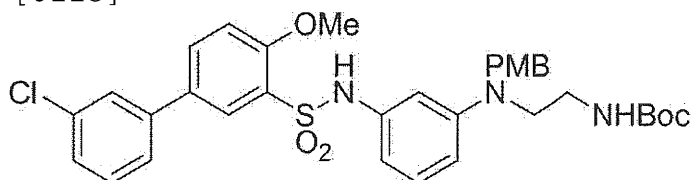
5 [0111]

Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (165 mg) in DME (4.0 mL) were added 3,5-dichlorophenylboronic acid (70.6 mg), sodium
10 carbonate (56.2 mg), water (0.270 mL) and tetrakis(triphenylphosphine)palladium (15.0 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative
15 thin layer chromatography (eluate: ammonia-saturated chloroform) to give tert-butyl (2-((3-(3',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (137 mg).

[0112]

20 Reference Example 17

tert-butyl (2-((3-(3'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate
[0113]



25 [0114]

Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (124 mg) in DME (4.0 mL) were added 3-chlorophenylboronic acid (43.8 mg), sodium
30 carbonate (42.4 mg), water (0.200 mL) and tetrakis(triphenylphosphine)palladium (11.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative
35 thin layer chromatography (eluate: ammonia-saturated

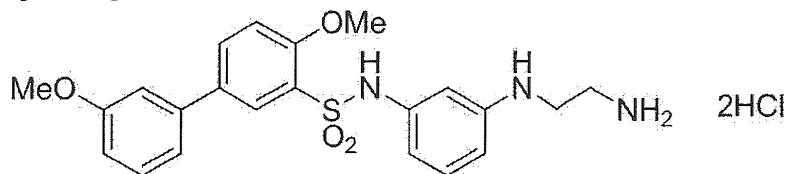
chloroform) to give tert-butyl (2-((3-(3'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (104 mg).

[0115]

5 Reference Example 18

N-(3-((2-aminoethyl)amino)phenyl)-3',4-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride

[0116]



10 [0117]

- (1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)amino)ethyl)carbamate (118 mg) in DME (4.0 mL) were added 3-methoxyphenylboronic acid (42.5 mg), sodium carbonate (42.4 mg), water (0.200 mL) and tetrakis(triphenylphosphine)palladium (11.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated chloroform). To a solution of the obtained residue in methanol (2.0 mL) was added 10% palladium/carbon (28.0 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) to give tert-butyl (2-((3-(3',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (43.0 mg).
- (2) To tert-butyl (2-((3-(3',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (43.0 mg) was added 10% hydrogen chloride-methanol (2.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-3',4-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (33.8 mg).

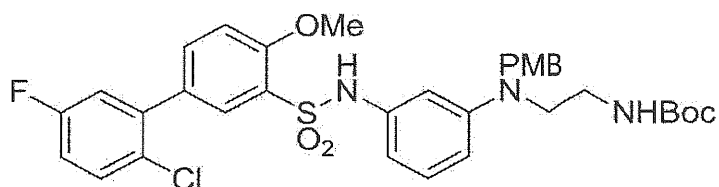
[0118]

Reference Example 19

tert-butyl (2-((3-(2'-chloro-5'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-

methoxybenzyl)amino)ethyl)carbamate

[0119]



[0120]

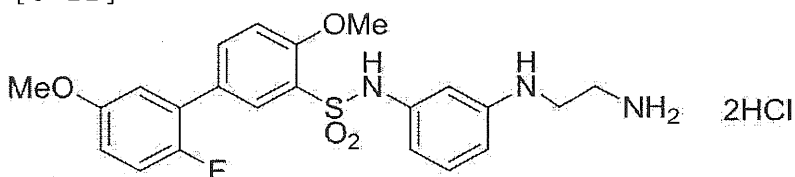
- 5 Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenyl)sulfonamido)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (124 mg) in DME (4.0 mL) were added 2-chloro-5-fluorophenylboronic acid (48.8 mg), sodium carbonate (42.4 mg), water (0.200 mL) and
- 10 tetrakis(triphenylphosphine)palladium (11.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2) to
- 15 give tert-butyl (2-((3-(2'-chloro-5'-fluoro-4-methoxy-[1,1'-biphenyl]-3-yl)sulfonamido)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (127 mg).

[0121]

Reference Example 20

- 20 N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride

[0122]



[0123]

- 25 (1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenyl)sulfonamido)phenyl)amino)ethyl)carbamate (118 mg) in DME (4.0 mL) were added 2-fluoro-5-methoxyphenylboronic acid (47.6 mg), sodium carbonate (42.4 mg), water (0.200 mL)
- 30 and tetrakis(triphenylphosphine)palladium (11.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated
- 35 chloroform) to give tert-butyl (2-((benzyl)(3-(2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-

ylsulfonamide)phenyl)amino)ethyl)carbamate (115 mg).

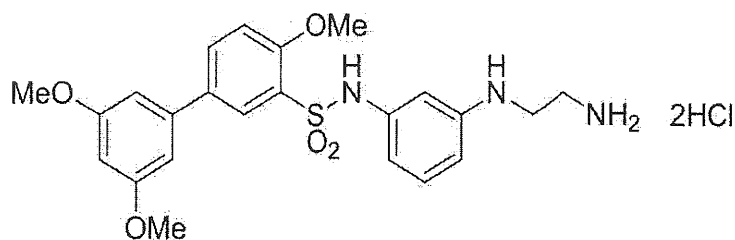
(2) To a solution of tert-butyl (2-((benzyl)(3-(2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (115 mg) in methanol (2.0 mL) was added 10% palladium/carbon (57.5 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature for 2 hr. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-(2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (95.4 mg).
(3) To tert-butyl (2-((3-(2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (95.4 mg) was added 10% hydrogen chloride-methanol (2.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (80.5 mg).

[0124]

Reference Example 21

N-(3-((2-aminoethyl)amino)phenyl)-3',4,5'-trimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride

[0125]



25

[0126]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (118 mg) in DME (4.0 mL) were added 3,5-dimethoxyphenylboronic acid (43.7 mg), sodium carbonate (42.4 mg), water (0.200 mL) and tetrakis(triphenylphosphine)palladium (11.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated chloroform) to give tert-butyl (2-((benzyl)(3-(3',4,5'-trimethoxy-[1,1'-biphenyl]-3-

ylsulfonamide)phenyl)amino)ethyl)carbamate (72.6 mg).

(2) To a solution of tert-butyl (2-((benzyl)(3-(3',4,5'-trimethoxy-[1,1'-biphenyl]-3-

ylsulfonamide)phenyl)amino)ethyl)carbamate (72.6 mg) in

5 methanol (2.0 mL) was added 10% palladium/carbon (21.8 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature for 2 hr. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column

10 chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-(3',4,5'-trimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (59.4 mg).

(3) To tert-butyl (2-((3-(3',4,5'-trimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (59.4 mg) was

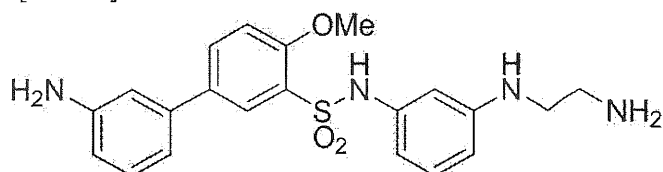
15 added 10% hydrogen chloride-methanol (2.0 mL), and the mixture was stirred at room temperature for 4 hr. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-3',4,5'-trimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (53.6 mg).

20 [0127]

Reference Example 22

3'-amino-N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-[1,1'-biphenyl]-3-sulfonamide

[0128]



25

[0129]

Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)amino)ethyl)carbamate (162 mg) in dioxane (3.0 mL) were added 3-(N-Boc-amino)phenylboronic acid pinacol ester (122 mg), sodium carbonate (58.3 mg), water (0.275 mL) and bis(diphenylphosphino)ferrocenepalladium dichloride dichloromethane complex (11.4 mg), and the mixture was heated under reflux overnight. The reaction mixture was

35 filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2). To a solution of the obtained solid (171 mg) in 10% hydrogen chloride-methanol (2.0 mL) was added 10% palladium/carbon

40 (51.3 mg), and the mixture was stirred under a hydrogen

atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: ammonia-saturated

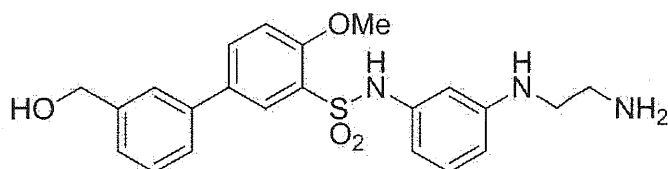
- 5 chloroform/methanol=10/1) to give 3'-amino-N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-[1,1'-biphenyl]-3-sulfonamide (81.5 mg).

[0130]

Reference Example 23

- 10 N-(3-((2-aminoethyl)amino)phenyl)-3'-(hydroxymethyl)-4-methoxy-[1,1'-biphenyl]-3-sulfonamide

[0131]



[0132]

- 15 (1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl) (3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (118 mg) in dioxane (4.0 mL) were added 2-(3-hydroxymethylphenyl)boronic acid pinacol ester (65.5 mg),
- 20 sodium carbonate (42.2 mg), water (0.200 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (8.2 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The
- 25 obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2→2/1) to give tert-butyl (2-((benzyl) (3-(3'-(hydroxymethyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (72.6 mg).
- 30 (2) To a solution of tert-butyl (2-((benzyl) (3-(3'-(hydroxymethyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (72.6 mg) in 10% hydrogen chloride-methanol (2.0 mL) was added 10% palladium/carbon (36.3 mg), and the mixture was stirred under
- 35 a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: ammonia-saturated chloroform/methanol=10/1) to give N-(3-((2-
- 40 aminoethyl)amino)phenyl)-3'-(hydroxymethyl)-4-methoxy-[1,1'-

biphenyl]-3-sulfonamide (51.0 mg).

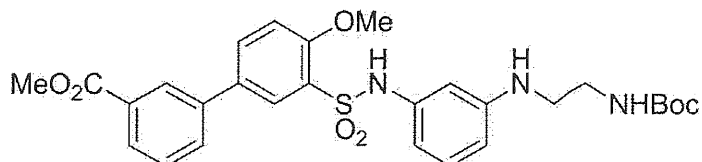
[0133]

Reference Example 24

methyl 3'-(N-(3-((2-((tert-

5 butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-
[1,1'-biphenyl]-3-carboxylate

[0134]



[0135]

10 (1) Under an argon atmosphere, to a solution of tert-butyl (2-
(benzyl)(3-(5-bromo-2-
methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (2.36 g)
in dioxane (20.0 mL) were added 3-methoxycarbonylphenylboronic
acid pinacol ester (1.26 g), sodium carbonate (828 mg), water
15 (4.0 mL) and bis(diphenylphosphino)ferrocenepalladium
dichloride dichloromethane complex (163 mg), and the mixture
was heated under reflux overnight. The reaction mixture was
filtered through celite, and the filtrate was concentrated.
The obtained residue was purified by silica gel column
20 chromatography (eluate: ethyl acetate/hexane=1/2) to give
methyl 3'-(N-(3-((benzyl)(2-((tert-
butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-
[1,1'-biphenyl]-3-carboxylate (2.25 g).

(2) To a suspension of methyl 3'-(N-(3-((benzyl)(2-((tert-
25 butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-
[1,1'-biphenyl]-3-carboxylate (2.25 g) in methanol (20 mL) was
added 10% palladium/carbon (675 mg), and the mixture was
stirred under a hydrogen atmosphere at room temperature for 5
hr. The reaction mixture was filtered through celite, and the
30 filtrate was concentrated. The obtained residue was purified
by amine silica gel column chromatography (eluate: ammonia-
saturated chloroform/methanol=10/1) to give methyl 3'-(N-(3-
((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-
4'-methoxy-[1,1'-biphenyl]-3-carboxylate (1.94 g).

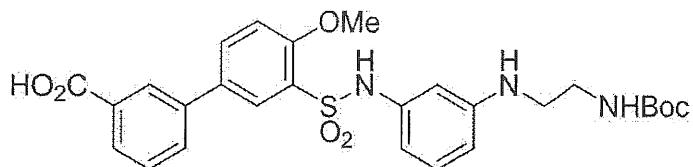
35 [0136]

Reference Example 25

3'-(N-(3-((2-((tert-

butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-
[1,1'-biphenyl]-3-carboxylic acid

40 [0137]



[0138]

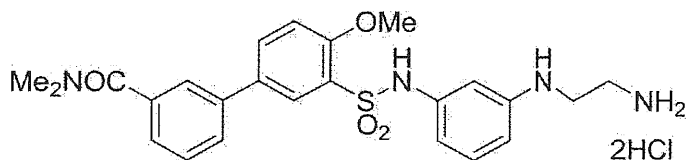
To a suspension of methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-
 5 [1,1'-biphenyl]-3-carboxylate (1.94 g) in methanol (30 mL) was added 1N aqueous sodium hydroxide solution (10.5 mL), and the mixture was stirred at 60°C for 3 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was extracted with a mixed solvent of chloroform and methanol (2/1). The
 10 organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated to give 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylic acid (1.77 g).

[0139]

15 Reference Example 26

3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride

[0140]



20 [0141]

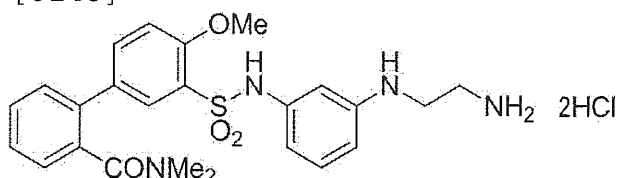
Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylic acid (1.77 g) in dichloromethane (30 mL) were added dimethylamine hydrochloride
 25 (533 mg), triethylamine (1.59 mL) and BOP reagent (1.59 g), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium
 30 sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/chloroform =4/1). To the obtained purified product was added 10% hydrogen chloride-methanol solution (10 mL), and the mixture was stirred at 50°C
 35 for 2 hr. The reaction mixture was concentrated, and the obtained residue was purified by amine silica gel column chromatography (eluate: ammonia-saturated

chloroform/methanol=1/0→100/1). To the obtained purified product was added 10% hydrogen chloride-methanol solution (5.0 mL), and the reaction mixture was concentrated to give 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (1.31 g).

[0142]

Reference Example 27

3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-2-carboxamide dihydrochloride
[0143]



[0144]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenyl)sulfonamido)phenyl)amino)ethyl)carbamate (118 mg) in dioxane (4.0 mL) were added 2-ethoxycarbonylphenylboronic acid pinacol ester (66.3 mg), sodium carbonate (42.4 mg), water (0.200 mL) and bis(diphenylphosphino)ferrocenepalladium dichloride dichloromethane complex (8.2 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2→1/1) to give ethyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-2-carboxylate (125 mg).

(2) To a suspension of ethyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-2-carboxylate (125 mg) in methanol (4.0 mL) was added 10% palladium/carbon (37.5 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature for 3 hr. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give ethyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-2-carboxylate (95.4 mg).

(3) To a suspension of ethyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-

[1,1'-biphenyl]-2-carboxylate (95.4 mg) in methanol (3.0 mL) was added 1N aqueous sodium hydroxide solution (0.510 mL), and the mixture was stirred at 60°C for 2 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was
5 extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, the filtrate was concentrated to give 3'-(N-(3-((2-((tert-

10 (4) Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-2-carboxylic acid (100 mg) in dichloromethane (2.0 mL) were added dimethylamine hydrochloride (30.2 mg), triethylamine (0.085 mL) and BOP
15 reagent (88.5 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was
20 concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) to give tert-butyl (2-((3-(2'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (65.0 mg).

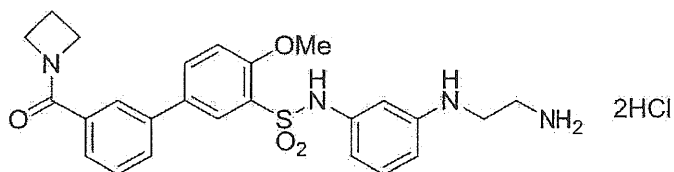
25 (5) To tert-butyl (2-((3-(2'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (65.0 mg) was added 10% hydrogen chloride-methanol solution (2.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give 3'-(N-(3-((2-
30 aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-2-carboxamide dihydrochloride (61.9 mg).

[0145]

Reference Example 28

N-(3-((2-aminoethyl)amino)phenyl)-3'-(azetidine-1-carbonyl)-4-
35 methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride

[0146]



[0147]

(1) Under an argon atmosphere, to a suspension of 3'-(N-(3-
40 ((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-

4'-methoxy-[1,1'-biphenyl]-3-carboxylic acid (130 mg) in dichloromethane (2.0 mL) were added azetidine hydrochloride (24.3 mg), triethylamine (0.074 mL) and BOP reagent (115 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl) carbamate (109 mg).

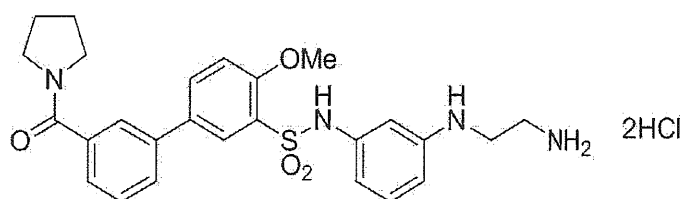
(2) To tert-butyl (2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl) carbamate (109 mg) was added 10% hydrogen chloride-methanol solution (2.0 mL), and the mixture was stirred at room temperature overnight. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (96.1 mg).

[0148]

Reference Example 29

N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride

[0149]



[0150]

(1) Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylic acid (130 mg) in dichloromethane (2.0 mL) were added pyrrolidine (0.020 mL), triethylamine (0.036 mL) and BOP reagent (115 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-(4-methoxy-

3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (154 mg).

(2) To tert-butyl (2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-

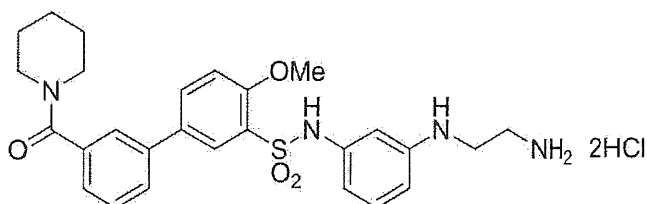
5 ylsulfonamide)phenyl)amino)ethyl)carbamate (154 mg) was added 10% hydrogen chloride-methanol solution (3.0 mL), and the mixture was stirred at 50°C overnight. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (125 mg).

[0151]

Reference Example 30

N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride

15 [0152]



[0153]

(1) Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylic acid (130 mg) in dichloromethane (2.0 mL) were added piperidine (0.024 mL), triethylamine (0.036 mL) and BOP reagent (115 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-(4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (129 mg).

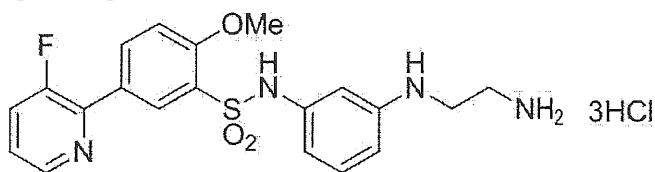
(2) To tert-butyl (2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (129 mg) was added 10% hydrogen chloride-methanol solution (3.0 mL), and the mixture was stirred at 50°C overnight. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (112 mg).

[0154]

Reference Example 31

N-(3-((2-aminoethyl)amino)phenyl)-5-(3-fluoropyridin-2-yl)-2-methoxybenzenesulfonamide trihydrochloride

5 [0155]



[0156]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl) (3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)amino)ethyl)carbamate (590 mg) in DMSO (2.0 mL) were added bis(pinacolato)diboron (305 mg), potassium acetate (294 mg) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (40.8 mg), and the mixture was stirred at 80°C overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was filtered through celite, and the filtrate was extracted with ethyl acetate. The organic layer was washed with water, dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/1). Under an argon atmosphere, to a solution of the obtained purified product (250 mg) in dioxane (4.0 mL) were added 2-bromo-3-fluoropyridine (0.056 mL), sodium carbonate (82.7 mg), water (0.390 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (16.3 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ethyl acetate/hexane=1/1) to give tert-butyl (2-((benzyl) (3-(5-(3-fluoropyridin-2-yl)-2-methoxyphenylsulfonamide)phenyl)amino)ethyl)carbamate (51.2 mg).

(2) To a suspension of tert-butyl (2-((benzyl) (3-(5-(3-fluoropyridin-2-yl)-2-methoxyphenylsulfonamide)phenyl)amino)ethyl)carbamate (51.2 mg) in methanol (2.0 mL) was added 10% palladium/carbon (15.4 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated.

The obtained residue was purified by amine silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) to give tert-butyl (2-((3-(5-(3-fluoropyridin-2-yl)-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (18.8 mg).

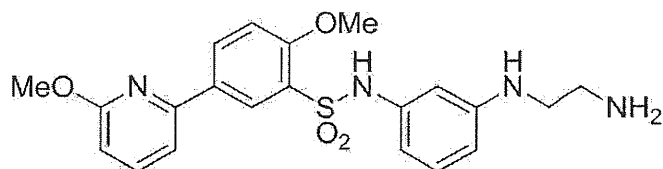
(3) To tert-butyl (2-((3-(5-(3-fluoropyridin-2-yl)-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (18.8 mg) was added 10% hydrogen chloride-methanol solution (1.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-5-(3-fluoropyridin-2-yl)-2-methoxybenzenesulfonamide trihydrochloride (16.8 mg).

[0157]

Reference Example 32

N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(6-methoxypyridin-2-yl)benzenesulfonamide

[0158]



[0159]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl) (3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (118 mg) in dioxane (2.5 mL) were added 6-methoxy-2-pyridineboronic acid MIDA ester (52.8 mg), potassium phosphate (318 mg), water (0.500 mL), palladium acetate (2.2 mg) and 2-dicyclohexylphosphino-2',6'-dimethoxybiphenyl (Sphos) (8.2 mg), and the mixture was stirred at 60°C overnight. To this reaction mixture was added 1N aqueous sodium hydroxide solution, and the mixture was extracted with ethyl acetate. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia saturated-chloroform) to give tert-butyl (2-((benzyl) (3-(2-methoxy-5-(6-methoxypyridin-2-yl)phenyl)sulfonamide)phenyl)amino)ethyl)carbamate (72.7 mg). (2) To a solution of tert-butyl (2-((benzyl) (3-(2-methoxy-5-(6-methoxypyridin-2-yl)phenyl)sulfonamide)phenyl)amino)ethyl)carbamate (72.7 mg) in 10% hydrogen chloride-methanol (4.0 mL) was added 10% palladium/carbon (36.4 mg), and the mixture was stirred under

a hydrogen atmosphere at room temperature for 4 hr. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate:

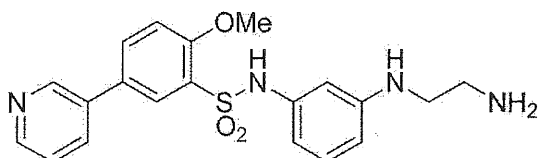
- 5 chloroform/methanol=10/1) to give N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(6-methoxypyridin-2-yl)benzenesulfonamide (54.9 mg).

[0160]

Reference Example 33

- 10 N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(pyridin-3-yl)benzenesulfonamide

[0161]



[0162]

- 15 (1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl) (3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (118 mg) in dioxane (4.0 mL) were added 3-pyridylboronic acid (29.5 mg), sodium carbonate (42.4 mg), water (0.200 mL) and
- 20 tetrakis(triphenylphosphine)palladium (11.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=3/2) to
- 25 give tert-butyl (2-((benzyl) (3-(2-methoxy-5-(pyridin-3-yl)phenyl)sulfonamide)phenyl)amino)ethyl)carbamate (46.0 mg).
- (2) To a solution of tert-butyl (2-((benzyl) (3-(2-methoxy-5-(pyridin-3-yl)phenyl)sulfonamide)phenyl)amino)ethyl)carbamate (46.0 mg) in 10% hydrogen chloride-methanol (2.0 mL) was added
- 30 10% palladium/carbon (23.0 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature for 3 hr. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate:
- 35 chloroform/methanol=10/1) to give N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(pyridin-3-yl)benzenesulfonamide (128 mg).

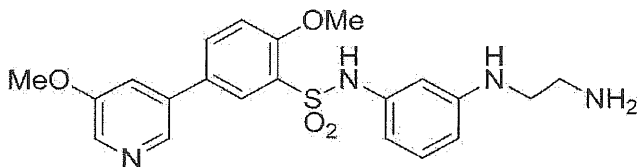
[0163]

Reference Example 34

- 40 N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(5-

methoxypyridin-3-yl)benzenesulfonamide

[0164]



[0165]

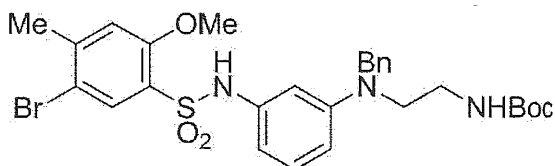
- 5 (1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl (3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)carbamate (118 mg) in dioxane (4.0 mL) were added 5-methoxy-3-pyridineboronic acid pinacol ester (65.8 mg), sodium carbonate (42.4 mg),
10 water (0.200 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (8.2 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column
15 chromatography (eluate: ethyl acetate/hexane=3/2→2/1) to give tert-butyl (2-((benzyl (3-(2-methoxy-5-(5-methoxypyridin-3-yl)phenyl)sulfonamide)phenyl)amino)ethyl)carbamate (108 mg).
(2) To a solution of tert-butyl (2-((benzyl (3-(2-methoxy-5-(5-methoxypyridin-3-yl)phenyl)sulfonamide)phenyl)amino)ethyl)carbamate (108 mg) in
20 10% hydrogen chloride-methanol (2.0 mL) was added 10% palladium/carbon (53.9 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate
25 was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: ammonia-saturated chloroform/methanol=10/1) to give N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(5-methoxypyridin-3-yl)benzenesulfonamide (52.1 mg).

30 [0166]

Reference Example 35

tert-butyl (2-((benzyl (3-(5-bromo-2-methoxy-4-methylphenyl)sulfonamide)phenyl)amino)ethyl)carbamate

[0167]



35

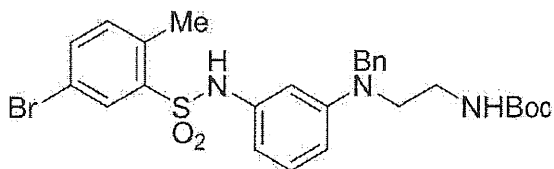
[0168]

Under an argon atmosphere, to a solution of tert-butyl

(2-((3-aminophenyl)(benzyl)amino)ethyl)carbamate (688 mg) in dichloromethane (10.0 mL) were added pyridine (0.164 mL) and 5-bromo-2-methoxy-4-methylbenzenesulfonyl chloride (581 mg), and the mixture was stirred at room temperature overnight. To
5 this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column
10 chromatography (eluate: ethyl acetate/hexane=1/2→2/3) to give tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxy-4-methylphenylsulfonamide)phenyl)amino)ethyl)carbamate (849 mg).
[0169]

Reference Example 36

15 tert-butyl (2-((benzyl)(3-(5-bromo-2-methylphenylsulfonamide)phenyl)amino)ethyl)carbamate
[0170]

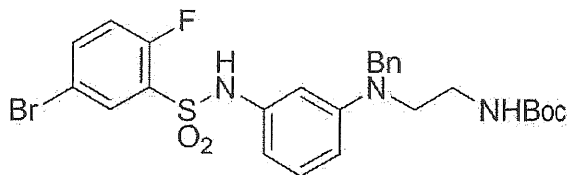


[0171]

20 Under an argon atmosphere, to a solution of tert-butyl (2-((3-aminophenyl)(benzyl)amino)ethyl)carbamate (275 mg) in dichloromethane (5.0 mL) were added pyridine (0.066 mL) and 5-bromo-2-methylbenzenesulfonyl chloride (200 mg), and the mixture was stirred at room temperature overnight. To this
25 reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography
30 (eluate: ethyl acetate/hexane=1/3) to give tert-butyl (2-((benzyl)(3-(5-bromo-2-methylphenylsulfonamide)phenyl)amino)ethyl)carbamate (386 mg).
[0172]

Reference Example 37

35 tert-butyl (2-((benzyl)(3-(5-bromo-2-fluorophenylsulfonamide)phenyl)amino)ethyl)carbamate
[0173]



[0174]

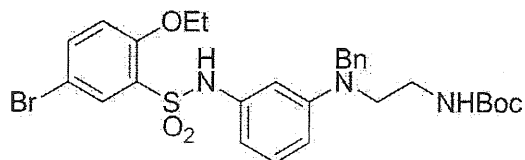
Under an argon atmosphere, to a solution of tert-butyl (2-((3-aminophenyl) (benzyl) amino) ethyl) carbamate (271 mg) in dichloromethane (6.0 mL) were added pyridine (0.064 mL) and 5-bromo-2-fluorobenzenesulfonyl chloride (200 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/3) to give tert-butyl (2-((benzyl) (3-(5-bromo-2-fluorophenylsulfonamide) phenyl) amino) ethyl) carbamate (405 mg).

[0175]

Reference Example 38

tert-butyl (2-((benzyl) (3-(5-bromo-2-ethoxyphenylsulfonamide) phenyl) amino) ethyl) carbamate

[0176]



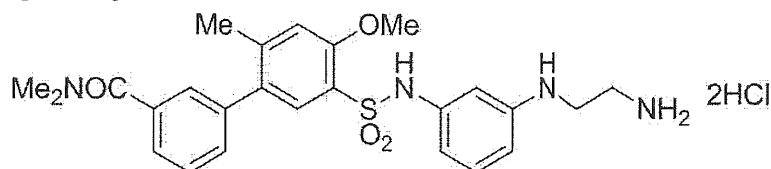
[0177]

Under an argon atmosphere, to a solution of tert-butyl (2-((3-aminophenyl) (benzyl) amino) ethyl) carbamate (249 mg) in dichloromethane (6.0 mL) were added pyridine (0.060 mL) and 5-bromo-2-ethoxybenzenesulfonyl chloride (200 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2) to give tert-butyl (2-((benzyl) (3-(5-bromo-2-ethoxyphenylsulfonamide) phenyl) amino) ethyl) carbamate (312 mg).

[0178]

Reference Example 39

5'- (N- (3- ((2-aminoethyl) amino) phenyl) sulfamoyl) -4'-methoxy-
N, N, 2'-trimethyl- [1, 1'-biphenyl]-3-carboxamide dihydrochloride
[0179]



5 [0180]

- (1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl) (3-(5-bromo-2-methoxy-4-methylphenyl)sulfonamide)phenyl) amino) ethyl) carbamate (424 mg) in dioxane (7.0 mL) were added 3-methoxycarbonylphenylboronic acid pinacol ester (220 mg), sodium carbonate (148 mg), water (0.700 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (28.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated.
- 15 The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=2/3) to give methyl 5'- (N- (3- ((benzyl) (2-((tert-butoxycarbonyl) amino) ethyl) amino) phenyl) sulfamoyl) -4'-methoxy-2'-methyl- [1, 1'-biphenyl]-3-carboxylate (461 mg).
- 20 (2) To a suspension of methyl 5'- (N- (3- ((benzyl) (2-((tert-butoxycarbonyl) amino) ethyl) amino) phenyl) sulfamoyl) -4'-methoxy-2'-methyl- [1, 1'-biphenyl]-3-carboxylate (461 mg) in methanol (7.0 mL) was added 10% palladium/carbon (138 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give methyl 5'- (N- (3- ((2-((tert-butoxycarbonyl) amino) ethyl) amino) phenyl) sulfamoyl) -4'-methoxy-2'-methyl- [1, 1'-biphenyl]-3-carboxylate (394 mg).
- 30 (3) To a suspension of methyl 5'- (N- (3- ((2-((tert-butoxycarbonyl) amino) ethyl) amino) phenyl) sulfamoyl) -4'-methoxy-2'-methyl- [1, 1'-biphenyl]-3-carboxylate (394 mg) in methanol (7.0 mL) was added 1N aqueous sodium hydroxide solution (2.07 mL), and the mixture was stirred at 60°C for 3 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, the filtrate was
- 40 concentrated to give 5'- (N- (3- ((2-((tert-

butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-2'-methyl-[1,1'-biphenyl]-3-carboxylic acid (383 mg).

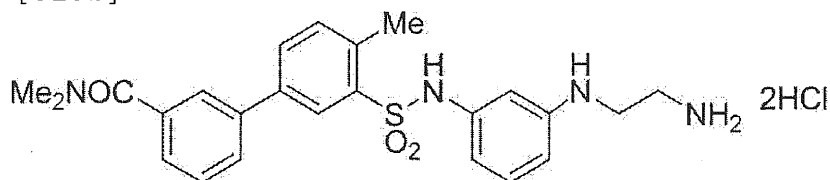
(4) Under an argon atmosphere, to a suspension of 5'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-2'-methyl-[1,1'-biphenyl]-3-carboxylic acid (383 mg) in dichloromethane (10 mL) were added dimethylamine hydrochloride (113 mg), triethylamine (0.337 mL) and BOP reagent (336 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→1/0) to give tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-6-methyl-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (299 mg).

(5) To tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-6-methyl-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (299 mg) was added 10% hydrogen chloride-methanol solution (4.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give 5'-(N-(3-(2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (290 mg).

[0181]

Reference Example 40

3'-(N-(3-(2-aminoethyl)amino)phenyl)sulfamoyl)-N,N,4'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride
[0182]



[0183]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl) (3-(5-bromo-2-methoxy-4-methylphenylsulfonamide)phenyl)amino)ethyl)carbamate (386 mg) in dioxane (7.0 mL) were added 3-methoxycarbonylphenylboronic acid pinacol ester (210 mg), sodium carbonate (142 mg), water (0.670 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (27.8 mg), and the mixture

was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=2/5) to give

5 methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methyl-[1,1'-biphenyl]-3-carboxylate (419 mg).

(2) To a suspension of methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methyl-

10 [1,1'-biphenyl]-3-carboxylate (419 mg) in methanol (7.0 mL) was added 10% palladium/carbon (126 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was

15 purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methyl-[1,1'-biphenyl]-3-carboxylate (353 mg).

(3) To a suspension of methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methyl-

20 [1,1'-biphenyl]-3-carboxylate (353 mg) in methanol (7.0 mL) was added 1N aqueous sodium hydroxide solution (1.96 mL), and the mixture was stirred at 60°C for 3 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was

25 extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated to give 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methyl-[1,1'-biphenyl]-3-carboxylic acid (341 mg).

(4) Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methyl-[1,1'-biphenyl]-3-carboxylic acid (341 mg) in

30 dichloromethane (10 mL) were added dimethylamine hydrochloride (106 mg), triethylamine (0.318 mL) and BOP reagent (318 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium

35 hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The

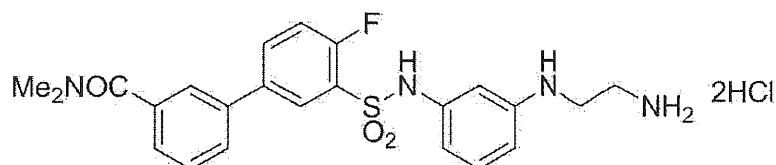
40 obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→1/0) to give tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-methyl-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (255

mg).

(5) To tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-methyl-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (255 mg) was added 10% hydrogen chloride-methanol solution (4.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-N,N,4'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (243 mg).
[0184]

10 Reference Example 41

3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-fluoro-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride
[0185]



15 [0186]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-fluorophenylsulfonamide)phenyl)amino)ethyl)carbamate (405 mg) in dioxane (7.0 mL) were added 3-methoxycarbonylphenylboronic acid pinacol ester (220 mg), sodium carbonate (148 mg), water (0.700 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (28.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated.
25 The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2) to give methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-fluoro-[1,1'-biphenyl]-3-carboxylate (458 mg).
30 (2) To a suspension of methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-fluoro-[1,1'-biphenyl]-3-carboxylate (458 mg) in methanol (7.0 mL) was added 10% palladium/carbon (137 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature
35 overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-fluoro-
40 [1,1'-biphenyl]-3-carboxylate (387 mg).

(3) To a suspension of methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-fluoro-[1,1'-biphenyl]-3-carboxylate (387 mg) in methanol (7.0 mL) was added 1N aqueous sodium hydroxide solution (2.13 mL), and the mixture was stirred at 60°C for 3 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated to give 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-fluoro-[1,1'-biphenyl]-3-carboxylic acid (346 mg).

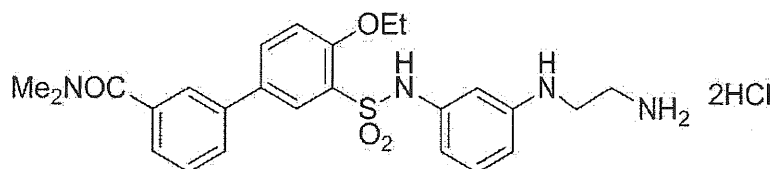
(4) Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-fluoro-[1,1'-biphenyl]-3-carboxylic acid (346 mg) in dichloromethane (10 mL) were added dimethylamine hydrochloride (116 mg), triethylamine (0.347 mL) and BOP reagent (345 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-fluoro-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (291 mg).

(5) To tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-fluoro-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (291 mg) was added 10% hydrogen chloride-methanol solution (4.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-fluoro-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (283 mg).

[0187]

Reference Example 42

3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride [0188]



[0189]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-

((benzyl)(3-(5-bromo-2-ethoxyphenylsulfonamide)phenyl)amino)ethyl)carbamate (312 mg) in dioxane (5.0 mL) were added 3-methoxycarbonylphenylboronic acid pinacol ester (163 mg), sodium carbonate (109 mg), water (0.520 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (21.2 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2→1/1) to give methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-[1,1'-biphenyl]-3-carboxylate (270 mg).

(2) To a suspension of methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-[1,1'-biphenyl]-3-carboxylate (270 mg) in methanol (7.0 mL) was added 10% palladium/carbon (81.0 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-[1,1'-biphenyl]-3-carboxylate (222 mg).

(3) To a suspension of methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-[1,1'-biphenyl]-3-carboxylate (222 mg) in methanol (7.0 mL) was added 1N aqueous sodium hydroxide solution (1.17 mL), and the mixture was stirred at 60°C for 2 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated to give 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-[1,1'-biphenyl]-3-carboxylic acid (215 mg).

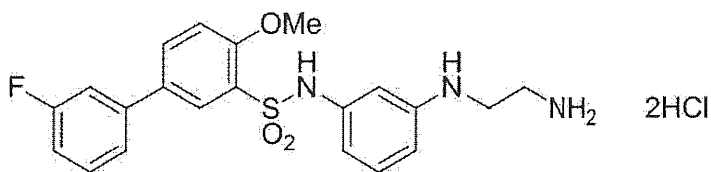
(4) Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-[1,1'-biphenyl]-3-carboxylic acid (215 mg) in dichloromethane (8.0 mL) were added dimethylamine hydrochloride (63.6 mg), triethylamine (0.190 mL) and BOP reagent (190 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the

mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-ethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (191 mg).

(5) To tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-ethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (191 mg) was added 10% hydrogen chloride-methanol solution (4.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (176 mg).
[0190]

15 Reference Example 43

N-(3-((2-aminoethyl)amino)phenyl)-3'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride
[0191]



20 [0192]

(1) Under an argon atmosphere, to a solution of tert-butyl (2-((3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (118 mg) in n-propanol (4.0 mL) were added potassium 3-fluorophenyltrifluoroborate (48.5 mg), triethylamine (0.084 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (4.9 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated chloroform) to give tert-butyl (2-((3-(3'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (67.5 mg).

(2) To a solution of tert-butyl (2-((3-(3'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (67.5 mg) in methanol (2.0 mL) was added 10% palladium/carbon (20.3 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite,

and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give tert-butyl (2-((3-(3'-fluoro-4-methoxy-[1,1'-biphenyl]-3-

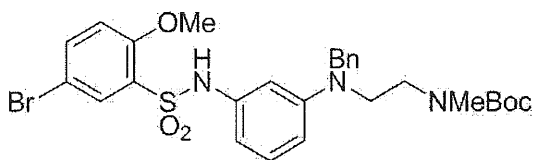
5 ylsulfonamide)phenyl)amino)ethyl)carbamate (28.5 mg).

(3) To tert-butyl (2-((3-(3'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (28.5 mg) was added 10% hydrogen chloride-methanol solution (2.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture
10 was concentrated to give N-(3-((2-aminoethyl)amino)phenyl)-3'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (26.7 mg).

[0193]

Reference Example 44

15 tert-butyl (2-((benzyl) (3-(5-bromo-2-methoxyphenylsulfonamide)phenyl)amino)ethyl) (methyl) carbamate
[0194]



[0195]

20 (1) Under an argon atmosphere, to a solution of tert-butyl (2-((3-nitrophenyl)amino)ethyl)carbamate (4.46 g) in dichloromethane (30 mL) were added 50 wt% aqueous sodium hydroxide solution (10 mL), benzyl bromide (2.84 mL) and tetrabutylammonium iodide (587 mg), 20 - 30°C for 48 hr. To
25 this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column
30 chromatography (eluate: ethyl acetate/hexane=1/6→1/4) to give tert-butyl (2-((benzyl) (3-nitrophenyl)amino)ethyl)carbamate (4.62 g).

(2) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl) (3-nitrophenyl)amino)ethyl)carbamate (297 mg) in DMF
35 (3.0 mL) were added potassium tert-butoxide (117 mg) and methyl iodide (0.065 mL), and the mixture was stirred at room temperature for 2 hr. To this reaction mixture was added saturated brine, the mixture was extracted with ethyl acetate, and the organic layer was washed with water. The organic layer
40 was dried over sodium sulfate, filtered, and the filtrate was

concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ethyl acetate/hexane=1/5) to give tert-butyl (2-((benzyl)(3-nitrophenyl)amino)ethyl)(methyl)carbamate (218 mg).

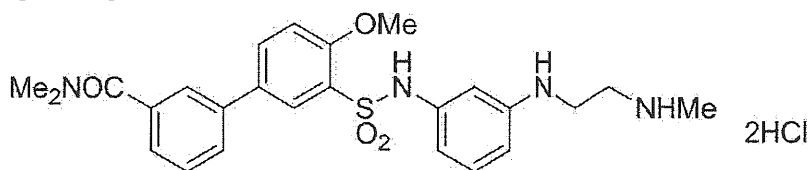
- 5 (3) Under an argon atmosphere, to a mixed suspension of (2-((benzyl)(3-nitrophenyl)amino)ethyl)(methyl)carbamate (218 mg) in ethanol (2.5 mL) and water (1.0 mL) were added ammonium chloride (305 mg) and iron powder (221 mg), and the mixture was heated under reflux for 2 hr. The reaction mixture was
10 filtered through celite, the filtrate was concentrated, and the obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give tert-butyl (2-((3-aminophenyl)(benzyl)amino)ethyl)(methyl)carbamate (200 mg).
- 15 (4) Under an argon atmosphere, to a solution of tert-butyl (2-((3-aminophenyl)(benzyl)amino)ethyl)(methyl)carbamate (200 mg) in dichloromethane (6.0 mL) were added pyridine (0.050 mL) and 5-bromo-2-methoxybenzenesulfonyl chloride (160 mg), and the mixture was stirred at room temperature overnight. To this
20 reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography
25 (eluate: ethyl acetate/hexane=1/2) to give tert-butyl (2-((benzyl)(3-(5-bromo-2-methoxyphenyl)sulfonamide)phenyl)amino)ethyl)(methyl)carbamate (260 mg).

[0196]

30 Reference Example 45

4'-methoxy-N,N-dimethyl-3'-(N-(3-((2-methylamino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxamide dihydrochloride

[0197]



[0198]

- (1) Under an argon atmosphere, to a solution of tert-butyl (2-((benzyl)(3-(5-bromo-2-ethoxyphenyl)sulfonamide)phenyl)amino)ethyl)(methyl)carbamate
40 (260 mg) in dioxane (4.0 mL) were added 3-

methoxycarbonylphenylboronic acid pinacol ester (136 mg), sodium carbonate (91.2 mg), water (0.430 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (17.6 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/2→3/5) to give methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)(methyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylate (246 mg).

(2) To a suspension of methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)(methyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-[1,1'-biphenyl]-3-carboxylate (246 mg) in methanol (5.0 mL) was added 10% palladium/carbon (73.8 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)(methyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylate (217 mg).

(3) To a suspension of methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)(methyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylate (217 mg) in methanol (7.0 mL) was added 1N aqueous sodium hydroxide solution (1.14 mL), and the mixture was stirred at 60°C for 2 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated to give 3'-(N-(3-((2-((tert-butoxycarbonyl)(methyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylic acid (208 mg).

(4) Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)(methyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylic acid (208 mg) in dichloromethane (5.0 mL) were added dimethylamine hydrochloride (61.1 mg), triethylamine (0.183 mL) and BOP reagent (181 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was

dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate) to give tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-

5 biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl) (methyl) carbamate (160 mg).

(5) To tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-

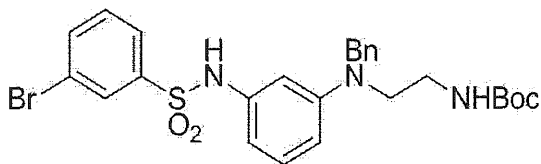
ylsulfonamide)phenyl)amino)ethyl) (methyl) carbamate (160 mg)
10 was added 10% hydrogen chloride-methanol solution (4.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give 4'-methoxy-N,N-dimethyl-3'-(N-(3-((2-methylamino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxamide dihydrochloride (150 mg).

15 [0199]

Reference Example 46

tert-butyl (2-((benzyl)(3-(3-bromophenylsulfonamide)phenyl)amino)ethyl) carbamate

[0200]



[0201]

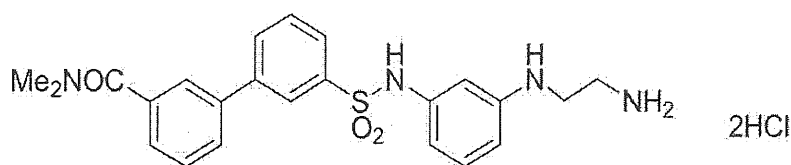
Under an argon atmosphere, to a solution of tert-butyl (2-((3-aminophenyl)(benzyl)amino)ethyl) carbamate (300 mg) in dichloromethane (8.0 mL) were added pyridine (0.078 mL) and 3-bromobenzenesulfonyl chloride (0.116 mL), and the mixture was
25 stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the
30 filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/3) to give tert-butyl (2-((benzyl)(3-(3-bromophenylsulfonamide)phenyl)amino)ethyl) carbamate (423 mg).

[0202]

35 Reference Example 47

3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride

[0203]



[0204]

(1) To a solution of tert-butyl (2-((benzyl)(3-(3-bromophenylsulfonamide)phenyl)amino)ethyl)carbamate (423 mg) in dioxane (7.0 mL) were added 3-methoxycarbonylphenylboronic acid pinacol ester (239 mg), sodium carbonate (160 mg), water (0.754 mL) and bis(diphenylphosphino)ferrocene palladium dichloride dichloromethane complex (30.8 mg), and the mixture was heated under reflux overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=1/3→1/2) to give methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylate (297 mg).

(2) To a suspension of methyl 3'-(N-(3-((benzyl)(2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylate (297 mg) in methanol (7.0 mL) was added 10% palladium/carbon (100 mg), and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The reaction mixture was filtered through celite, and the filtrate was concentrated. The obtained residue was purified by amine silica gel column chromatography (eluate: chloroform/methanol=10/1) to give methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylate (238 mg).

(3) To a suspension of methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylate (238 mg) in methanol (5.0 mL) was added 1N aqueous sodium hydroxide solution (1.36 mL), and the mixture was stirred at 60°C for 2 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated to give 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (222 mg).

(4) Under an argon atmosphere, to a suspension of 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (222 mg) in dichloromethane

(6.0 mL) were added dimethylamine hydrochloride (70.1 mg), triethylamine (0.210 mL) and BOP reagent (208 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=4/1) to give tert-butyl (2-((3-
5 (3'-(dimethylcarbamoyl)-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (185 mg).

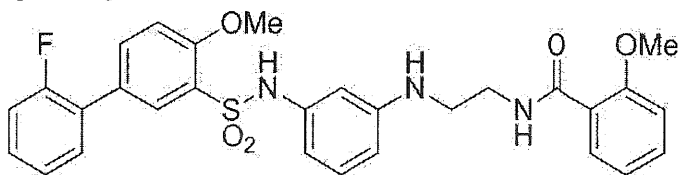
(5) To tert-butyl (2-((3-(3'-(dimethylcarbamoyl)-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (185 mg) was added 10% hydrogen chloride-methanol solution (4.0 mL),
15 and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (142 mg).

[0205]

20 Example 1

N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

[0206]



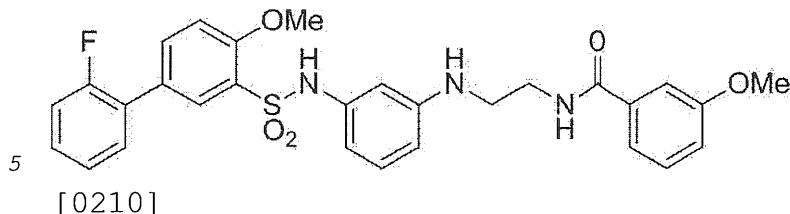
25 [0207]

Under an argon atmosphere, to a suspension of N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) in dichloromethane (1.0 mL) were added o-anisic acid (6.1 mg), triethylamine
30 (0.018 mL) and BOP reagent (19.5 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the
35 filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=3/2→3/1) to give N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (22.0 mg).

40 [0208]

Example 2

N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide
[0209]

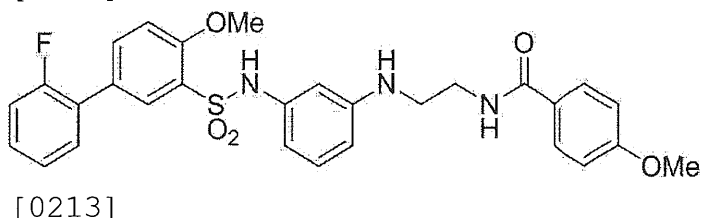


In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with m-anisic acid (6.1 mg) to give N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (21.9 mg).

[0211]

Example 3

15 N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-4-methoxybenzamide
[0212]

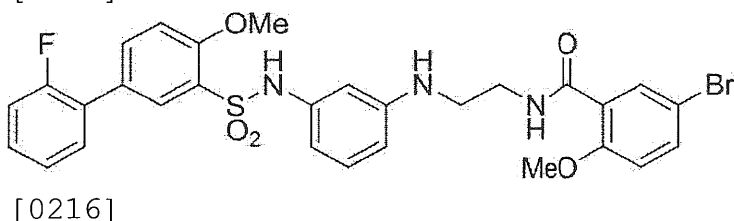


20 In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with p-anisic acid (6.1 mg) to give N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-4-methoxybenzamide (16.8 mg).

[0214]

Example 4

5-bromo-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide
30 [0215]



In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with 5-bromo-2-methoxybenzoic acid (6.1 mg) to give 5-bromo-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (16.8 mg).

3-sulfonamide dihydrochloride (19.5 mg) was amidated with 5-bromo-2-methoxybenzoic acid (9.2 mg) to give 5-bromo-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (24.8 mg).

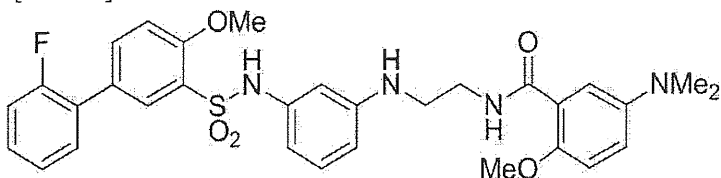
5 For purification, silica gel column chromatography (eluate: chloroform/ethyl acetate=1/0→3/1) was used.

[0217]

Example 5

5-(dimethylamino)-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

[0218]



[0219]

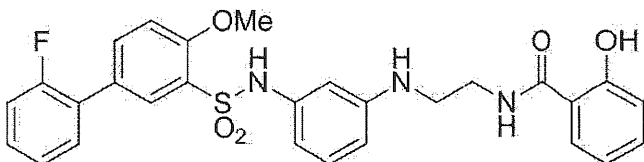
15 In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with 5-dimethylamino-2-methoxybenzoic acid (7.8 mg) to give 5-(dimethylamino)-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (21.3 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0220]

Example 6

25 N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-hydroxybenzamide

[0221]



[0222]

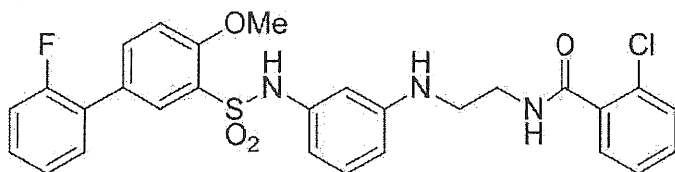
30 In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with salicylic acid (5.5 mg) to give N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-hydroxybenzamide (18.8 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) was used.

[0223]

Example 7

2-chloro-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide

5 [0224]



[0225]

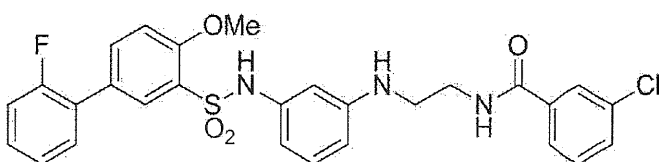
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with 2-chlorobenzoic acid (6.3 mg) to give 2-chloro-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide (20.3 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=2/1) was used.

[0226]

Example 8

3-chloro-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide

20 [0227]



[0228]

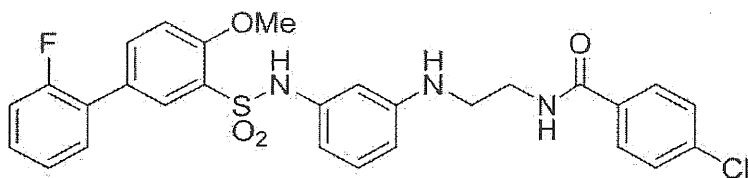
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with 3-chlorobenzoic acid (6.3 mg) to give 3-chloro-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide (15.0 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0229]

Example 9

4-chloro-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide

35 [0230]



[0231]

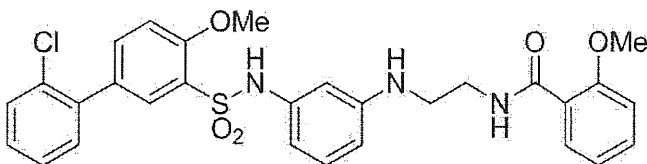
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with 4-chlorobenzoic acid (6.3 mg) to give 4-chloro-N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide (21.3 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0232]

Example 10

N-(2-((3-(2'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

[0233]



[0234]

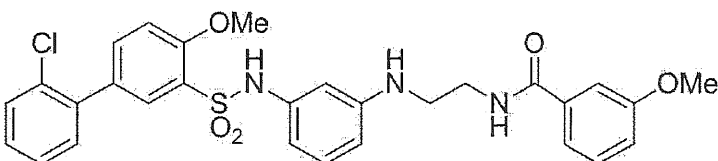
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-chloro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (23.0 mg) was amidated with o-anisic acid (7.0 mg) to give N-(2-((3-(2'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (20.2 mg). For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=40/1) was used.

[0235]

Example 11

N-(2-((3-(2'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide

[0236]



[0237]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-chloro-4-methoxy-[1,1'-biphenyl]-

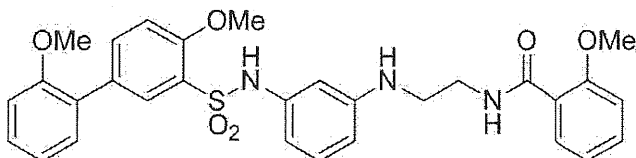
3-sulfonamide dihydrochloride (26.9 mg) was amidated with m-anisic acid (8.1 mg) to give N-(2-((3-(2'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (21.5 mg). For purification, silica gel
5 column chromatography (eluate: ethyl acetate) was used.

[0238]

Example 12

N-(2-((3-(2',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

10 [0239]



[0240]

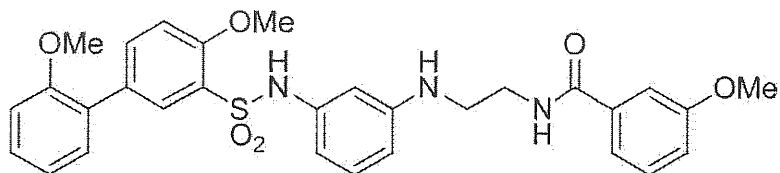
To a solution of tert-butyl (2-((3-(2',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl)amino)ethyl)carbamate (136 mg) in
15 dichloromethane (3.0 mL) was added trifluoroacetic acid (TFA) (0.500 mL), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture
20 was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. Under an argon atmosphere, to a solution of the obtained residue in dichloromethane (2.0 mL) were added o-anisic acid (18.3 mg), triethylamine (0.020 mL) and BOP
25 reagent (61.9 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was
30 concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=2/1→1/0) to give N-(2-((3-(2',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (20.3 mg).

[0241]

35 Example 13

N-(2-((3-(2',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide

[0242]

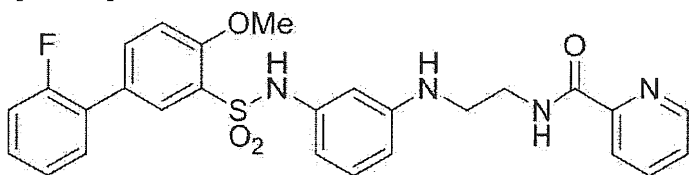


[0243]

In the same manner as in Example 12, tert-butyl (2-((3-(2',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl)amino)ethyl)carbamate (136 mg) was deprotected with TFA, and amidated with m-anisic acid (18.3 mg) to give N-(2-((3-(2',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (22.3 mg).
[0244]

10 Example 14

N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)picolinamide
[0245]

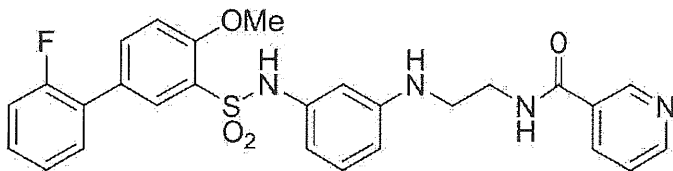


15 [0246]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with picolinic acid (4.9 mg) to give N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)picoline amide (19.2 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=2/1→1/0) was used.
[0247]

Example 15

25 N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)nicotineamide
[0248]



[0249]

30 In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with nicotinic acid (4.9 mg) to give N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-

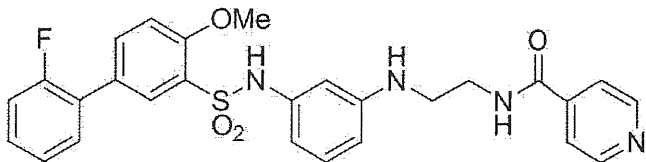
ylsulfoneamide)phenyl)amino)ethyl)nicotinamide (20.2 mg). For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=40/1) was used.

[0250]

5 Example 16

N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)isonicotinamide

[0251]



10 [0252]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with isonicotinic acid (4.9 mg) to give N-(2-((3-(2'-fluoro-4-methoxy-[1,1'-biphenyl]-3-

15 methoxy-[1,1'-biphenyl]-3-

ylsulfonamide)phenyl)amino)ethyl)isonicotinamide (21.6 mg).

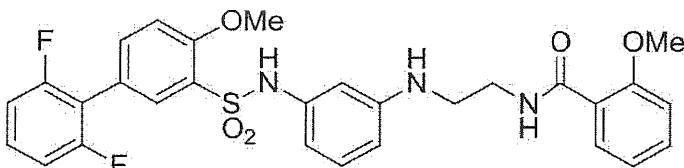
For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=40/1) was used.

[0253]

20 Example 17

N-(2-((3-(2',6'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

[0254]



25 [0255]

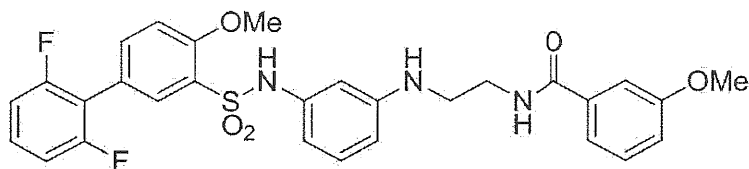
In the same manner as in Example 12, tert-butyl (2-((3-(2',6'-difluoro-4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)(4-methoxybenzyl)amino)ethyl)carbamate (69.6 mg) was deprotected with TFA, and amidated with o-anisic acid (11.4 mg) to give N-(2-((3-(2',6'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (15.2 mg).

[0256]

Example 18

35 N-(2-((3-(2',6'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide

[0257]



[0258]

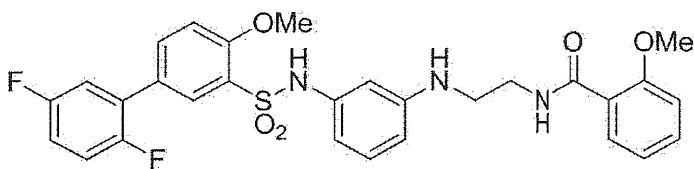
In the same manner as in Example 12, tert-butyl (2-((3-(2',6'-difluoro-4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino) ethyl) carbamate (69.6 mg) was deprotected with TFA, and amidated with m-anisic acid (11.4 mg) to give N-(2-((3-(2',6'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino) ethyl)-3-methoxybenzamide (13.0 mg).

[0259]

Example 19

N-(2-((3-(2',5'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino) ethyl)-2-methoxybenzamide

[0260]



15

[0261]

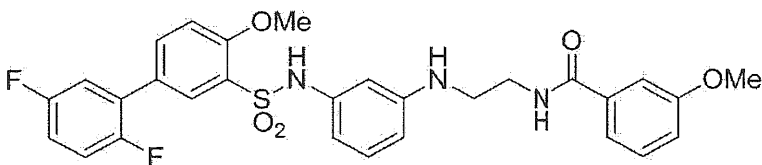
In the same manner as in Example 12, tert-butyl (2-((3-(2',5'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino) ethyl) carbamate (70.1 mg) was deprotected with TFA, and amidated with o-anisic acid (11.4 mg) to give N-(2-((3-(2',5'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino) ethyl)-2-methoxybenzamide (9.8 mg).

[0262]

Example 20

N-(2-((3-(2',5'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino) ethyl)-3-methoxybenzamide

[0263]



30 [0264]

In the same manner as in Example 12, tert-butyl (2-((3-(2',5'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino) ethyl) carbamate (70.1 mg) was deprotected with TFA, and amidated with m-anisic

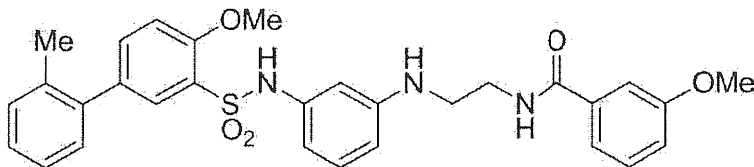
acid (11.4 mg) to give N-(2-((3-(2',5'-difluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (11.9 mg).

[0265]

5 Example 21

3-methoxy-N-(2-((3-(4-methoxy-2'-methyl-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide

[0266]



10 [0267]

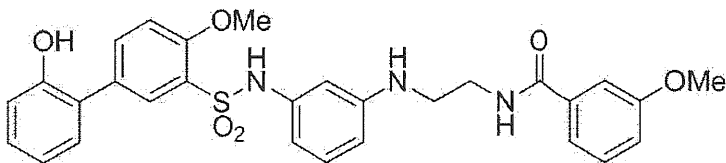
To tert-butyl (2-((3-(4-methoxy-2'-methyl-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (11.7 mg) was added 10% hydrogen chloride-methanol solution (2.0 mL), and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated, to a suspension of the obtained residue in dichloromethane (1.0 mL) were added m-anisic acid (3.5 mg), triethylamine (0.011 mL) and BOP reagent (11.1 mg), and the mixture was stirred at room temperature overnight under an argon atmosphere. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ethyl acetate/hexane=3/2) to give 3-methoxy-N-(2-((3-(4-methoxy-2'-methyl-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide (5.0 mg).

[0268]

Example 22

30 N-(2-((3-(2'-hydroxy-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide

[0269]



[0270]

35 In the same manner as in Example 21, tert-butyl (2-((3-(2'-hydroxy-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)carbamate (7.2 mg) was

deprotected with 10% hydrogen chloride-methanol solution, and amidated with m-anisic acid (2.1 mg) to give N-(2-((3-(2'-hydroxy-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (8.2 mg).

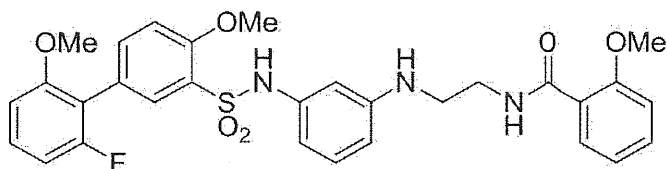
5 For purification, silica gel column chromatography (eluate: ethyl acetate/chloroform =1/0→1/1) was used.

[0271]

Example 23

N-(2-((3-(2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

[0272]



[0273]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.5 mg) was amidated with o-anisic acid (5.8 mg) to give N-(2-((3-(2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (15.8 mg). For purification, silica gel

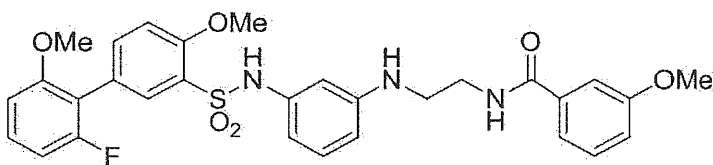
20 column chromatography (eluate: ethyl acetate/hexane=1/2→1/1) was used.

[0274]

Example 24

N-(2-((3-(2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide

[0275]



[0276]

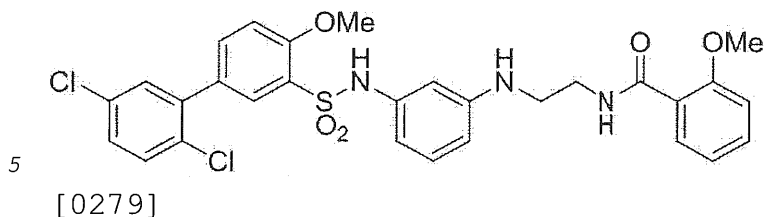
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (17.1 mg) was amidated with m-anisic acid (5.0 mg) to give N-(2-((3-(2'-fluoro-4,6'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (14.6 mg). For purification, silica gel

35 column chromatography (eluate: ethyl acetate/hexane=1/2→1/1) was used.

[0277]

Example 25

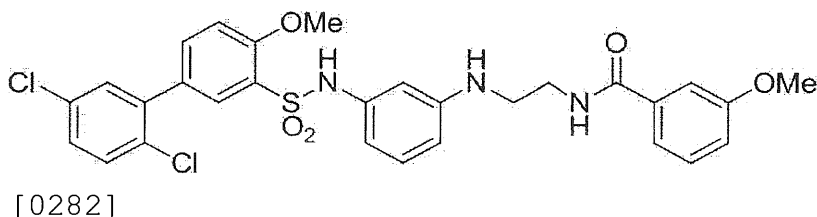
N-(2-((3-(2',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide
[0278]



In the same manner as in Example 21, tert-butyl (2-((3-(2',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino)ethyl) carbamate
10 (177 mg) was deprotected with 10% hydrogen chloride-methanol solution, and amidated with o-anisic acid (19.8 mg) to give N-(2-((3-(2',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (35.8 mg). For purification, silica gel column chromatography (eluate:
15 ethyl acetate/hexane=3/2→2/1) was used.
[0280]

Example 26

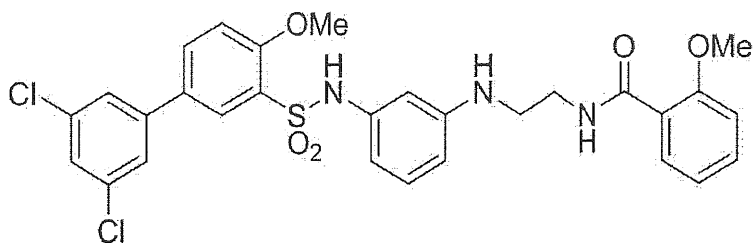
N-(2-((3-(2',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide
20 [0281]



In the same manner as in Example 21, tert-butyl (2-((3-(2',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino)ethyl) carbamate
25 (177 mg) was deprotected with 10% hydrogen chloride-methanol solution, and amidated with m-anisic acid (19.8 mg) to give N-(2-((3-(2',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (13.8 mg).
30 For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/2→2/1) was used.
[0283]

Example 27

N-(2-((3-(3',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide
35 [0284]



[0285]

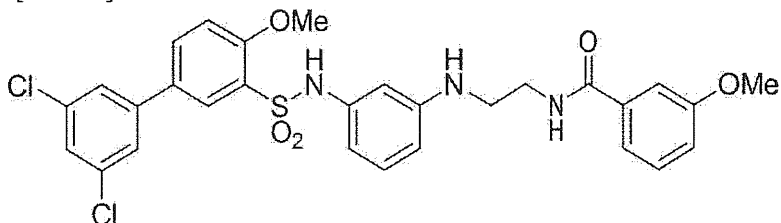
In the same manner as in Example 21, tert-butyl (2-((3-(3',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino) ethyl) carbamate (137 mg) was deprotected with 10% hydrogen chloride-methanol solution, and amidated with o-anisic acid (15.2 mg) to give N-(2-((3-(3',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino) ethyl)-2-methoxybenzamide (42.5 mg).
 10 For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/2→2/1) was used.

[0286]

Example 28

N-(2-((3-(3',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino) ethyl)-3-methoxybenzamide

[0287]



[0288]

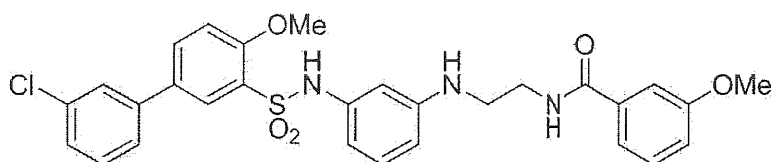
In the same manner as in Example 21, tert-butyl (2-((3-(3',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino) ethyl) carbamate (137 mg) was deprotected with 10% hydrogen chloride-methanol solution, and amidated with m-anisic acid (15.2 mg) to give N-(2-((3-(3',5'-dichloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino) ethyl)-3-methoxybenzamide (20.5 mg).
 25 For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/2→2/1) was used.

[0289]

Example 29

N-(2-((3-(3'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino) ethyl)-3-methoxybenzamide

[0290]



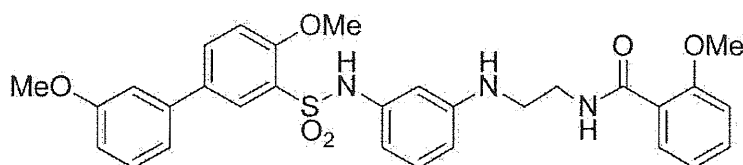
[0291]

In the same manner as in Example 21, tert-butyl (2-((3-(3'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino)ethyl) carbamate (104 mg) was deprotected with 10% hydrogen chloride-methanol solution, and amidated with m-anisic acid (8.5 mg) to give N-(2-((3-(3'-chloro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino)ethyl)-3-methoxybenzamide (13.5 mg).
 For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=1/1) was used.

[0292]

Example 30

N-(2-((3-(3',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino)ethyl)-2-methoxybenzamide
 [0293]



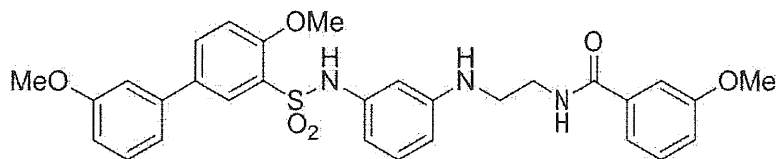
[0294]

In the same manner as in Example 1, N-(3-((2-aminoethyl) amino)phenyl)-3',4-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (16.4 mg) was amidated with o-anisic acid (5.0 mg) to give N-(2-((3-(3',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino)ethyl)-2-methoxybenzamide (4.1 mg).

[0295]

Example 31

N-(2-((3-(3',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) amino)ethyl)-3-methoxybenzamide
 [0296]



30

[0297]

In the same manner as in Example 1, N-(3-((2-aminoethyl) amino)phenyl)-3',4-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (16.4 mg) was amidated with m-

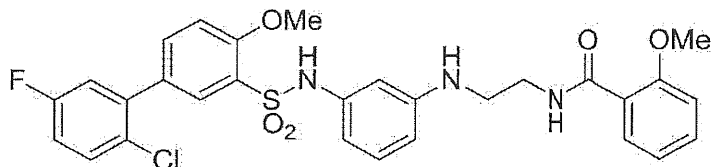
anisic acid (5.0 mg) to give N-(2-((3-(3',4-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (6.9 mg).

[0298]

5 Example 32

N-(2-((3-(2'-chloro-5'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

[0299]



10 [0300]

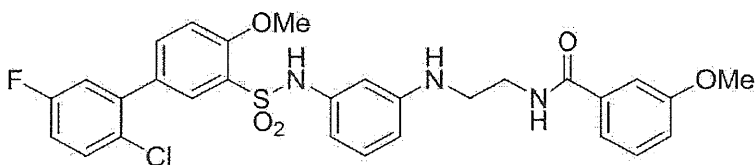
In the same manner as in Example 21, tert-butyl (2-((3-(2'-chloro-5'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino)ethyl) carbamate (127 mg) was deprotected with 10% hydrogen chloride-methanol solution, and amidated with o-anisic acid (16.7 mg) to give N-(2-((3-(2'-chloro-5'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (15.6 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/2) was used.

20 [0301]

Example 33

N-(2-((3-(2'-chloro-5'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide

[0302]



25

[0303]

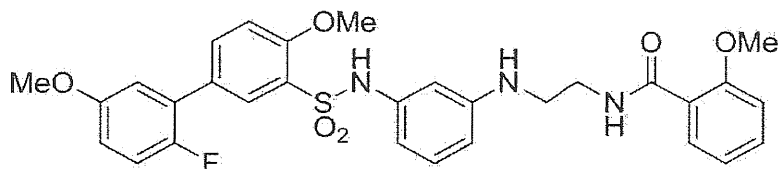
In the same manner as in Example 21, tert-butyl (2-((3-(2'-chloro-5'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl) (4-methoxybenzyl) amino)ethyl) carbamate (127 mg) was deprotected with 10% hydrogen chloride-methanol solution, and amidated with m-anisic acid (16.7 mg) to give N-(2-((3-(2'-chloro-5'-fluoro-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (10.4 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/2) was used.

35

[0304]

Example 34

N-(2-((3-(2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide
[0305]



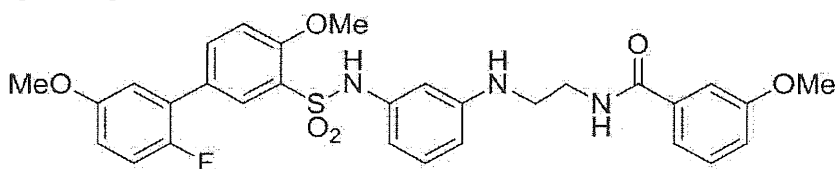
5 [0306]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (18.1 mg) was amidated with o-anisic acid (5.3 mg) to give N-(2-((3-(2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (8.2 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1) was used.

[0307]

15 Example 35

N-(2-((3-(2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide
[0308]



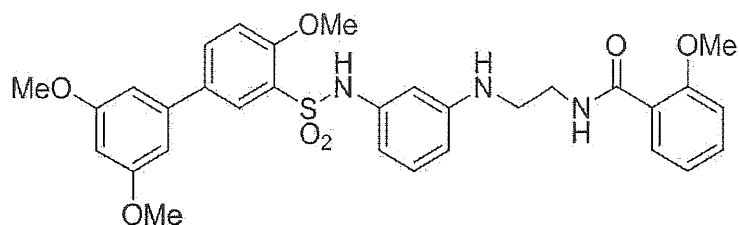
20 [0309]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (18.1 mg) was amidated with m-anisic acid (5.3 mg) to give N-(2-((3-(2'-fluoro-4,5'-dimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-3-methoxybenzamide (12.9 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1) was used.

[0310]

30 Example 36

2-methoxy-N-(2-((3-(3',4,5'-trimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide
[0311]



[0312]

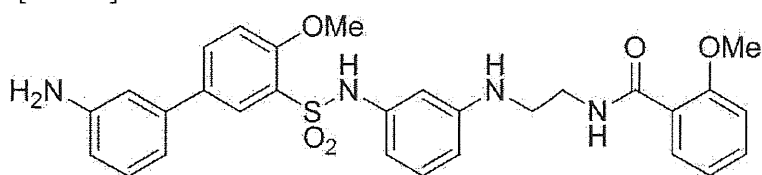
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-3',4,5'-trimethoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (16.2 mg) was amidated with o-anisic acid (4.7 mg) to give 2-methoxy-N-(2-((3-(3',4,5'-trimethoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)benzamide (12.4 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=2/1→1/0) was used.

[0313]

Example 37

N-(2-((3-(3'-amino-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

[0314]



[0315]

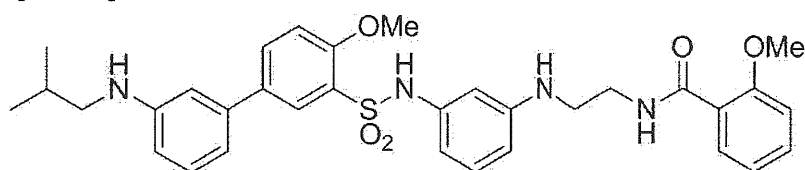
In the same manner as in Example 1, 3'-amino-N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-[1,1'-biphenyl]-3-sulfonamide (44.9 mg) was amidated with o-anisic acid (16.7 mg) to give N-(2-((3-(3'-amino-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (53.1 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0316]

Example 38

N-(2-((3-(3'-(isobutylamino)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide

[0317]



30

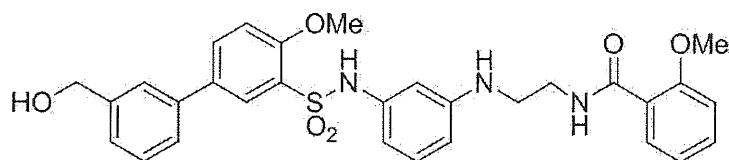
[0318]

To a solution of N-(2-((3-(3'-amino-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-

methoxybenzamide (53.6 mg) in 1,2-dichloroethane (2.0 mL) were added isobutylaldehyde (0.01 mL) and sodium triacetoxyborohydride (31.8 mg), and the mixture was stirred at room temperature for 4 hr. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (ammonia-saturated chloroform/methanol=40/1) to give N-(2-((3-(3'-(isobutylamino)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (32.3 mg).
[0319]

Example 39

N-(2-((3-(3'-(hydroxymethyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide
[0320]

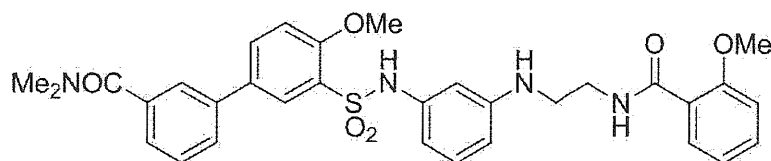


[0321]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-3'-hydroxymethyl-4-methoxy-[1,1'-biphenyl]-3-sulfonamide (25.3 mg) was amidated with o-anisic acid (9.0 mg) to give N-(2-((3-(3'-(hydroxymethyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxybenzamide (20.6 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.
[0322]

Example 40

4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide
[0323]



[0324]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (86.6 mg) was

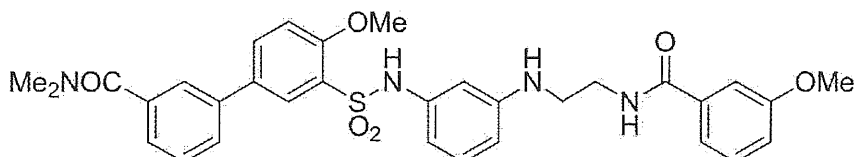
amidated with o-anisic acid (24.3 mg) to give 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (85.3 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0325]

Example 41

4'-methoxy-3'-(N-(3-((2-(3-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

[0326]



[0327]

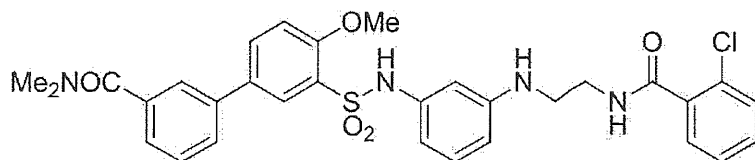
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with m-anisic acid (5.3 mg) to give 4'-methoxy-3'-(N-(3-((2-(3-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (17.9 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0328]

Example 42

3'-(N-(3-((2-(2-chlorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

[0329]



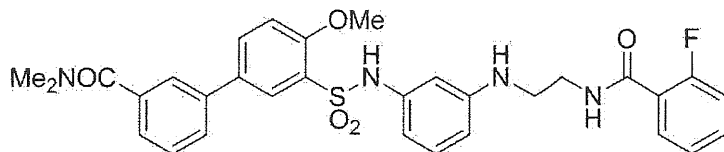
[0330]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 2-chlorobenzoic acid (5.5 mg) to give 3'-(N-(3-((2-(2-chlorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (17.2 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0331]

Example 43

3'-(N-(3-((2-(2-fluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide
[0332]



[0333]

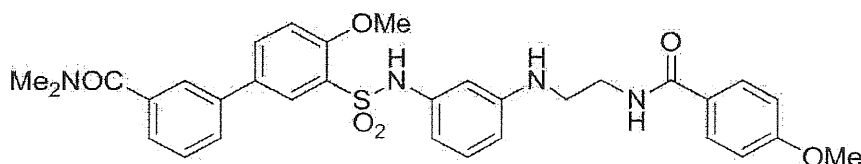
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
10 amidated with 2-fluorobenzoic acid (4.9 mg) to give 3'-(N-(3-((2-(2-fluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (19.9 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

15 [0334]

Example 44

4'-methoxy-3'-(N-(3-((2-(4-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

20 [0335]



[0336]

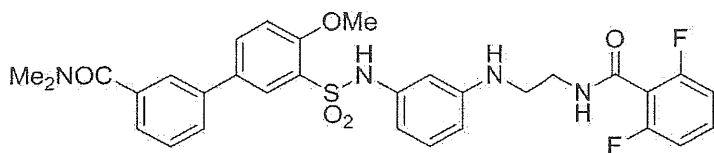
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
25 amidated with p-anisic acid (5.5 mg) to give 4'-methoxy-3'-(N-(3-((2-(4-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (20.6 mg). For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=40/1) was used.

30 [0337]

Example 45

3'-(N-(3-((2-(2,6-difluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

35 [0338]



[0339]

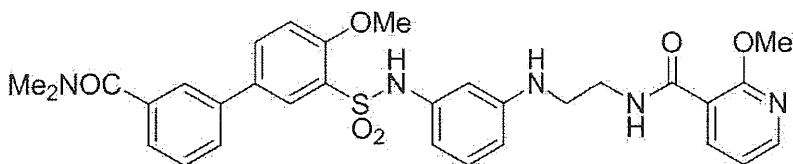
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (22.3 mg) was
 5 amidated with 2,6-difluorobenzoic acid (6.5 mg) to give 3'-(N-(3-((2-(2,6-difluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (22.1 mg). For purification, silica gel column chromatography
 10 (eluate: ethyl acetate) was used.

[0340]

Example 46

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxynicotinamide

15 [0341]



[0342]

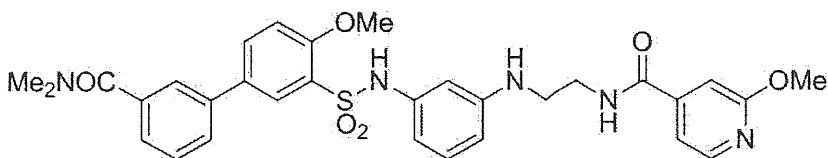
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
 20 amidated with 2-methoxypyridine-3-carboxylic acid (5.4 mg) to give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxynicotinamide (20.6 mg). For purification, preparative
 25 thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=40/1) was used.

[0343]

Example 47

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-methoxyisonicotinamide

30 [0344]



[0345]

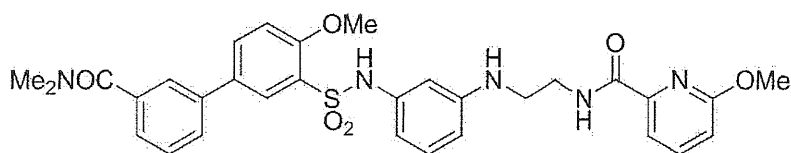
In the same manner as in Example 1, 3'-(N-(3-((2-

aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
amidated with 2-methoxypyridine-4-carboxylic acid (5.4 mg) to
give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-
5 biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-
methoxynicotinamide (20.1 mg). For purification, preparative
thin layer chromatography (eluate: ammonia-saturated
chloroform/methanol=40/1) was used.

[0346]

10 Example 48

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-
ylsulfonamide)phenyl)amino)ethyl)-6-methoxypicolinamide
[0347]



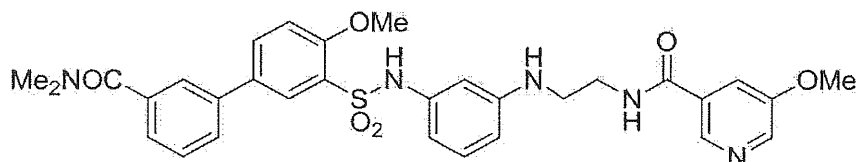
15 [0348]

In the same manner as in Example 1, 3'-(N-(3-((2-
aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
amidated with 6-methoxypyridine-2-carboxylic acid (5.4 mg) to
20 give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-
biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-
methoxynicotinamide (18.3 mg). For purification, preparative
thin layer chromatography (eluate: ammonia saturated
chloroform/methanol=40/1) was used.

25 [0349]

Example 49

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-
ylsulfonamide)phenyl)amino)ethyl)-5-methoxynicotinamide
[0350]



30

[0351]

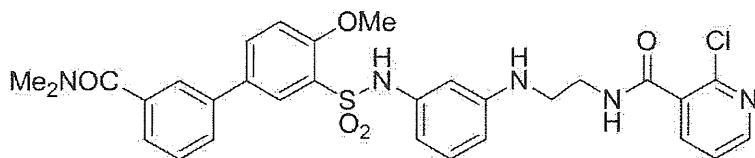
In the same manner as in Example 1, 3'-(N-(3-((2-
aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
35 amidated with 5-methoxypyridine-3-carboxylic acid (5.4 mg) to
give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-
biphenyl]-3-ylsulfonamide)phenyl)amino)ethyl)-2-

methoxynicotinamide (18.3 mg). For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=40/1) was used.

[0352]

5 Example 50

2-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)nicotinamide
[0353]

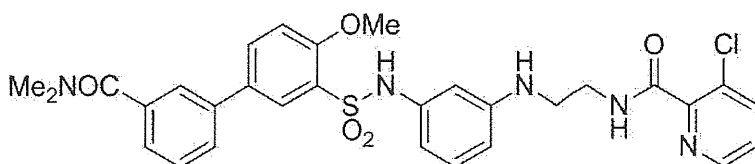


10 [0354]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 2-chloronicotinic acid (5.5 mg) to give 2-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)nicotinamide (20.1 mg). For purification, amine silica gel column chromatography (eluate: chloroform/methanol=1/0→50/1) was used.
[0355]

20 Example 51

3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)picolinamide
[0356]



25 [0357]

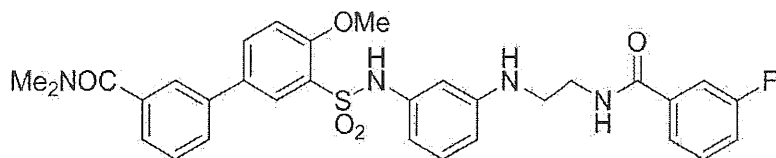
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 3-chloropyridine-2-carboxylic acid (5.5 mg) to give 3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)picolinamide (18.2 mg). For purification, amine silica gel column chromatography (eluate: chloroform/methanol=1/0→100/1) was used.

35 [0358]

Example 52

3'-(N-(3-((2-(3-fluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-

4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide
[0359]



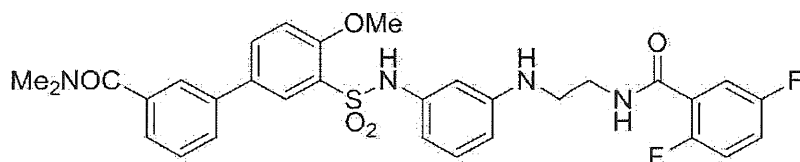
[0360]

5 In the same manner as in Example 1, 3'-((N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 3-fluorobenzoic acid (4.9 mg) to give 3'-((N-(3-((2-(3-fluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (13.5 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0361]

Example 53

15 3'-((N-(3-((2-(2,5-difluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide
[0362]



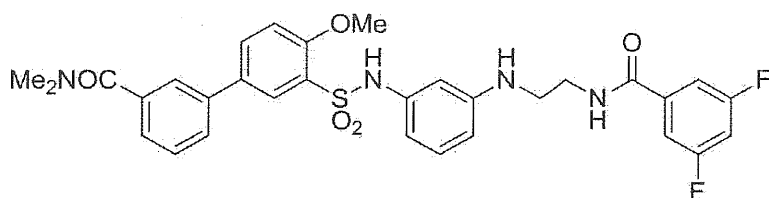
20 [0363]

In the same manner as in Example 1, 3'-((N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 2,5-difluorobenzoic acid (5.5 mg) to give 3'-((N-(3-((2-(2,5-difluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (15.3 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0364]

30 Example 54

3'-((N-(3-((2-(3,5-difluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide
[0365]



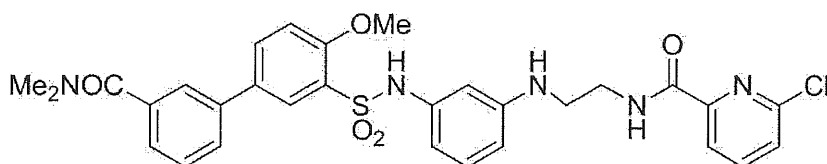
[0366]

In the same manner as in Example 1, 3'-((N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
 5 [1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
 amidated with 3,5-difluorobenzoic acid (5.5 mg) to give 3'-((N-
 (3-((2-(3,5-difluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-
 4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (18.3
 mg). For purification, silica gel column chromatography
 10 (eluate: ethyl acetate) was used.

[0367]

Example 55

6-chloro-N-(2-((3-(3'-((dimethylcarbamoyl)-4-methoxy-[1,1'-
 biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)picolinamide
 15 [0368]



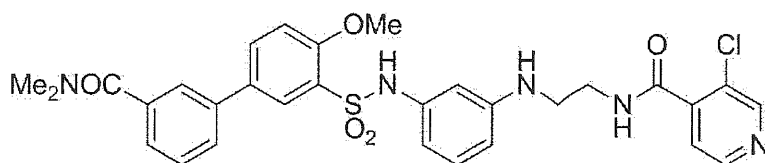
[0369]

In the same manner as in Example 1, 3'-((N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
 20 [1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
 amidated with 6-chloropyridine-2-carboxylic acid (5.5 mg) to
 give 6-chloro-N-(2-((3-(3'-((dimethylcarbamoyl)-4-methoxy-
 [1,1'-biphenyl]-3-
 ylsulfonamido)phenyl)amino)ethyl)picolinamide (20.3 mg). For
 25 purification, amine silica gel column chromatography (eluate:
 chloroform/methanol=1/0→100/1) was used.

[0370]

Example 56

3-chloro-N-(2-((3-(3'-((dimethylcarbamoyl)-4-methoxy-[1,1'-
 30 biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)isonicotinamide
 [0371]



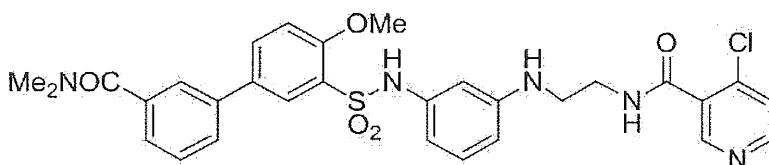
[0372]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 3-chloropyridine-4-carboxylic acid (5.5 mg) to give 3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)isonicotinamide (18.5 mg). For purification, amine silica gel column chromatography (eluate: chloroform/methanol=1/0→100/1) was used.

[0373]

Example 57

4-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)nicotinamide [0374]



15

[0375]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 4-chloropyridine-3-carboxylic acid (5.5 mg) to give 3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)nicotinamide (12.9 mg). For purification, amine silica gel column chromatography (eluate: chloroform/methanol=1/0→100/1) was used.

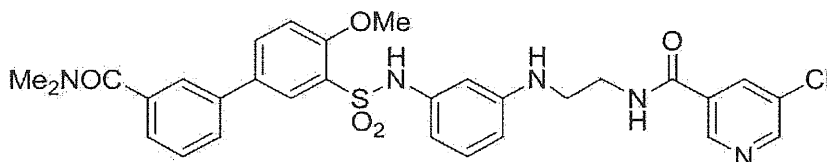
25

[0376]

Example 58

5-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)nicotinamide

30 [0377]



[0378]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 5-chloropyridine-3-carboxylic acid (5.5 mg) to give 3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-

35

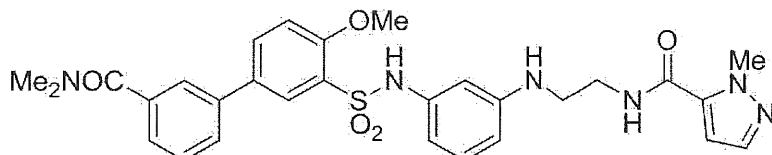
[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)nicotinamide (18.0 mg). For purification, amine silica gel column chromatography (eluate: chloroform/methanol=1/0→100/1) was used.

5 [0379]

Example 59

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-1-methyl-1H-pyrazole-5-carboxamide

10 [0380]



[0381]

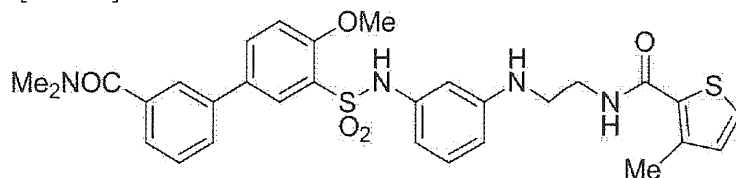
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
15 amidated with 1-methyl-1H-pyrazole-5-carboxylic acid (4.4 mg) to give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-1-methyl-1H-pyrazole-5-carboxamide (15.7 mg). For purification, amine
20 silica gel column chromatography (eluate: chloroform/methanol=1/0→100/1) was used.

[0382]

Example 60

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-3-methylthiophene-2-
25 carboxamide

[0383]



[0384]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
30 amidated with 3-methylthiophene-2-carboxylic acid (5.0 mg) to give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-3-methylthiophene-2-carboxamide (16.4 mg). For purification,
35 silica gel column chromatography (eluate: ethyl acetate) was

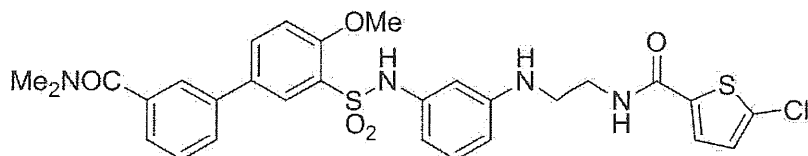
used.

[0385]

Example 61

5-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-
5 biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)thiophene-2-
carboxamide

[0386]



[0387]

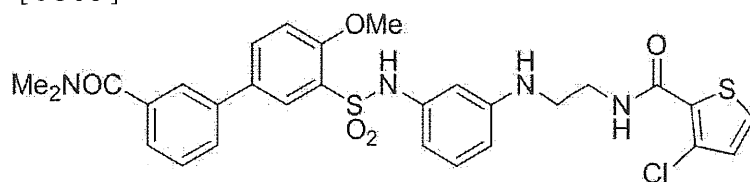
10 In the same manner as in Example 1, 3'-(N-(3-((2-
aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
amidated with 5-chlorothiophene-2-carboxylic acid (5.7 mg) to
give 5-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-
15 [1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)thiophene-
2-carboxamide (15.8 mg). For purification, silica gel column
chromatography (eluate: ethyl acetate) was used.

[0388]

Example 62

20 3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-
biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)thiophene-2-
carboxamide

[0389]



25 [0390]

In the same manner as in Example 1, 3'-(N-(3-((2-
aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
amidated with 3-chlorothiophene-2-carboxylic acid (5.7 mg) to
30 give 3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-
[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)thiophene-
2-carboxamide (19.7 mg). For purification, silica gel column
chromatography (eluate: ethyl acetate) was used.

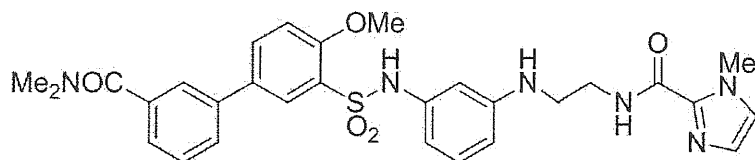
[0391]

35 Example 63

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-
ylsulfonamido)phenyl)amino)ethyl)-1-methyl-1H-imidazole-2-

carboxamide

[0392]



[0393]

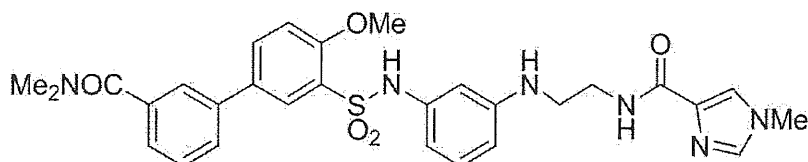
5 In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 1-methyl-1H-imidazole-2-carboxylic acid (4.9 mg) to give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-
10 biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-1-methyl-1H-imidazole-2-carboxamide (20.3 mg). For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=40/1) was used.

[0394]

15 Example 64

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-1-methyl-1H-imidazole-4-carboxamide

[0395]



20

[0396]

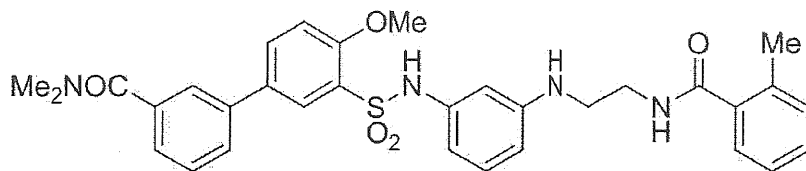
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
25 amidated with 1-methyl-1H-imidazole-4-carboxylic acid (4.4 mg) to give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-1-methyl-1H-imidazole-4-carboxamide (19.7 mg). For purification, preparative thin layer chromatography (eluate: ammonia-
30 saturated chloroform/methanol=40/1) was used.

[0397]

Example 65

4'-methoxy-N,N-dimethyl-3'-(N-(3-((2-(2-methylbenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-
35 3-carboxamide

[0398]



[0399]

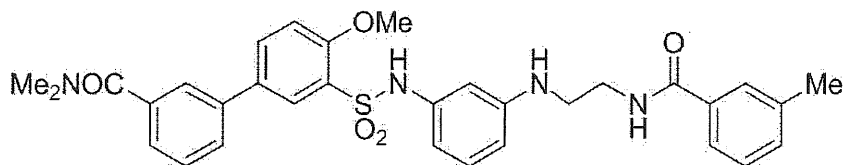
In the same manner as in Example 1, 3'-((N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
 5 amidated with 2-toluic acid (4.8 mg) to give 4'-methoxy-N,N-dimethyl-3'-((N-(3-((2-(2-methylbenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxamide (20.1 mg). For purification, silica gel column
 10 chromatography (eluate: ethyl acetate) was used.

[0400]

Example 66

4'-methoxy-N,N-dimethyl-3'-((N-(3-((2-(3-methylbenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-
 15 3-carboxamide

[0401]



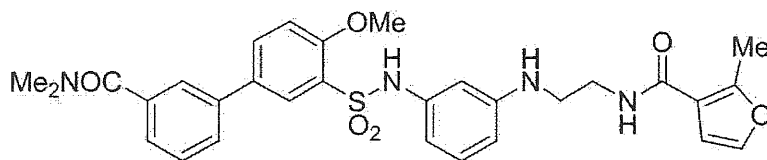
[0402]

In the same manner as in Example 1, 3'-((N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
 20 [1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 3-toluic acid (4.8 mg) to give 4'-methoxy-N,N-dimethyl-3'-((N-(3-((2-(3-methylbenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxamide (15.8 mg). For purification, silica gel column
 25 chromatography (eluate: ethyl acetate) was used.

[0403]

Example 67

N-(2-((3-(3'-((dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-2-methylfuran-3-carboxamide
 30 [0404]



[0405]

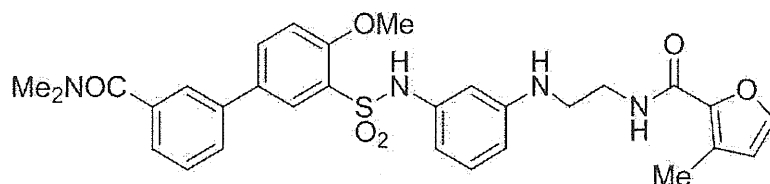
In the same manner as in Example 1, 3'-((N-(3-((2-

aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
amidated with 2-methylfuran-3-carboxylic acid (4.4 mg) to give
N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-
5 ylsulfonamido)phenyl)amino)ethyl)-2-methylfuran-3-carboxamide
(18.0 mg). For purification, silica gel column chromatography
(eluate: ethyl acetate) was used.

[0406]

Example 68

10 N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-
ylsulfonamido)phenyl)amino)ethyl)-3-methylfuran-2-carboxamide
[0407]



[0408]

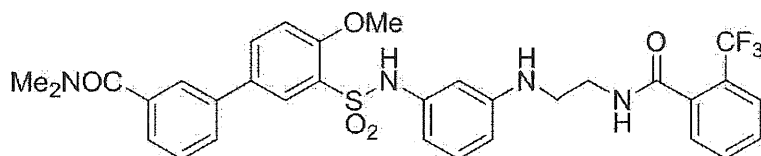
15 In the same manner as in Example 1, 3'-(N-(3-((2-
aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
amidated with 3-methylfuran-2-carboxylic acid (4.4 mg) to give
N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-[1,1'-biphenyl]-3-
20 ylsulfonamido)phenyl)amino)ethyl)-3-methylfuran-2-carboxamide
(20.0 mg). For purification, silica gel column chromatography
(eluate: ethyl acetate) was used.

[0409]

Example 69

25 4'-methoxy-N,N-dimethyl-3'-(N-(3-((2-(2-
(trifluoromethyl)benzamido)ethyl)amino)phenyl)sulfamoyl)-
[1,1'-biphenyl]-3-carboxamide

[0410]



30 [0411]

In the same manner as in Example 1, 3'-(N-(3-((2-
aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-
[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
amidated with 2-trifluoromethylbenzoic acid (6.7 mg) to give
35 4'-methoxy-N,N-dimethyl-3'-(N-(3-((2-(2-
(trifluoromethyl)benzamido)ethyl)amino)phenyl)sulfamoyl)-

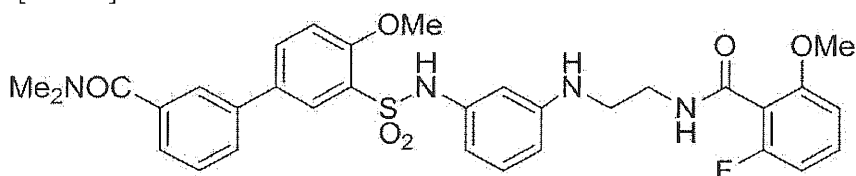
[1,1'-biphenyl]-3-carboxamide (21.0 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0412]

5 Example 70

3'-(N-(3-((2-(2-fluoro-6-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

[0413]



[0414]

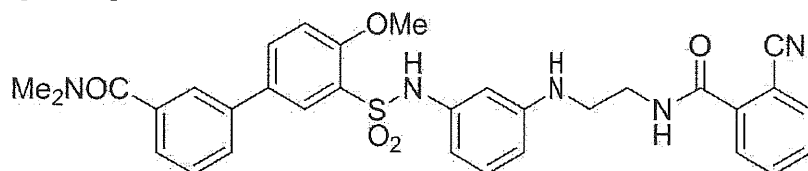
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
15 amidated with 2-fluoro-6-methoxybenzoic acid (6.0 mg) to give 3'-(N-(3-((2-(2-fluoro-6-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (23.6 mg). For purification, preparative thin layer chromatography (eluate:
20 ammonia-saturated chloroform/methanol=80/1) was used.

[0415]

Example 71

3'-(N-(3-((2-(2-cyanobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

25 [0416]



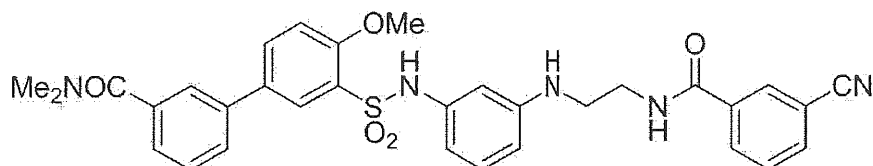
[0417]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was
30 amidated with 2-cyanobenzoic acid (5.1 mg) to give 3'-(N-(3-((2-(2-cyanobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (19.2 mg). For purification, amine silica gel column chromatography
35 (eluate: chloroform/methanol=1/0→100/1) was used.

[0418]

Example 72

3'-(N-(3-((2-(3-cyanobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide
[0419]

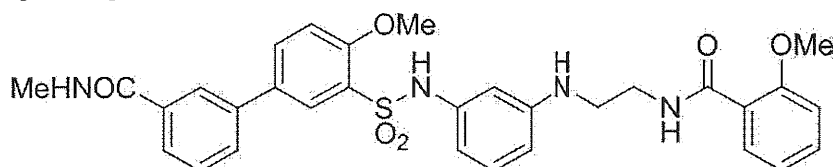


5 [0420]

In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.0 mg) was amidated with 3-cyanobenzoic acid (5.1 mg) to give 3'-(N-(3-((2-(3-cyanobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (21.3 mg). For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=100/1) was used.
[0421]

15 Example 73

4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N-methyl-[1,1'-biphenyl]-3-carboxamide
[0422]



20

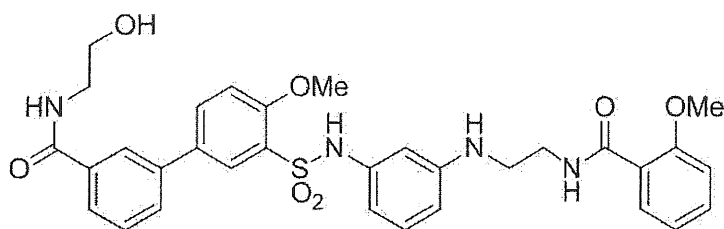
[0423]

To a suspension of 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (19.0 mg) in dichloromethane (2.0 mL) were added methylamine hydrochloride (7.4 mg), triethylamine (0.021 mL) and BOP reagent (17.2 mg), and the mixture was stirred at room temperature overnight. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate) to give 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N-methyl-[1,1'-biphenyl]-3-carboxamide (21.8 mg).
[0424]

35

Example 74

N-(2-hydroxyethyl)-4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxamide
[0425]



[0426]

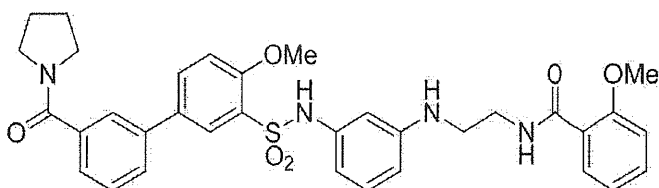
In the same manner as in Example 73, 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (20.1 mg) was amidated with 2-aminoethanol (0.006 mL) to give N-(2-hydroxyethyl)-4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxamide (18.2 mg). For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=20/1) was used.

[0427]

Example 75

2-methoxy-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-1-[1,1'-biphenyl]-3-yl)sulfonamido)phenyl)amino)ethyl)benzamide

[0428]



[0429]

In the same manner as in Example 73, 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (20.1 mg) was amidated with pyrrolidine (0.009 mL) to give 2-methoxy-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-1-[1,1'-biphenyl]-3-yl)sulfonamido)phenyl)amino)ethyl)benzamide (18.4 mg). For purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=40/1) was used.

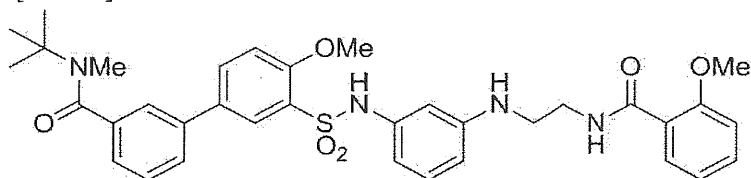
[0430]

Example 76

N-(tert-butyl)-4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N-methyl-[1,1'-

biphenyl]-3-carboxamide

[0431]



[0432]

5 In the same manner as in Example 73, 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (20.1 mg) was amidated with tert-butylmethylamine (0.008 mL) to give N-(tert-butyl)-4'-methoxy-3'-(N-(3-((2-(2-

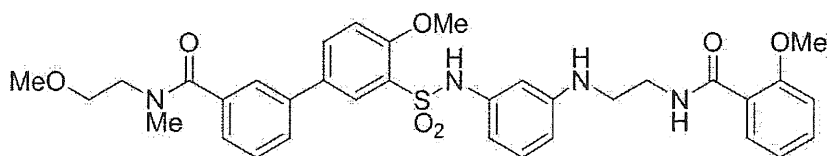
10 methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N-methyl-[1,1'-biphenyl]-3-carboxamide (21.6 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0433]

Example 77

15 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N-(2-methoxyethyl)-N-methyl-[1,1'-biphenyl]-3-carboxamide

[0434]



20 [0435]

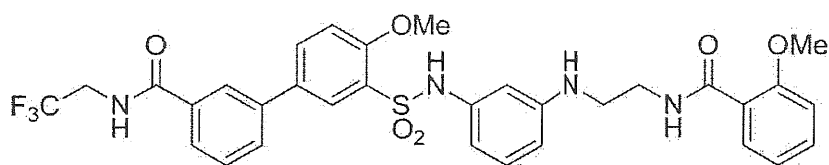
In the same manner as in Example 73, 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (20.1 mg) was amidated with N-(2-methoxyethyl)-N-methylamine (0.004 mL) to give 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N-(2-methoxyethyl)-N-methyl-[1,1'-biphenyl]-3-carboxamide (21.3 mg). For purification, preparative thin layer chromatography (eluate: ethyl acetate) was used.

[0436]

30 Example 78

4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N-(2,2,2-trifluoroethyl)-[1,1'-biphenyl]-3-carboxamide

[0437]



[0438]

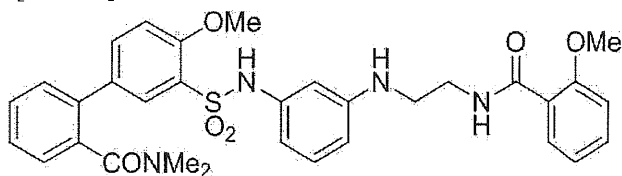
In the same manner as in Example 73, 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (22.4 mg) was amidated with 2,2,2-trifluoroethylamine hydrochloride (5.3 mg) to give 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N-(2,2,2-trifluoroethyl)-[1,1'-biphenyl]-3-carboxamide (20.1 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→1/0) was used.

[0439]

Example 79

4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-2-carboxamide

[0440]



[0441]

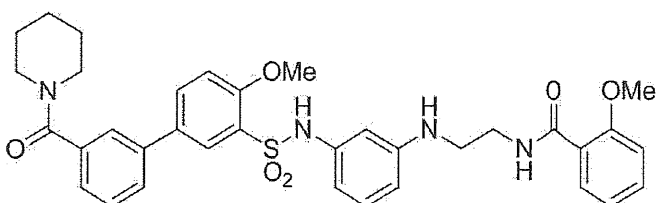
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N-dimethyl-[1,1'-biphenyl]-2-carboxamide dihydrochloride (19.0 mg) was amidated with o-anisic acid (5.3 mg) to give 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-2-carboxamide (19.7 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→1/0) was used.

[0442]

Example 80

2-methoxy-N-(2-((3-(4-methoxy-3'-(piperidine-1-carbonyl)-1-[1,1'-biphenyl]-3-yl)sulfonamido)phenyl)amino)ethyl)benzamide

[0443]



[0444]

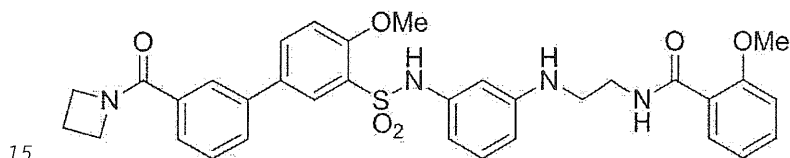
In the same manner as in Example 73, 4'-methoxy-3'-(N-(3-
((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-
biphenyl]-3-carboxylic acid (20.7 mg) was amidated with
5 piperidine (0.004 mL) to give 2-methoxy-N-(2-((3-(4-methoxy-
3'-(piperidine-1-carbonyl)-1-[1,1'-biphenyl]-3-
ylsulfonamido)phenyl)amino)ethyl)benzamide (14.0 mg). For
purification, silica gel column chromatography (eluate: ethyl
acetate) was used.

10 [0445]

Example 81

N-(2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-
3-ylsulfonamido)phenyl)amino)ethyl)-2-methoxybenzamide

[0446]



[0447]

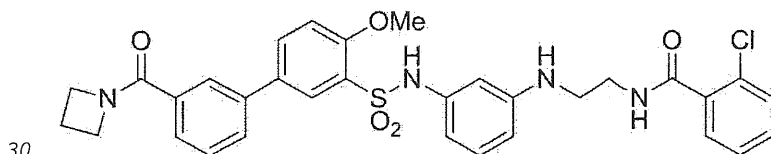
In the same manner as in Example 73, 4'-methoxy-3'-(N-(3-
((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-
biphenyl]-3-carboxylic acid (20.7 mg) was amidated with
20 azetidine hydrochloride (3.4 mg) to give N-(2-((3-(3'-
(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-
ylsulfonamido)phenyl)amino)ethyl)-2-methoxybenzamide (12.0 mg).
For purification, preparative thin layer chromatography
(eluate: ammonia-saturated chloroform/methanol=40/1) was used.

25 [0448]

Example 82

N-(2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-
3-ylsulfonamido)phenyl)amino)ethyl)-2-chlorobenzamide

[0449]



[0450]

In the same manner as in Example 1, N-(3-((2-
aminoethyl)amino)phenyl)-3'-(azetidine-1-carbonyl)-4-methoxy-
[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.4 mg) was
35 amidated with 2-chlorobenzoic acid (5.5 mg) to give N-(2-((3-
(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-
ylsulfonamido)phenyl)amino)ethyl)-2-chlorobenzamide (13.2 mg).

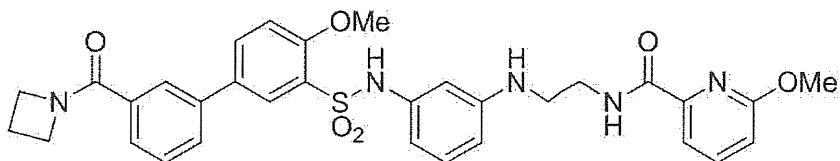
For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→4/1) was used.

[0451]

Example 83

- 5 N-(2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-6-methoxypicolinamide

[0452]



[0453]

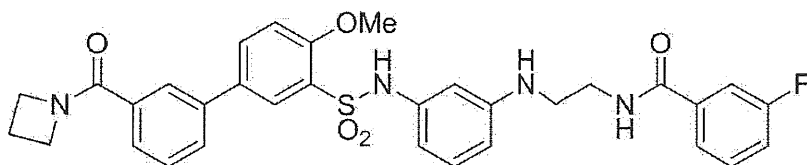
- 10 In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.4 mg) was amidated with 6-methoxypyridine-2-carboxylic acid (5.4 mg) to give N-(2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-6-methoxypicolinamide (16.1 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=4/1) was used.

[0454]

20 Example 84

N-(2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-3-fluorobenzamide

[0455]



25 [0456]

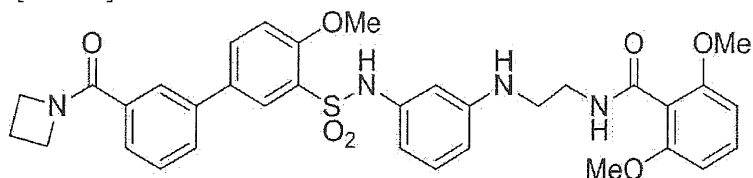
- In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.4 mg) was amidated with 3-fluorobenzoic acid (4.9 mg) to give N-(2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-3-fluorobenzamide (15.0 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→1/0) was used.

[0457]

35 Example 85

N-(2-((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-2,6-dimethoxybenzamide

[0458]



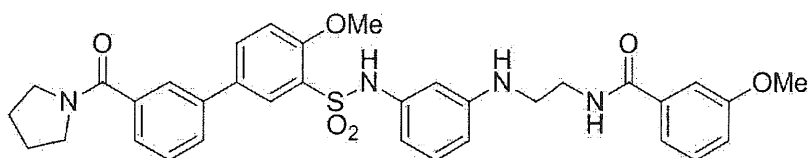
[0459]

In the same manner as in Example 1, N-(3-((2-
5 aminoethyl)amino)phenyl)-3'-(azetidine-1-carbonyl)-4-methoxy-
[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.4 mg) was
amidated with 2,6-dimethoxybenzoic acid (6.4 mg) to give N-(2-
((3-(3'-(azetidine-1-carbonyl)-4-methoxy-[1,1'-biphenyl]-3-
ylsulfonamido)phenyl)amino)ethyl)-2,6-dimethoxybenzamide (15.2
10 mg). For purification, silica gel column chromatography
(eluate: ethyl acetate) was used.

[0460]

Example 86

3-methoxy-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-
15 [1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide
[0461]



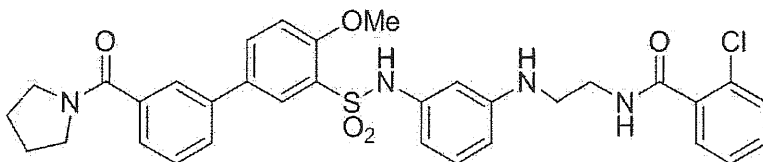
[0462]

In the same manner as in Example 1, N-(3-((2-
20 aminoethyl)amino)phenyl)-4-methoxy-3'-(pyrrolidine-1-
carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.9
mg) was amidated with m-anisic acid (5.3 mg) to give 3-
methoxy-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-
biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide (16.3
25 mg). For purification, silica gel column chromatography
(eluate: ethyl acetate/hexane=3/1→1/0) was used.

[0463]

Example 87

2-chloro-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-
30 [1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide
[0464]



[0465]

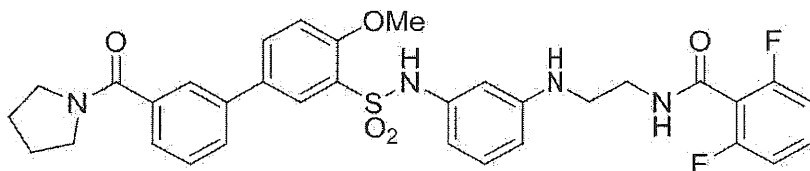
In the same manner as in Example 1, N-(3-((2-

aminoethyl)amino)phenyl)-4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.9 mg) was amidated with 2-chlorobenzoic acid (5.5 mg) to give 2-chloro-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-
5 biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide (16.7 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0466]

Example 88

10 2,6-difluoro-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide
[0467]



[0468]

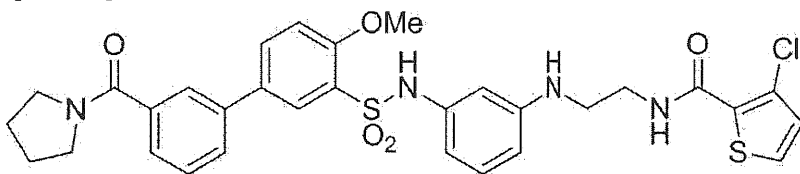
15 In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.9 mg) was amidated with 2,6-difluorobenzoic acid (5.5 mg) to give 2,6-difluoro-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-
20 carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide (16.8 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0469]

25 Example 89

3-chloro-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)thiophene-2-carboxamide

[0470]



30

[0471]

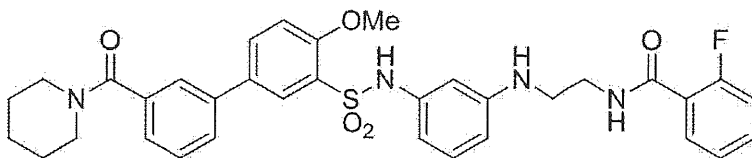
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (19.9
35 mg) was amidated with 3-chlorothiophene-2-carboxylic acid (5.7 mg) to give 3-chloro-N-(2-((3-(4-methoxy-3'-(pyrrolidine-1-carbonyl)-[1,1'-biphenyl]-3-

ylsulfonamido)phenyl)amino)ethyl)thiophene-2-carboxamide (15.9 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0472]

5 Example 90

2-fluoro-N-(2-((3-(4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide
[0473]



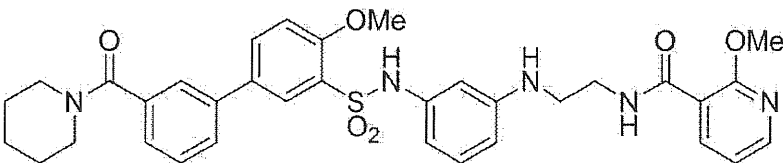
10 [0474]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (20.4 mg) was amidated with 2-fluorobenzoic acid (4.9 mg) to give 2-fluoro-N-(2-((3-(4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide (14.4 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→1/0) was used.

[0475]

20 Example 91

2-methoxy-N-(2-((3-(4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)nicotinamide
[0476]



25

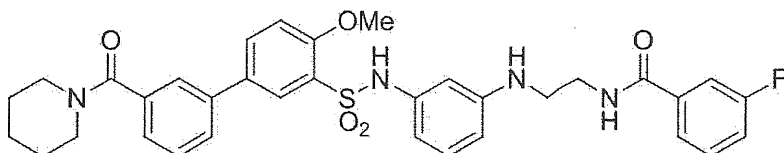
[0477]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (20.4 mg) was amidated with 2-methoxypyridine-3-carboxylic acid (5.4 mg) to give 2-methoxy-N-(2-((3-(4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)nicotinamide (16.7 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0478]

Example 92

3-fluoro-N-(2-((3-(4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide
[0479]



5 [0480]

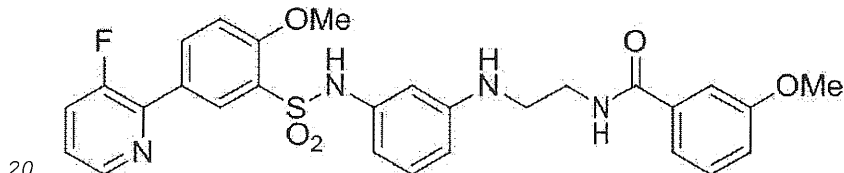
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-sulfonamide dihydrochloride (20.4 mg) was amidated with 3-fluorobenzoic acid (4.9 mg) to give 3-fluoro-N-(2-((3-(4-methoxy-3'-(piperidine-1-carbonyl)-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)benzamide (17.4 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→1/0) was used.

[0481]

15 Example 93

N-(2-((3-(5-(3-fluoropyridin-2-yl)-2-methoxyphenylsulfonamido)phenyl)amino)ethyl)-3-methoxybenzamide

[0482]



20

[0483]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-5-(3-fluoropyridin-2-yl)-2-methoxybenzenesulfonamide trihydrochloride (16.7 mg) was amidated with m-anisic acid (5.5 mg) to give N-(2-((3-(5-(3-fluoropyridin-2-yl)-2-methoxyphenylsulfonamido)phenyl)amino)ethyl)-3-methoxybenzamide (12.2 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1→1/0) was used.

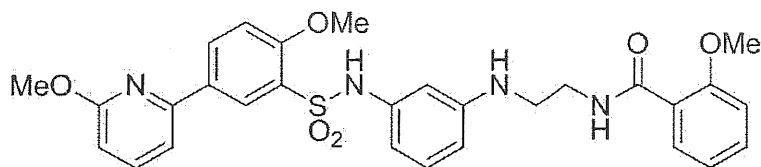
30

[0484]

Example 94

2-methoxy-N-(2-((3-(2-methoxy-5-(6-methoxypyridin-2-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide

35 [0485]



[0486]

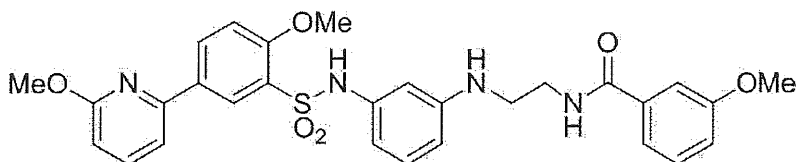
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(6-methoxypyridin-2-yl)benzenesulfonamide (27.0 mg) was amidated with o-anisic acid (9.6 mg) to give 2-methoxy-N-(2-((3-(2-methoxy-5-(6-methoxypyridin-2-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide (28.4 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=3/1) was used.

[0487]

Example 95

3-methoxy-N-(2-((3-(2-methoxy-5-(6-methoxypyridin-2-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide

[0488]



[0489]

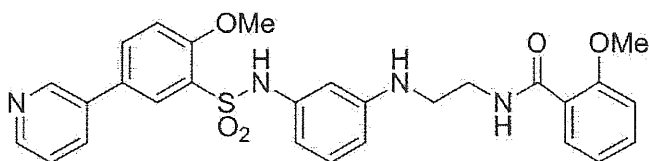
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(6-methoxypyridin-2-yl)benzenesulfonamide (27.0 mg) was amidated with m-anisic acid (9.6 mg) to give 3-methoxy-N-(2-((3-(2-methoxy-5-(6-methoxypyridin-2-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide (32.3 mg). For purification, silica gel column chromatography (eluate: ethyl acetate/hexane=1/1→3/1) was used.

[0490]

Example 96

2-methoxy-N-(2-((3-(2-methoxy-5-(pyridin-3-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide

[0491]



[0492]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(pyridin-3-

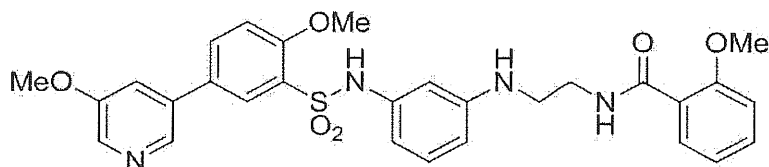
yl)benzenesulfonamide (14.0 mg) was amidated with o-anisic acid (5.3 mg) to give 2-methoxy-N-(2-((3-(2-methoxy-5-(pyridin-3-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide (13.5 mg). For purification, silica gel column chromatography
5 (eluate: ethyl acetate/chloroform =9/1) was used.

[0493]

Example 97

2-methoxy-N-(2-((3-(2-methoxy-5-(5-methoxypyridin-3-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide

10 [0494]



[0495]

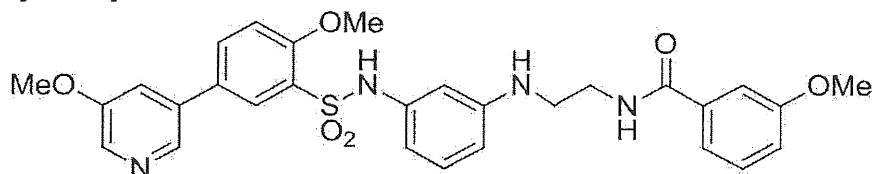
In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(5-methoxypyridin-3-yl)benzenesulfonamide (17.0 mg) was amidated with o-anisic acid (6.1 mg) to give 2-methoxy-N-(2-((3-(2-methoxy-5-(5-methoxypyridin-3-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide (19.0 mg). For purification, silica gel column chromatography (eluate:
20 ethyl acetate) was used.

[0496]

Example 98

3-methoxy-N-(2-((3-(2-methoxy-5-(5-methoxypyridin-3-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide

25 [0497]



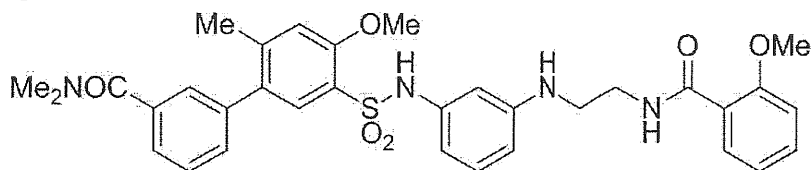
[0498]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-2-methoxy-5-(5-methoxypyridin-3-yl)benzenesulfonamide (17.0 mg) was amidated with m-anisic acid (6.1 mg) to give 3-methoxy-N-(2-((3-(2-methoxy-5-(5-methoxypyridin-3-yl)phenylsulfonamido)phenyl)amino)ethyl)benzamide (18.3 mg). For purification, silica gel column chromatography (eluate:
35 ethyl acetate) was used.

[0499]

Example 99

4'-methoxy-5'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide
[0500]



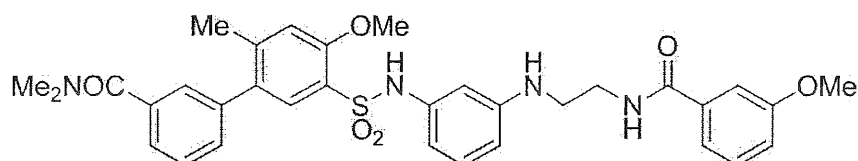
[0501]

In the same manner as in Example 1, 5'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.4 mg) was amidated with o-anisic acid (5.3 mg) to give 4'-methoxy-5'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide (9.3 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0502]

Example 100

4'-methoxy-5'-(N-(3-((2-(3-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide
[0503]



[0504]

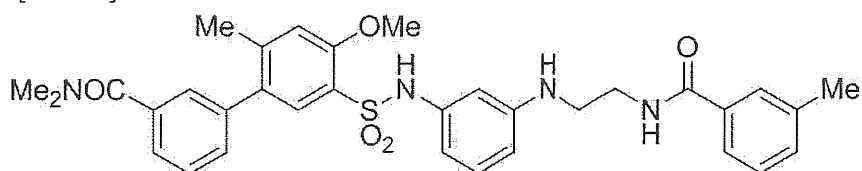
In the same manner as in Example 1, 5'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (27.0 mg) was amidated with m-anisic acid (7.5 mg) to give 4'-methoxy-5'-(N-(3-((2-(3-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide (19.5 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0505]

Example 101

4'-methoxy-N,N,2'-trimethyl-5'-(N-(3-((2-(3-methylbenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxamide

[0506]



[0507]

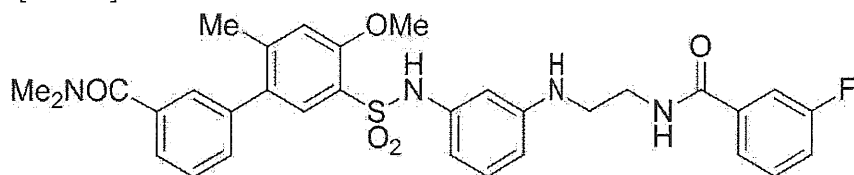
In the same manner as in Example 1, 5'-(N-(3-((2-
5 aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-
trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.4
mg) was amidated with 3-toluic acid (4.8 mg) to give 4'-
methoxy-N,N,2'-trimethyl-5'-(N-(3-((2-(3-
methylbenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-
10 3-carboxamide (16.6 mg). For purification, silica gel column
chromatography (eluate: ethyl acetate) was used.

[0508]

Example 102

5'-(N-(3-((2-(3-fluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-
15 4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide

[0509]



[0510]

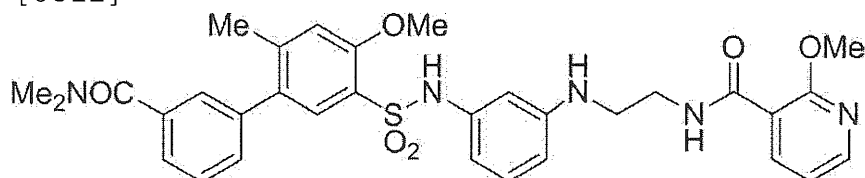
In the same manner as in Example 1, 5'-(N-(3-((2-
20 aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-
trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.4
mg) was amidated with 3-fluorobenzoic acid (4.9 mg) to give
5'-(N-(3-((2-(3-fluorobenzamido)ethyl)amino)phenyl)sulfamoyl)-
4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide
25 (22.0 mg). For purification, silica gel column chromatography
(eluate: ethyl acetate) was used.

[0511]

Example 103

N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-6-methyl-[1,1'-
30 biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-2-
methoxynicotinamide

[0512]



[0513]

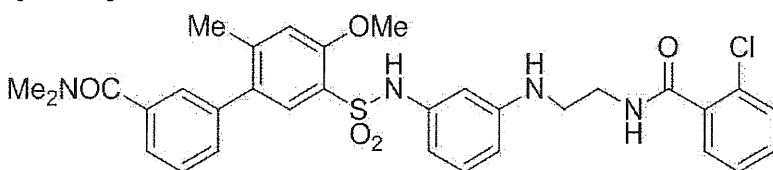
In the same manner as in Example 1, 5'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.4 mg) was amidated with 2-methoxypyridine-3-carboxylic acid (5.4 mg) to give N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-6-methyl-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)-2-methoxynicotinamide (21.1 mg). For purification, amine silica gel column chromatography (eluate: chloroform/ethyl acetate=1/1) was used.

10 [0514]

Example 104

5'-(N-(3-((2-(2-chlorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide
[0515]

15



[0516]

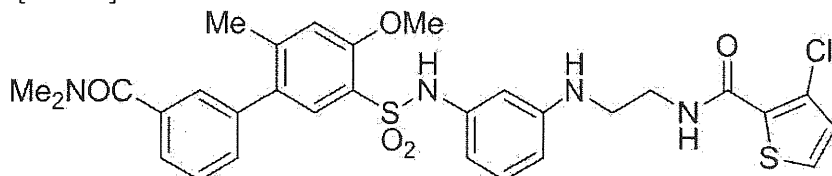
In the same manner as in Example 1, 5'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.4 mg) was amidated with 2-chlorobenzoic acid (5.5 mg) to give 5'-(N-(3-((2-(2-chlorobenzamido)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide (17.4 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

25 [0517]

Example 105

3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-methoxy-6-methyl-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)thiophene-2-carboxamide

30 [0518]



[0519]

In the same manner as in Example 1, 5'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-N,N,2'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.4 mg) was amidated with 3-chlorothiophene-2-carboxylic acid (5.7 mg) to give 3-chloro-N-(2-((3-(3'-(dimethylcarbamoyl)-4-

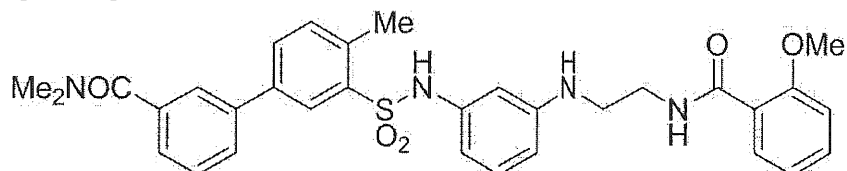
methoxy-6-methyl-[1,1'-biphenyl]-3-ylsulfonamido)phenyl)amino)ethyl)thiophene-2-carboxamide (20.4 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

5 [0520]

Example 106

3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N,4'-trimethyl-[1,1'-biphenyl]-3-carboxamide

10 [0521]



[0522]

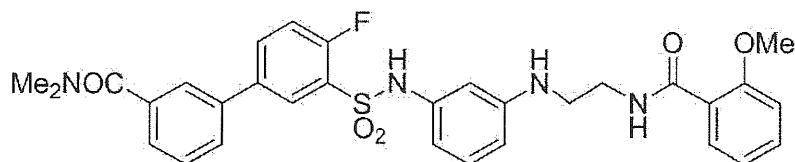
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-N,N,4'-trimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (18.4 mg) was amidated with o-anisic acid (5.3 mg) to give 3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N,4'-trimethyl-[1,1'-biphenyl]-3-carboxamide (20.3 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0523]

Example 107

4'-fluoro-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

25 [0524]



[0525]

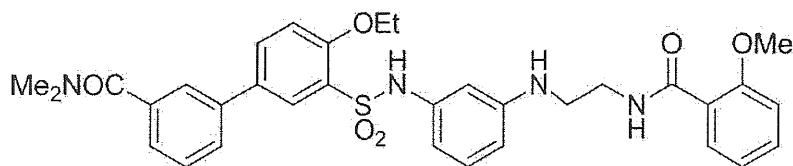
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-fluoro-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (18.5 mg) was amidated with o-anisic acid (5.3 mg) to give 4'-fluoro-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (20.1 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was used.

[0526]

Example 108

4'-ethoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

5 [0527]



[0528]

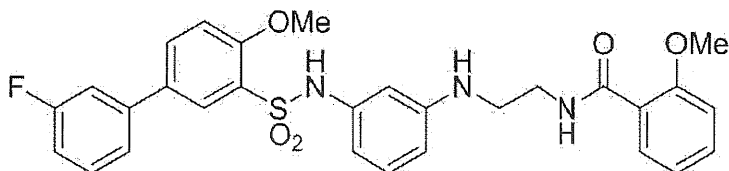
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-ethoxy-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.4 mg) was
10 amidated with o-anisic acid (5.3 mg) to give 4'-ethoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (15.5 mg). For purification, silica gel column chromatography (eluate: ethyl
15 acetate) was used.

[0529]

Example 109

N-(2-((3-(3'-fluoro-4-methoxy-[1,1'-biphenyl]-3-yl)sulfonamido)phenyl)amino)ethyl)-2-methoxybenzamide

20 [0530]



[0531]

In the same manner as in Example 1, N-(3-((2-aminoethyl)amino)phenyl)-3'-fluoro-4-methoxy-[1,1'-biphenyl]-
25 3-sulfonamide dihydrochloride (26.9 mg) was amidated with o-anisic acid (8.4 mg) to give N-(2-((3-(3'-fluoro-4-methoxy-[1,1'-biphenyl]-3-yl)sulfonamido)phenyl)amino)ethyl)-2-methoxybenzamide (13.0 mg). For purification, silica gel column chromatography (eluate: ammonia-saturated chloroform)
30 was used.

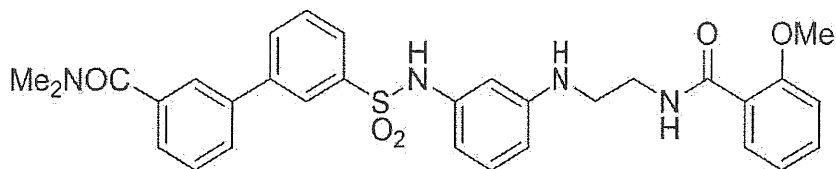
[0532]

Example 110

3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide

35 [0533]

[0533]



[0534]

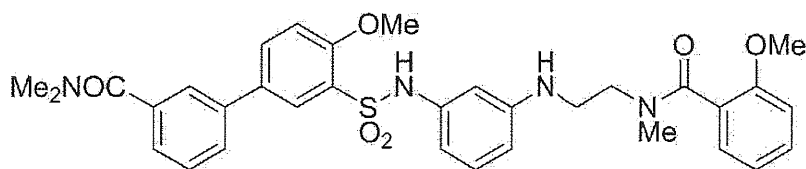
In the same manner as in Example 1, 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-
 5 biphenyl]-3-carboxamide dihydrochloride (17.9 mg) was amidated with o-anisic acid (5.3 mg) to give 3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (16.9 mg). For purification, silica gel column chromatography (eluate: ethyl acetate) was
 10 used.

[0535]

Example 111

4'-methoxy-3'-(N-(3-((2-(2-methoxy-N-methylbenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-
 15 [1,1'-biphenyl]-3-carboxamide

[0536]



[0537]

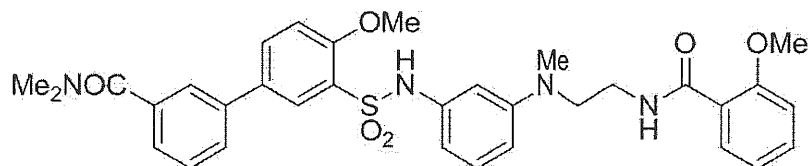
In the same manner as in Example 1, 4'-methoxy-N,N-dimethyl-3'-(N-(3-((2-methylamino)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxamide dihydrochloride (19.4 mg) was amidated with o-anisic acid (5.3 mg) to give 4'-methoxy-3'-(N-(3-((2-(2-methoxy-N-methylbenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (19.3 mg). For
 25 purification, preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=50/1) was used.

[0538]

Example 112

4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)(methyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide
 30

[0539]



To a solution of 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (17.2 mg) in acetonitrile (1.0 mL) were added 37% formalin aqueous solution (0.035 mL), acetic acid (0.007 mL) and sodium cyanoborohydride (8.2 mg), and the mixture was stirred at room temperature for 2 days. To this reaction mixture was added saturated aqueous sodium hydrogen carbonate solution, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated. The obtained residue was purified by preparative thin layer chromatography (eluate: ammonia-saturated chloroform/methanol=80/1) to give 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)(methyl)amino)phenyl)sulfamoyl)-N,N-dimethyl-[1,1'-biphenyl]-3-carboxamide (12.8 mg).

Example 113

4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid

COc1ccc(cc1)C(=O)NCCNC2=CC=C(N2)NS(=O)(=O)c3ccc(cc3C(=O)O)C4=CC=C(C=C4)OC

(1) To methyl 3'-(N-(3-((2-((tert-butoxycarbonyl)amino)ethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylate was added 10% hydrogen chloride-methanol (6.0 mL) solution, and the mixture was stirred at 50°C for 2 hr. The reaction mixture was concentrated to give methyl 3'-(N-(3-((2-aminoethyl)amino)phenyl)sulfamoyl)-4'-methoxy-[1,1'-biphenyl]-3-carboxylate dihydrochloride (356 mg).

124

concentrated. The obtained residue was purified by silica gel column chromatography (eluate: ethyl acetate/hexane=2/1→1/0) to give methyl 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylate (176 mg).

(3) To a suspension of methyl 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylate (176 mg) in methanol (3.0 mL) was added 1N aqueous sodium hydroxide solution (0.900 mL), and the mixture was stirred at 60°C for 3 hr. To this reaction mixture was added 1N hydrochloric acid, and the mixture was extracted with chloroform. The organic layer was dried over sodium sulfate, filtered, and the filtrate was concentrated to give 4'-methoxy-3'-(N-(3-((2-(2-methoxybenzamido)ethyl)amino)phenyl)sulfamoyl)-[1,1'-biphenyl]-3-carboxylic acid (161 mg).

[0544]

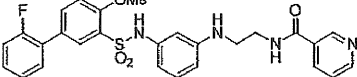
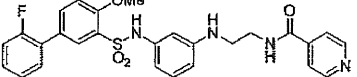
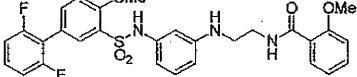
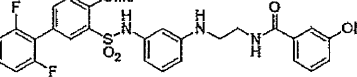
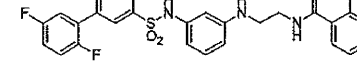
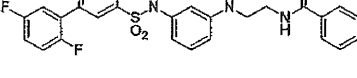
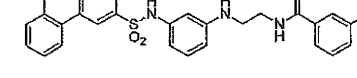
The structural formulas and physical property data of the Example compounds are shown in Table 9 to Table 24.

Example No.	Structure	ESI-MS (M+H) ⁺	¹ H-NMR Chemical Shift (400MHz, CDCl ₃)
1		δ 3.31 (2H, t, J = 6.0 Hz), 3.62-3.65 (2H, m), 3.81 (3H, s), 4.06 (3H, s), 4.25 (1H, bs), 6.31-6.36 (2H, m), 6.48 (1H, t, J = 7.6, 2.0 Hz), 7.25-7.37 (2H, m), 7.44 (1H, td, J = 7.6, 2.0 Hz), 7.68 (1H, td, J = 8.4, 2.0 Hz), 8.03 (1H, dd, J = 8.4, 2.0 Hz), 8.08-8.16 (1H, m), 8.20 (1H, dd, J = 8.0, 2.0 Hz)	
2		δ 3.31 (2H, t, J = 5.6 Hz), 3.62-3.65 (2H, m), 3.83 (3H, s), 4.06 (3H, s), 6.31-6.36 (4H, m), 6.50 (1H, t, J = 2.4 Hz), 6.88 (1H, bs), 6.97 (1H, t, J = 8.0 Hz), 7.01-7.08 (3H, m), 7.17-7.22 (2H, m), 7.26-7.35 (4H, m), 7.68 (1H, td, J = 9.2, 2.0 Hz), 8.04 (1H, d, J = 0.8 Hz)	
3		δ 3.30 (2H, t, J = 6.0 Hz), 3.59-3.65 (2H, m), 3.84 (3H, s), 4.06 (3H, s), 4.15 (1H, bs), 6.32-6.48 (3H, m), 6.49 (1H, t, J = 2.0 Hz), 6.88-6.90 (3H, m), 6.96 (1H, t, J = 8.0 Hz), 7.04-7.11 (2H, m), 7.17 (1H, t, J = 7.2 Hz), 7.28-7.35 (2H, m), 7.67-7.69 (3H, m), 8.04 (1H, dd, J = 4.8, 0.8 Hz)	
4		δ 3.29-3.32 (2H, m), 3.60-3.65 (2H, m), 3.78 (3H, s), 4.07 (3H, s), 4.18 (1H, bs), 6.30-6.35 (2H, m), 6.50 (1H, t, J = 2.4 Hz), 6.79 (1H, d, J = 8.8 Hz), 6.89 (1H, bs), 6.96 (1H, t, J = 8.0 Hz), 7.04-7.18 (3H, m), 7.26-7.34 (2H, m), 7.52 (1H, dd, J = 8.0 Hz), 7.69 (1H, td, J = 8.4, 2.0 Hz), 8.02-8.03 (2H, m), 8.29 (1H, d, J = 4.2 Hz)	
5		δ 1.73 (1H, bs), 2.97 (6H, s), 3.30 (2H, t, J = 6.0 Hz), 3.61-3.66 (2H, m), 3.76 (3H, s), 4.06 (3H, s), 6.32-6.36 (2H, m), 6.49 (1H, t, J = 8.0 Hz), 7.00-7.18 (4H, m), 7.26-7.37 (2H, m), 7.68 (1H, td, J = 8.8, 2.0 Hz), 7.73-7.75 (1H, m), 8.02 (1H, d, J = 0.6 Hz), 8.22-8.25 (1H, m)	
6		δ 3.34 (2H, t, J = 6.0 Hz), 3.59-3.64 (2H, m), 4.00 (1H, bs), 4.06 (3H, s), 4.25 (1H, bs), 6.31-6.36 (2H, m), 6.53 (1H, t, J = 2.0 Hz), 6.58 (1H, bs), 6.77 (1H, t, J = 8.0 Hz), 7.27-7.40 (4H, m), 7.68 (1H, td, J = 8.8, 2.4 Hz), 8.04 (1H, d, J = 1.2 Hz), 12.21 (1H, bs)	
7		δ 3.33 (2H, t, J = 6.0 Hz), 3.61-3.66 (2H, m), 4.07 (3H, s), 4.09 (1H, bs), 6.32-6.36 (3H, m), 6.49 (1H, t, J = 2.4 Hz), 6.89 (1H, bs), 6.97 (1H, t, J = 8.0 Hz), 7.05-7.19 (3H, m), 7.26-7.39 (2H, m), 7.59 (1H, td, J = 8.0, 1.2 Hz), 7.68 (1H, td, J = 10.8, 2.0 Hz), 8.04 (1H, dd, J = 2.4, 0.8 Hz)	

8		554	δ 3.32 (2H, t, J = 6.0 Hz), 3.60-3.64 (2H, m), 4.06 (3H, s), 4.09 (1H, bs), 6.30-6.36 (3H, m), 6.52 (1H, t, J = 2.0 Hz), 6.87 (1H, bs), 6.97 (1H, t, J = 8.0 Hz), 7.05-7.19 (3H, m), 7.26-7.36 (3H, m), 7.45 (1H, ddd, J = 8.0, 2.0, 0.8 Hz), 7.56 (1H, dt, J = 5.6, 1.2 Hz), 7.67-7.70 (2H, m), 8.04 (1H, dd, J = 2.0, 0.8 Hz)
9		554	δ 3.32 (2H, t, J = 6.0 Hz), 3.60-3.64 (2H, m), 4.07 (3H, s), 4.09 (1H, bs), 6.30-6.36 (3H, m), 6.49 (1H, t, J = 2.0 Hz), 6.89 (1H, bs), 6.97 (1H, t, J = 8.0 Hz), 7.05-7.19 (3H, m), 7.26-7.38 (3H, m), 7.62-7.65 (2H, m), 7.68 (1H, dt, J = 8.8, 2.0 Hz), 8.04 (1H, dd, J = 2.0, 0.8 Hz)
10		566	δ 3.31 (2H, t, J = 6.0 Hz), 3.62-3.67 (2H, m), 3.83 (3H, s), 4.06 (3H, s), 4.25 (1H, bs), 6.31 (1H, dd, J = 8.0, 1.6 Hz), 6.35 (1H, dt, J = 8.0, 1.6 Hz), 6.47 (1H, t, J = 2.4 Hz), 6.83 (1H, bs), 6.92-7.10 (4H, m), 7.21-7.26 (3H, m), 7.37-7.39 (1H, m), 7.44 (1H, dt, J = 8.4, 2.0 Hz), 7.55 (1H, dd, J = 8.4, 2.0 Hz), 7.92 (1H, d, J = 2.4 Hz), 8.08-8.12 (1H, m), 8.20 (1H, dd, J = 8.0, 2.0 Hz)
11		566	δ 3.33 (2H, t, J = 6.0 Hz), 3.64-3.69 (2H, m), 3.84 (3H, s), 4.06 (3H, s), 4.24 (1H, bs), 6.32 (1H, dd, J = 7.2, 1.6 Hz), 6.41-6.44 (3H, m), 6.54 (1H, bs), 7.00-7.06 (2H, m), 7.24-7.35 (7H, m), 7.63 (1H, ddd, J = 8.0, 2.0, 1.2 Hz), 7.94 (1H, d, J = 2.0 Hz), 8.0, 2.0, 0.8 Hz)
12		562	δ 3.31 (2H, t, J = 6.0 Hz), 3.61-3.65 (2H, m), 3.81 (6H, s), 4.08 (3H, s), 6.34 (1H, d, J = 8.4 Hz), 6.47-6.48 (1H, m), 6.80-6.84 (1H, m), 6.90-7.10 (6H, m), 7.21-7.31 (4H, m), 7.44 (1H, dt, J = 7.6, 2.0 Hz), 7.66 (1H, dd, J = 8.4, 2.0 Hz), 8.03 (1H, dd, J = 5.6, 2.4 Hz), 8.06-8.13 (1H, m), 8.18 (1H, dd, J = 8.0, 1.6 Hz)
13		562	δ 3.31 (2H, t, J = 6.0 Hz), 3.61-3.65 (2H, m), 3.83 (6H, s), 4.08 (3H, s), 4.25 (1H, bs), 6.33-6.36 (3H, m), 6.47-6.49 (1H, m), 6.81-6.86 (2H, m), 6.92-7.08 (5H, m), 7.20-7.23 (1H, m), 7.26-7.32 (3H, m), 7.67 (1H, dd, J = 8.8, 2.4 Hz), 8.02 (1H, dd, J = 7.2, 2.0 Hz)
14		521	δ 3.31 (2H, t, J = 6.0 Hz), 3.63-3.68 (2H, m), 4.06 (3H, s), 6.33-6.36 (2H, m), 6.47 (1H, t, J = 2.0 Hz), 6.86 (1H, bs), 6.96 (1H, t, J = 8.0 Hz), 7.05-7.19 (3H, m), 7.26-7.29 (2H, m), 7.34 (1H, dt, J = 9.2, 1.6 Hz), 7.43 (1H, ddd, J = 8.0, 4.8, 1.2 Hz), 7.68 (1H, dt, J = 8.8, 2.0 Hz), 7.85 (1H, dt, J = 9.6, 2.0 Hz), 8.04 (1H, t, J = 1.6 Hz), 8.18 (1H, d, J = 7.6 Hz), 8.20-8.28 (1H, m), 8.52 (1H, ddd, J = 4.0, 1.6, 0.8 Hz)

[0547]

Table 11

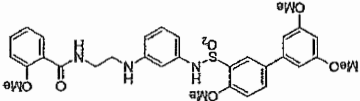
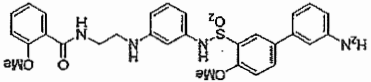
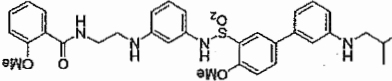
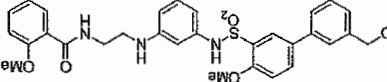
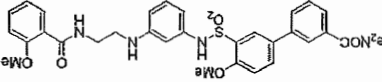
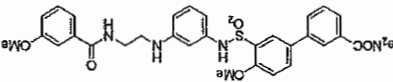
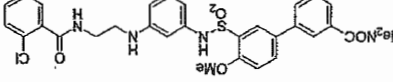
15		521	δ 3.34 (2H, t, $J = 6.0$ Hz), 3.63-3.70 (2H, m), 4.06 (3H, s), 4.25 (1H, brs), 6.30-6.36 (2H, m), 6.41-6.50 (1H, m), 6.54 (1H, t, $J = 2.0$ Hz), 6.93-6.99 (2H, m), 7.05-7.19 (3H, m), 7.26-7.36 (3H, m), 7.68 (1H, dt, $J = 8.8, 2.0$ Hz), 8.02-8.05 (2H, m), 8.71 (1H, dd, $J = 4.8, 2.0$ Hz), 8.90 (1H, d, $J = 1.6$ Hz)
16		521	δ 3.34 (2H, t, $J = 6.0$ Hz), 3.62-3.66 (2H, m), 4.02 (1H, brs), 4.06 (3H, s), 6.29 (1H, dd, $J = 8.4, 2.0$ Hz), 6.34 (1H, dd, $J = 8.0, 2.0$ Hz), 6.46-6.53 (1H, m), 6.56 (1H, t, $J = 2.0$ Hz), 6.92-6.99 (2H, m), 7.05-7.19 (3H, m), 7.26-7.36 (2H, m), 7.52-7.53 (2H, m), 7.68 (1H, dt, $J = 8.8, 2.0$ Hz), 8.04 (1H, t, $J = 0.8$ Hz), 8.68-8.70 (2H, m)
17		568	δ 3.31 (2H, t, $J = 6.0$ Hz), 3.62-3.67 (2H, m), 3.81 (3H, s), 4.05 (3H, s), 4.20 (1H, brs), 6.30-6.36 (2H, m), 6.48 (1H, t, $J = 2.0$ Hz), 6.84 (1H, brs), 6.89-7.00 (3H, m), 7.04-7.10 (2H, m), 7.22-7.28 (2H, m), 7.42-7.47 (1H, m), 7.58 (1H, dt, $J = 8.8, 1.2$ Hz), 7.99-8.01 (1H, m), 8.06-8.14 (1H, m), 8.19 (1H, dd, $J = 8.0, 2.0$ Hz)
18		568	δ 3.30 (2H, t, $J = 6.0$ Hz), 3.60-3.65 (2H, m), 3.83 (3H, s), 4.07 (3H, s), 4.15 (1H, brs), 6.30-6.35 (2H, m), 6.37-6.43 (1H, m), 6.49 (1H, t, $J = 2.0$ Hz), 6.70-7.07 (6H, m), 7.20-7.32 (4H, m), 7.57-7.60 (1H, m), 8.00 (1H, dd, $J = 2.4, 1.2$ Hz)
19		568	δ 3.31 (2H, t, $J = 6.0$ Hz), 3.63-3.69 (2H, m), 3.84 (3H, s), 4.07 (3H, s), 4.25 (1H, brs), 6.31-6.37 (2H, m), 6.48 (1H, t, $J = 2.0$ Hz), 6.86 (1H, brs), 6.93-6.99 (3H, m), 7.02-7.10 (4H, m), 7.44 (1H, td, $J = 6.4, 1.2$ Hz), 7.65 (1H, dt, $J = 8.4, 2.0$ Hz), 8.01 (1H, dd, $J = 2.0, 0.8$ Hz), 8.07-8.13 (1H, m), 8.19 (1H, dd, $J = 8.0, 1.6$ Hz)
20		568	δ 3.31 (2H, t, $J = 6.0$ Hz), 3.61-3.65 (2H, m), 3.83 (3H, s), 4.07 (3H, s), 4.20 (1H, brs), 6.31-6.36 (2H, m), 6.37-6.44 (1H, m), 6.49 (1H, t, $J = 2.0$ Hz), 6.91 (1H, brs), 6.95-7.07 (5H, m), 7.20-7.32 (4H, m), 7.65 (1H, dt, $J = 8.4, 2.0$ Hz), 8.01 (1H, dd, $J = 2.0, 1.2$ Hz)
21		546	δ 2.91 (3H, s), 3.50-3.53 (2H, m), 3.56-3.59 (2H, m), 3.83 (3H, s), 4.02 (3H, s), 4.15 (1H, brs), 6.29-6.32 (2H, m), 6.42-6.49 (1H, m), 6.59 (1H, brs), 6.89 (1H, brs), 6.94-7.13 (4H, m), 7.13-7.14 (1H, m), 7.26-7.28 (5H, m), 7.45 (1H, td, $J = 7.2, 1.6$ Hz), 7.80 (1H, dd, $J = 8.0, 1.6$ Hz)

22		548 δ 3.28 (2H, t, J = 6.0 Hz), 3.66-3.70 (2H, m), 3.82 (3H, s), 4.10 (3H, s), 4.13 (1H, bs), 6.17 (1H, dd, J = 8.0, 1.6 Hz), 6.23 (1H, dd, J = 8.0, 1.6 Hz), 6.66 (1H, t, J = 2.0 Hz), 6.85 (1H, bs), 6.91-7.09 (5H, m), 7.18-7.35 (5H, m), 7.46-7.52 (1H, m), 7.55 (1H, dd, J = 8.4, 2.4 Hz), 8.32 (1H, d, J = 2.4 Hz)
23		580 δ 3.31 (2H, t, J = 6.0 Hz), 3.60-3.65 (2H, m), 3.82 (6H, s), 4.09 (3H, s), 4.11-4.15 (1H, m), 6.34 (1H, d, J = 8.0 Hz), 6.47-6.48 (1H, m), 6.80-6.86 (1H, m), 6.91-7.11 (4H, m), 7.21-7.31 (4H, m), 7.44 (1H, td, J = 7.6, 2.0 Hz), 7.68 (1H, dd, J = 8.4, 2.0 Hz), 8.06 (1H, dd, J = 5.6, 2.0 Hz), 8.08-8.15 (1H, m), 8.20 (1H, dd, J = 8.0, 1.2 Hz)
24		580 δ 3.32 (2H, t, J = 6.0 Hz), 3.60-3.66 (2H, m), 3.81 (6H, s), 4.08 (3H, s), 4.20 (1H, bs), 6.33-6.36 (3H, m), 6.47-6.49 (1H, m), 6.80-6.84 (2H, m), 6.90-7.05 (4H, m), 7.18-7.20 (1H, m), 7.24-7.30 (3H, m), 7.70 (1H, dd, J = 8.8, 2.4 Hz), 8.02 (1H, dd, J = 7.2, 2.0 Hz)
25		600 δ 3.31 (2H, t, J = 6.0 Hz), 3.63-3.68 (2H, m), 3.86 (3H, s), 4.07 (3H, s), 4.23 (1H, bs), 6.30 (1H, dd, J = 8.0, 1.6 Hz), 6.35 (1H, dd, J = 8.0, 1.6 Hz), 6.50 (1H, t, J = 2.0 Hz), 6.86 (1H, bs), 6.93-7.11 (4H, m), 7.20-7.32 (3H, m), 7.43-7.47 (1H, m), 7.52 (1H, dd, J = 8.4, 2.0 Hz), 7.90 (1H, d, J = 8.4, 2.0 Hz), 8.13 (1H, m), 8.19 (1H, dd, J = 8.4, 2.0 Hz)
26		600 δ 3.31 (2H, t, J = 6.0 Hz), 3.62-3.68 (2H, m), 3.84 (3H, s), 4.08 (3H, s), 4.13 (1H, bs), 6.29-6.36 (2H, m), 6.37-6.44 (1H, m), 6.49 (1H, t, J = 2.0 Hz), 6.88 (1H, bs), 6.96 (1H, t, J = 8.0 Hz), 7.02-7.05 (2H, m), 7.21-7.33 (6H, m), 7.52 (1H, dd, J = 8.4, 2.4 Hz), 7.90 (1H, d, J = 2.0 Hz)
27		600 δ 3.30 (2H, t, J = 6.0 Hz), 3.62-3.68 (2H, m), 3.84 (3H, s), 4.08 (3H, s), 4.28 (1H, bs), 6.31-6.37 (2H, m), 6.45-6.47 (1H, m), 6.65-6.68 (1H, m), 6.84 (1H, bs), 6.93-7.10 (4H, m), 7.29-7.35 (2H, m), 7.45 (1H, td, J = 8.0, 2.0 Hz), 7.61-7.65 (1H, m), 7.99 (1H, d, J = 2.4 Hz), 8.06-8.12 (1H, m), 8.17-8.21 (1H, m)
28		600 δ 3.32-3.35 (2H, m), 3.64-3.68 (2H, m), 3.85 (3H, s), 4.08 (3H, s), 4.18 (1H, bs), 6.31-6.37 (3H, m), 6.48 (1H, t, J = 2.0 Hz), 6.87 (1H, bs), 6.95-7.08 (3H, m), 7.20-7.35 (6H, m), 7.63 (1H, dd, J = 8.8, 2.4 Hz), 8.00 (1H, d, J = 2.4 Hz)

[0549]

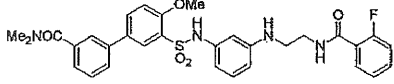
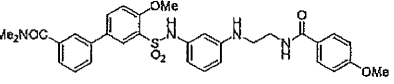
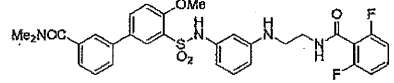
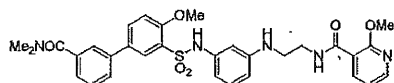
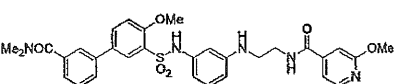
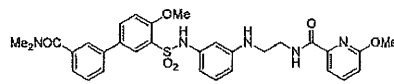
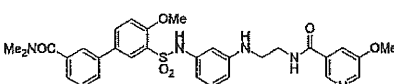
Table 13

29		566	δ 3.31 (2H, t, $J = 6.0$ Hz), 3.61-3.67 (2H, m), 3.84 (3H, s), 4.06 (3H, s), 4.18 (1H, brs), 6.32-6.36 (3H, m), 6.48 (1H, t, $J = 2.0$ Hz), 6.88 (1H, brs), 6.94-7.07 (3H, m), 7.19-7.37 (6H, m), 7.46 (1H, t, $J = 1.6$ Hz), 7.65 (1H, dd, $J = 8.8, 2.4$ Hz), 8.03 (1H, d, $J = 2.4$ Hz)
30		562	δ 3.30 (2H, t, $J = 6.0$ Hz), 3.62-3.66 (2H, m), 3.80 (3H, s), 3.88 (3H, s), 4.06 (3H, s), 4.15 (1H, brs), 6.32-6.36 (2H, m), 6.48 (1H, t, $J = 2.0$ Hz), 6.84-7.10 (7H, m), 7.28-7.32 (2H, m), 7.42-7.46 (1H, m), 7.66 (1H, dd, $J = 8.4, 2.4$ Hz), 8.05 (1H, d, $J = 2.4$ Hz), 8.06-8.12 (1H, m), 8.18 (1H, dd, $J = 7.6, 2.0$ Hz)
31		562	δ 3.30 (2H, t, $J = 6.0$ Hz), 3.59-3.64 (2H, m), 3.81 (3H, s), 3.88 (3H, s), 4.06 (3H, s), 4.16 (1H, brs), 6.31-6.35 (2H, m), 6.36-6.41 (1H, m), 6.48-6.50 (1H, m), 6.85-7.08 (7H, m), 7.18-7.34 (4H, m), 7.67 (1H, dd, $J = 8.8, 2.4$ Hz), 8.06 (1H, d, $J = 2.4$ Hz)
32		584	δ 3.31 (2H, t, $J = 6.0$ Hz), 3.63-3.68 (2H, m), 3.86 (3H, s), 4.07 (3H, s), 4.25 (1H, brs), 6.33-6.36 (2H, m), 6.47 (1H, t, $J = 2.4$ Hz), 6.86 (1H, brs), 6.94-6.98 (3H, m), 7.03-7.11 (3H, m), 7.32-7.35 (1H, m), 7.43-7.47 (1H, m), 7.53 (1H, dd, $J = 8.4, 2.4$ Hz), 7.91 (1H, d, $J = 2.4$ Hz), 8.07-8.13 (1H, m), 8.20 (1H, dd, $J = 8.0, 2.0$ Hz)
33		584	δ 3.32 (2H, t, $J = 6.0$ Hz), 3.62-3.68 (2H, m), 3.84 (3H, s), 4.07 (3H, s), 4.15 (1H, brs), 6.30-6.36 (2H, m), 6.36-6.44 (1H, m), 6.49 (1H, t, $J = 2.0$ Hz), 6.85-6.86 (1H, m), 6.94-7.04 (5H, m), 7.06-7.35 (4H, m), 7.52-7.55 (1H, m), 7.91 (1H, d, $J = 2.4$ Hz)
34		580	δ 3.31 (2H, t, $J = 6.0$ Hz), 3.62-3.66 (2H, m), 3.79 (3H, s), 3.87 (3H, s), 4.06 (3H, s), 4.24 (1H, brs), 6.31-6.36 (2H, m), 6.45-6.49 (1H, m), 6.77-6.85 (3H, m), 6.91-7.11 (5H, m), 7.42-7.48 (1H, m), 7.66 (1H, dt, $J = 8.4, 2.0$ Hz), 8.01-8.03 (1H, m), 8.06-8.13 (1H, m), 8.10-8.20 (1H, m)
35		580	δ 3.31 (2H, t, $J = 6.0$ Hz), 3.60-3.65 (2H, m), 3.79 (3H, s), 3.84 (3H, s), 4.05 (3H, s), 4.25 (1H, brs), 6.33-6.35 (2H, m), 6.39-6.46 (1H, brs), 6.50 (1H, t, $J = 2.0$ Hz), 6.78-6.85 (2H, m), 6.91-7.05 (5H, m), 7.20-7.34 (3H, m), 7.67 (1H, dt, $J = 8.4, 2.0$ Hz), 8.03 (1H, dd, $J = 2.4, 1.2$ Hz)

36		δ 2.97 (3H, s), 3.30 (2H, t, J = 6.0 Hz), 3.62-3.66 (2H, m), 3.80 (3H, s), 3.83 (3H, s), 4.06 (3H, s), 4.22 (1H, bs), 6.31-6.36 (2H, m), 6.46-6.48 (1H, m), 6.60-6.62 (1H, m), 6.86 (1H, bs), 7.42-7.46 (1H, td, J = 8.4, 2.0 Hz), 7.64-7.68 (1H, m), 8.04-8.06 (1H, m), 8.06-8.12 (1H, m), 8.18-8.20 (1H, m)
37		δ 3.30 (2H, t, J = 6.0 Hz), 3.62-3.67 (2H, m), 3.76 (2H, bs), 3.80 (3H, s), 4.07 (3H, s), 4.21 (1H, bs), 6.29-6.35 (2H, m), 6.49 (1H, t, J = 2.0 Hz), 6.63 (1H, ddd, J = 8.0, 2.4, 0.8 Hz), 6.80-7.10 (7H, m), 7.16 (1H, t, J = 8.0 Hz), 7.44 (1H, ddd, J = 8.8, 5.2, 1.6 Hz), 7.66 (1H, dd, J = 8.8, 2.4 Hz), 8.03 (1H, d, J = 2.4 Hz), 8.07-8.10 (1H, m), 8.19 (1H, dd, J = 8.0, 2.0 Hz)
38		δ 0.98 (6H, d, J = 6.8 Hz), 1.86-1.90 (1H, m), 2.94 (2H, d, J = 6.8 Hz), 3.30 (2H, t, J = 6.0 Hz), 3.61-3.66 (2H, m), 3.78 (3H, s), 4.05 (3H, s), 4.18 (1H, bs), 6.31-6.35 (2H, m), 6.47 (1H, t, J = 2.04 Hz), 6.56 (1H, ddd, J = 8.0, 2.4, 0.8 Hz), 6.69 (1H, t, J = 2.0 Hz), 6.76 (1H, dd, J = 8.4, 1.6 Hz), 6.88-6.96 (4H, m), 7.01-7.09 (2H, m), 7.17 (1H, t, J = 8.0 Hz), 7.41-7.46 (1H, m), 7.66 (1H, dd, J = 8.8, 2.4 Hz), 8.04 (1H, d, J = 2.4 Hz), 8.12 (1H, m), 8.19 (1H, dd, J = 8.0, 2.0 Hz)
39		δ 2.37 (2H, bs), 2.39 (1H, bs), 3.30-3.32 (2H, m), 3.61-3.66 (2H, m), 3.79 (3H, s), 4.23 (1H, bs), 6.32-6.35 (2H, m), 6.47 (1H, t, J = 2.0 Hz), 6.88-6.97 (3H, m), 7.03-7.14 (3H, m), 7.26-7.30 (3H, m), 7.44 (1H, ddd, J = 9.2, 7.2, 1.6 Hz), 7.66 (1H, dd, J = 8.8, 2.4 Hz), 8.05 (1H, d, J = 2.4 Hz), 8.11 (1H, m), 8.18 (1H, dd, J = 8.0, 2.0 Hz)
40		δ 2.98 (3H, bs), 3.12 (3H, bs), 3.30 (2H, t, J = 6.0 Hz), 3.63-3.67 (2H, m), 3.82 (3H, s), 4.07 (3H, s), 4.31 (1H, bs), 6.31-6.36 (2H, m), 6.47 (1H, t, J = 2.4 Hz), 6.88 (1H, bs), 6.92-6.97 (2H, m), 7.04-7.09 (2H, m), 7.33-7.46 (3H, m), 7.51-7.54 (2H, m), 7.68 (1H, dd, J = 8.8, 2.4 Hz), 8.05 (1H, d, J = 2.4 Hz), 8.08-8.14 (1H, m), 8.18 (1H, dd, J = 8.0, 2.0 Hz)
41		δ 3.02 (3H, bs), 3.13 (3H, bs), 3.21 (2H, t, J = 5.2 Hz), 3.58-3.63 (2H, m), 3.79 (3H, s), 4.04 (3H, s), 4.48 (1H, bs), 6.21 (1H, dd, J = 7.6, 1.6 Hz), 6.31 (1H, dd, J = 8.0, 1.6 Hz), 6.47 (1H, t, J = 8.0 Hz), 6.92 (1H, t, J = 2.4 Hz), 6.87 (1H, bs), 6.92 (1H, t, J = 8.0 Hz), 6.97-7.05 (2H, m), 7.18-7.28 (3H, m), 7.30-7.36 (2H, m), 7.42 (1H, t, J = 8.0 Hz), 8.11 (1H, d, J = 2.4 Hz), 8.4, 2.4 Hz)
42		δ 2.99 (3H, bs), 3.08 (3H, bs), 3.28 (2H, t, J = 6.0 Hz), 3.61-3.65 (2H, m), 4.06 (3H, s), 4.28 (1H, bs), 6.27 (1H, dd, J = 8.0, 1.6 Hz), 6.33 (1H, dd, J = 8.0 Hz), 6.47 (1H, t, J = 2.0 Hz), 6.85-6.87 (2H, m), 6.92 (1H, t, J = 8.0 Hz), 7.07 (1H, d, J = 8.8 Hz), 7.23-7.43 (5H, m), 7.50-7.55 (3H, m), 7.69 (1H, dd, J = 8.4, 2.4 Hz), 8.09 (1H, d, J = 2.4 Hz)

[0551]

Table 15

43		591	δ 2.99 (3H, brs), 3.12 (3H, brs), 3.29 (2H, t, $J = 6.0$ Hz), 3.64-3.67 (2H, m), 4.07 (3H, s), 4.24 (1H, brs), 6.31 (1H, dd, $J = 8.0, 1.6$ Hz), 6.35 (1H, dd, $J = 8.4, 2.0$ Hz), 6.46 (1H, t, $J = 2.0$ Hz), 6.84 (1H, brs), 6.96 (1H, t, $J = 8.0$ Hz), 7.05-7.11 (3H, m), 7.23-7.35 (2H, m), 7.40-7.47 (2H, m), 7.52-7.55 (2H, m), 7.68 (1H, dd, $J = 8.8, 2.0$ Hz), 8.02-8.08 (2H, m)
44		603	δ 3.02 (3H, brs), 3.14 (3H, brs), 3.20 (2H, t, $J = 5.6$ Hz), 3.57-3.62 (2H, m), 3.82 (3H, s), 3.86 (1H, brs), 4.04 (3H, s), 4.50 (1H, brs), 6.22 (1H, dd, $J = 8.0, 1.2$ Hz), 6.31 (1H, dd, $J = 8.0, 1.6$ Hz), 6.45 (1H, t, $J = 2.0$ Hz), 6.84-6.95 (4H, m), 7.04 (1H, d, $J = 8.8$ Hz), 7.09-7.17 (1H, m), 7.32-7.34 (1H, m), 7.43 (1H, t, $J = 7.6$ Hz), 7.54-7.57 (2H, m), 7.67-7.79 (2H, m), 8.12 (1H, d, $J = 2.4$ Hz)
45		609	δ 2.99 (3H, brs), 3.07 (3H, brs), 3.24-3.26 (2H, m), 3.58-3.62 (2H, m), 4.06 (3H, s), 4.17 (1H, brs), 6.28 (1H, dd, $J = 8.0, 1.6$ Hz), 6.33 (1H, dd, $J = 8.0, 1.6$ Hz), 6.45 (1H, t, $J = 2.0$ Hz), 6.85-6.97 (4H, m), 7.07 (1H, d, $J = 8.8$ Hz), 7.26-7.35 (3H, m), 7.41 (1H, t, $J = 7.6$ Hz), 7.53-7.55 (2H, m), 7.70 (1H, dd, $J = 8.4, 2.4$ Hz), 8.09 (1H, d, $J = 2.0$ Hz)
46		604	δ 2.99 (3H, brs), 3.13 (3H, brs), 3.31 (2H, t, $J = 6.0$ Hz), 3.63-3.73 (2H, m), 3.97 (3H, s), 4.07 (3H, s), 4.30 (1H, brs), 6.31 (1H, dd, $J = 8.0, 1.2$ Hz), 6.35 (1H, dd, $J = 8.0, 1.2$ Hz), 6.49 (1H, t, $J = 2.0$ Hz), 6.93-6.97 (2H, m), 7.03-7.07 (2H, m), 7.33-7.35 (1H, m), 7.42 (1H, t, $J = 7.6$ Hz), 7.51-7.55 (2H, m), 7.69 (1H, dd, $J = 8.4, 2.0$ Hz), 8.07 (1H, d, $J = 2.4$ Hz), 8.18-8.25 (1H, m), 8.26 (1H, dd, $J = 8.8, 2.0$ Hz), 8.49 (1H, dd, $J = 7.6, 2.0$ Hz)
47		604	δ 3.04 (3H, brs), 3.15 (3H, brs), 3.15-3.17 (2H, m), 3.55-3.59 (2H, m), 3.92 (3H, s), 4.02 (3H, s), 4.44 (1H, brs), 6.17 (1H, dd, $J = 8.0, 1.2$ Hz), 6.29 (1H, dd, $J = 8.0, 1.6$ Hz), 6.46 (1H, t, $J = 2.0$ Hz), 6.86 (1H, brs), 6.92 (1H, t, $J = 8.0$ Hz), 7.03 (3H, m), 7.32 (1H, dt, $J = 7.6, 1.2$ Hz), 7.43 (1H, t, $J = 8.0$ Hz), 7.55-7.57 (2H, m), 7.70 (1H, dd, $J = 8.8, 2.4$ Hz), 7.74-7.81 (1H, m), 8.12-8.16 (2H, m)
48		604	δ 2.99 (3H, brs), 3.12 (3H, brs), 3.31 (2H, t, $J = 6.0$ Hz), 3.64-3.69 (2H, m), 3.86 (3H, s), 4.07 (3H, s), 4.33 (1H, brs), 6.30 (1H, dd, $J = 8.0, 1.6$ Hz), 6.35 (1H, dd, $J = 8.0, 2.0$ Hz), 6.48 (1H, t, $J = 2.0$ Hz), 6.87-6.89 (2H, m), 6.94 (1H, t, $J = 8.0$ Hz), 7.06 (1H, d, $J = 8.8$ Hz), 7.32-7.35 (1H, m), 7.44 (1H, t, $J = 8.0$ Hz), 7.51-7.55 (2H, m), 7.68-7.79 (3H, m), 8.06 (1H, d, $J = 2.4$ Hz), 8.07-8.13 (1H, m)
49		604	δ 3.04 (3H, brs), 3.14 (3H, brs), 3.14-3.18 (2H, m), 3.58-3.61 (2H, m), 3.82 (3H, s), 4.02 (3H, s), 4.56 (1H, brs), 6.15 (1H, dd, $J = 8.0, 1.2$ Hz), 6.29 (1H, dd, $J = 8.4, 1.6$ Hz), 6.48 (1H, t, $J = 2.0$ Hz), 6.89-6.94 (2H, m), 7.03 (1H, d, $J = 8.8$ Hz), 7.25-7.31 (2H, m), 7.42 (1H, t, $J = 8.0$ Hz), 7.54-7.57 (1H, m), 7.64-7.70 (2H, m), 7.91-7.97 (1H, m), 8.15 (1H, d, $J = 2.4$ Hz), 8.35 (1H, d, $J = 2.8$ Hz), 8.53 (1H, d, $J = 1.6$ Hz)

[0552]

Table 16

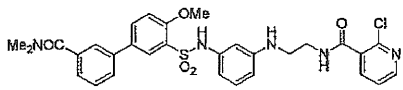
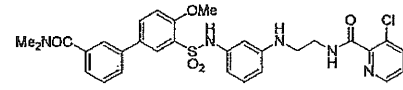
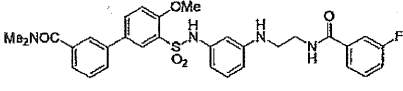
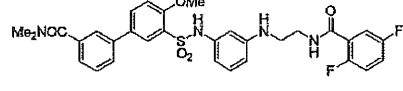
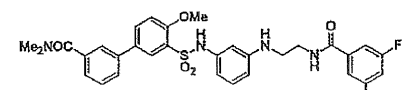
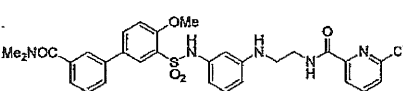
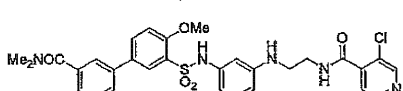
50		608	δ 2.99 (3H, brs), 3.06 (3H, brs), 3.23-3.27 (2H, m), 3.59-3.64 (2H, m), 4.05 (3H, s), 4.23-4.29 (1H, m), 6.22 (1H, dd, J = 7.6, 1.2 Hz), 6.31 (1H, dd, J = 8.0, 1.6 Hz), 6.48 (1H, t, J = 2.0 Hz), 6.88 (1H, brs), 6.94 (1H, t, J = 8.0 Hz), 7.05 (1H, d, J = 8.8 Hz), 7.21-7.26 (2H, m), 7.40 (1H, dd, J = 7.6, 3.2 Hz), 7.43-7.50 (1H, m), 7.54-7.57 (2H, m), 7.72 (1H, dd, J = 8.8, 2.4 Hz), 7.85 (1H, dd, J = 7.6, 2.0 Hz), 8.11 (1H, d, J = 2.4 Hz), 8.39 (1H, dd, J = 8.8, 2.0 Hz)
51		608	δ 2.99 (3H, brs), 3.12 (3H, brs), 3.29-3.31 (2H, m), 3.61-3.66 (2H, m), 4.07 (3H, s), 4.18 (1H, brs), 6.33-6.36 (2H, m), 6.45 (1H, t, J = 2.0 Hz), 6.90 (1H, brs), 6.95 (1H, t, J = 8.0 Hz), 7.07 (1H, d, J = 8.8 Hz), 7.32-7.44 (3H, m), 7.52-7.54 (2H, m), 7.70 (1H, dd, J = 8.4, 2.0 Hz), 7.80 (1H, dd, J = 8.4, 1.6 Hz), 8.00-8.06 (1H, m), 8.06 (1H, d, J = 2.0 Hz), 8.41 (1H, dd, J = 8.4, 1.2 Hz)
52		591	δ 3.04 (3H, brs), 3.15 (3H, brs), 3.15-3.18 (2H, m), 3.57-3.61 (2H, m), 4.03 (3H, s), 4.54 (1H, brs), 6.17 (1H, dd, J = 8.0, 1.6 Hz), 6.30 (1H, dd, J = 8.0, 2.4 Hz), 6.46 (1H, t, J = 2.0 Hz), 6.82 (1H, brs), 6.92 (1H, t, J = 8.0 Hz), 7.04 (1H, d, J = 8.4 Hz), 7.12-7.13 (1H, m), 7.26-7.34 (3H, m), 7.41-7.61 (5H, m), 7.68 (1H, dd, J = 8.8, 2.4 Hz), 8.15 (1H, d, J = 2.4 Hz)
53		609	δ 3.00 (3H, brs), 3.12 (3H, brs), 3.29 (2H, t, J = 6.0 Hz), 3.63-3.66 (2H, m), 4.07 (3H, s), 4.22 (1H, brs), 6.30 (1H, dd, J = 8.0, 1.6 Hz), 6.34 (1H, dd, J = 8.4, 1.6 Hz), 6.48 (1H, t, J = 2.0 Hz), 6.86 (1H, brs), 6.96 (1H, d, J = 8.0 Hz), 7.05-7.16 (4H, m), 7.32-7.34 (1H, m), 7.42 (1H, t, J = 7.6 Hz), 7.52-7.55 (2H, m), 7.69-7.72 (2H, m), 8.08 (1H, d, J = 2.4 Hz)
54		609	δ 3.06 (3H, brs), 3.11 (2H, t, J = 5.6 Hz), 3.16 (3H, brs), 3.53-3.57 (2H, m), 4.02 (3H, s), 4.52 (1H, brs), 6.15 (1H, dd, J = 8.0, 1.6 Hz), 6.28 (1H, dd, J = 8.4, 2.4 Hz), 6.43 (1H, t, J = 2.0 Hz), 6.84-6.93 (3H, m), 7.03 (1H, d, J = 8.4 Hz), 7.31-7.34 (3H, m), 7.44 (1H, t, J = 8.0 Hz), 7.57-7.59 (2H, m), 7.68 (1H, dd, J = 8.4, 2.4 Hz), 7.97-8.03 (1H, m), 8.17 (1H, d, J = 2.4 Hz)
55		608	δ 3.00 (3H, brs), 3.13 (3H, brs), 3.28-3.32 (2H, m), 3.63-3.67 (2H, m), 4.07 (3H, s), 4.17-4.23 (1H, m), 6.33-6.36 (2H, m), 6.45 (1H, t, J = 2.0 Hz), 6.86 (1H, brs), 6.96 (1H, t, J = 8.0 Hz), 7.08 (1H, d, J = 8.8 Hz), 7.35 (1H, dt, J = 7.2, 1.2 Hz), 7.41-7.55 (4H, m), 7.71 (1H, dd, J = 8.8, 2.4 Hz), 7.82 (1H, t, J = 8.0 Hz), 8.07-8.10 (2H, m), 8.11 (1H, d, J = 1.2 Hz)
56		608	δ 2.99 (3H, brs), 3.05 (3H, brs), 3.22-3.24 (2H, m), 3.59-3.63 (2H, m), 4.05 (3H, s), 4.22-4.29 (1H, m), 6.19 (1H, dd, J = 8.0, 1.6 Hz), 6.30 (1H, dd, J = 8.0, 2.0 Hz), 6.50 (1H, brs), 6.85 (1H, brs), 6.93 (1H, t, J = 8.0 Hz), 6.94 (1H, d, J = 8.8 Hz), 7.19 (1H, dt, J = 7.6, 1.6 Hz), 7.26-7.29 (1H, m), 7.40 (1H, t, J = 8.0 Hz), 7.52-7.58 (2H, m), 7.66-7.72 (1H, m), 7.72 (1H, dd, J = 8.4, 2.4 Hz), 8.13 (1H, d, J = 2.4 Hz), 8.42 (1H, d, J = 8.8 Hz), 8.51 (1H, s)

Table 17

57		δ 2.98 (3H, bs), 3.06 (3H, bs), 3.28 (2H, t, J = 5.6 Hz), 3.69-3.74 (2H, m), 4.04 (3H, s), 4.36-4.41 (1H, m), 6.18 (1H, dd, J = 8.0, 1.6 Hz), 6.28-6.32 (2H, m), 6.45 (1H, bs), 6.87 (1H, t, J = 8.0 Hz), 7.02 (1H, d, J = 8.8 Hz), 7.14-7.17 (1H, m), 7.24-7.33 (1H, m), 7.41-7.52 (2H, m), 7.65 (1H, dd, J = 8.4, 2.4 Hz), 7.96-8.03 (1H, m), 8.09-8.13 (2H, m), 8.47 (1H, d, J = 6.0 Hz), 9.06 (1H, s)
58		δ 3.07 (3H, bs), 3.07-3.10 (2H, m), 3.17 (3H, bs), 3.53-3.58 (2H, m), 4.01 (3H, s), 4.55-4.61 (1H, m), 6.12 (1H, dd, J = 7.6, 1.2 Hz), 6.29 (1H, dd, J = 8.0, 1.6 Hz), 6.43 (1H, t, J = 6.0 Hz), 6.86 (1H, bs), 6.91 (1H, t, J = 8.0 Hz), 7.03 (1H, d, J = 8.8 Hz), 7.29-7.31 (1H, m), 7.42 (1H, td, J = 7.6, 0.8 Hz), 7.57-7.59 (2H, m), 7.69 (1H, dd, J = 8.4, 2.0 Hz), 8.11 (1H, t, J = 2.0 Hz), 8.18 (1H, d, J = 2.4 Hz), 8.36-8.42 (1H, m), 8.59 (1H, d, J = 2.4 Hz), 8.84 (1H, d, J = 1.6 Hz)
59		δ 3.05 (3H, bs), 3.07-3.17 (5H, m), 3.51-3.56 (2H, m), 4.03 (3H, s), 4.10 (3H, s), 4.22-4.29 (1H, m), 6.16 (1H, d, J = 7.6 Hz), 6.28 (1H, dd, J = 8.0, 2.0 Hz), 6.48-6.51 (2H, m), 6.86 (1H, bs), 6.92 (1H, t, J = 8.0 Hz), 7.04 (1H, d, J = 8.8 Hz), 7.30-7.33 (2H, m), 7.42 (2H, m), 7.56-7.58 (2H, m), 7.68 (1H, dd, J = 8.4, 2.4 Hz), 8.14 (1H, d, J = 2.4 Hz)
60		δ 2.44 (3H, s), 3.00 (3H, bs), 3.13 (3H, bs), 3.24 (2H, t, J = 5.6 Hz), 3.57-3.61 (2H, m), 4.06 (3H, s), 4.31-4.38 (1H, m), 6.28 (1H, dd, J = 8.0, 2.0 Hz), 6.33 (1H, dd, J = 8.0, 2.0 Hz), 6.44-6.46 (2H, m), 6.84-6.87 (2H, m), 6.95 (1H, t, J = 8.0 Hz), 7.05 (1H, d, J = 8.8 Hz), 7.23 (1H, d, J = 4.8 Hz), 7.33 (1H, dt, J = 7.6, 1.2 Hz), 7.42 (1H, t, J = 8.0 Hz), 7.53-7.56 (2H, m), 7.69 (1H, dd, J = 8.4, 2.0 Hz), 8.09 (1H, d, J = 2.4 Hz)
61		δ 3.06 (3H, bs), 3.12-3.16 (2H, m), 3.16 (3H, bs), 3.50-3.54 (2H, m), 4.03 (3H, s), 4.49 (1H, bs), 6.15 (1H, dd, J = 8.0, 1.2 Hz), 6.28 (1H, dd, J = 8.0, 1.6 Hz), 6.46 (1H, t, J = 2.0 Hz), 6.78 (1H, d, J = 3.6 Hz), 6.82 (1H, bs), 6.91 (1H, t, J = 8.0 Hz), 7.04 (1H, d, J = 8.8 Hz), 7.26-7.35 (3H, m), 7.45 (1H, t, J = 7.6 Hz), 7.56-7.60 (2H, m), 7.68 (1H, dd, J = 8.8, 2.4 Hz), 8.15 (1H, d, J = 2.4 Hz)
613		δ 3.00 (3H, bs), 3.13 (3H, bs), 3.29-3.32 (2H, m), 3.62-3.67 (2H, m), 4.07 (3H, s), 4.20 (1H, bs), 6.32-6.36 (2H, m), 6.47 (1H, t, J = 2.0 Hz), 6.86 (1H, bs), 6.93-6.98 (2H, m), 7.07 (1H, d, J = 8.8 Hz), 7.26-7.46 (2H, m), 7.52-7.55 (2H, m), 7.69 (1H, dd, J = 8.8, 2.4 Hz), 8.07 (1H, d, J = 2.4 Hz)
613		δ 2.99 (3H, bs), 3.13 (3H, bs), 3.23-3.26 (2H, m), 3.56-3.61 (2H, m), 3.73 (3H, s), 4.06 (3H, s), 4.29 (1H, bs), 6.31-6.40 (3H, m), 6.93-6.97 (2H, m), 7.07 (1H, d, J = 8.8 Hz), 7.26-7.38 (3H, m), 7.41-7.45 (1H, m), 7.50-7.55 (3H, m), 7.69 (1H, dd, J = 8.4, 2.4 Hz), 8.05 (1H, d, J = 2.4 Hz)

Table 18

135

[0555]

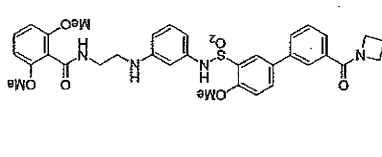
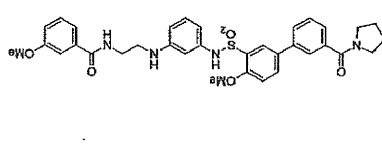
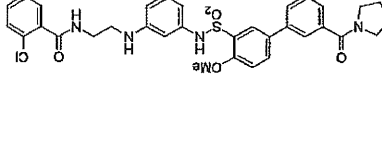
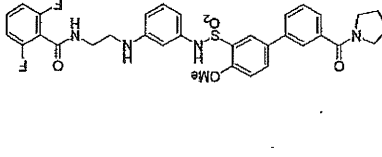
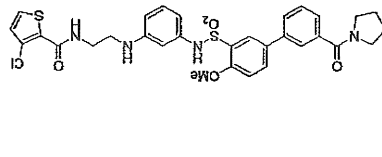
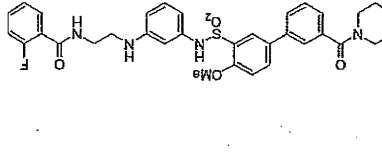
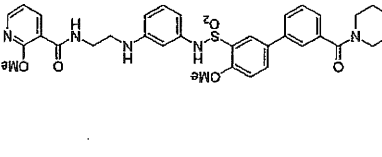
Table 19

71		598	δ 2.99 (3H, brs), 3.13 (3H, brs), 3.34-3.38 (2H, m), 4.06 (3H, s), 4.07-4.11 (2H, m), 4.50-4.55 (1H, m), 6.30 (2H, d, J = 8.0 Hz), 6.39 (1H, brs), 6.81 (1H, brs), 6.92 (1H, t, J = 8.0 Hz), 7.05 (1H, d, J = 8.8 Hz), 7.33-7.35 (1H, m), 7.43 (1H, t, J = 7.6 Hz), 7.52-7.55 (2H, m), 7.63-7.70 (4H, m), 7.81-7.83 (1H, m), 8.07 (1H, d, J = 2.0 Hz), 8.68 (1H, brs)
72		598	δ 3.07 (3H, brs), 3.07-3.11 (2H, m), 3.18 (3H, brs), 3.54-3.58 (2H, m), 4.01 (3H, s), 4.63 (1H, brs), 6.11 (1H, d, J = 7.6 Hz), 6.28 (1H, dd, J = 8.0, 1.6 Hz), 6.44 (1H, t, J = 2.0 Hz), 6.81 (1H, brs), 6.91 (1H, t, J = 8.0 Hz), 7.03 (1H, d, J = 8.8 Hz), 7.33-7.36 (1H, m), 7.42-7.48 (2H, m), 7.58-7.61 (2H, m), 7.68-7.71 (2H, m), 8.02-8.05 (1H, m), 8.10 (1H, t, J = 1.2 Hz), 8.19-8.20 (1H, m), 8.25-8.30 (1H, m)
73		589	δ 3.01 (3H, d, J = 4.8 Hz), 3.24-3.28 (2H, m), 3.55-3.60 (2H, m), 3.80 (3H, s), 4.06 (3H, s), 4.22-4.33 (1H, m), 6.30-6.35 (2H, m), 6.46-6.48 (2H, m), 6.87 (1H, brs), 6.91-6.98 (2H, m), 7.04-7.08 (2H, m), 7.41-7.46 (2H, m), 7.58-7.60 (1H, m), 7.68-7.71 (2H, m), 7.88 (1H, t, J = 1.6 Hz), 8.09-8.12 (2H, m), 8.13 (1H, dd, J = 8.0, 2.0 Hz)
74		619	δ 3.22-3.25 (2H, m), 3.47-3.51 (2H, m), 3.70-3.74 (2H, m), 3.76 (3H, s), 3.82 (3H, s), 3.92-3.95 (2H, m), 4.06-4.16 (2H, m), 6.26 (1H, dd, J = 7.2, 1.6 Hz), 6.46-6.48 (2H, m), 6.73 (1H, d, J = 8.8 Hz), 6.91-6.97 (2H, m), 7.06-7.10 (1H, m), 7.40-7.48 (2H, m), 7.52-7.55 (3H, m), 7.85-7.87 (2H, m), 7.98 (1H, brs), 8.06-8.12 (3H, m)
75		629	δ 1.85-1.89 (2H, m), 1.95-1.99 (2H, m), 3.30 (2H, t, J = 6.0 Hz), 3.40-3.44 (2H, m), 3.62-3.67 (4H, m), 3.81 (3H, s), 4.07 (3H, s), 4.38 (1H, brs), 6.32-6.36 (2H, m), 6.47 (1H, t, J = 2.0 Hz), 6.90-6.97 (3H, m), 7.04-7.09 (2H, m), 7.40-7.46 (3H, m), 7.52 (1H, dt, J = 7.2, 1.6 Hz), 7.63 (1H, d, J = 1.2 Hz), 7.69 (1H, dd, J = 8.8, 1.6 Hz), 8.06 (1H, d, J = 2.4 Hz), 8.09-8.15 (1H, m), 8.18 (1H, dd, J = 8.0, 2.0 Hz)
76		645	δ 1.52 (9H, s), 2.87 (3H, s), 3.28-3.31 (2H, m), 3.62-3.67 (2H, m), 3.81 (3H, s), 4.07 (3H, s), 4.36 (1H, brs), 6.31-6.36 (2H, m), 6.47 (1H, t, J = 2.0 Hz), 6.89 (1H, brs), 6.91-6.95 (2H, m), 7.04-7.09 (2H, m), 7.32-7.50 (4H, m), 7.54 (1H, d, J = 1.2 Hz), 7.68 (1H, dd, J = 8.8, 2.4 Hz), 8.06 (1H, d, J = 2.4 Hz), 8.11-8.14 (1H, m), 8.18 (1H, dd, J = 7.6, 2.0 Hz)
77		647	δ 3.03-3.13 (3H, m), 3.30-3.49 (7H, m), 3.64-3.73 (4H, m), 3.82 (3H, s), 4.07 (3H, s), 4.30 (1H, brs), 6.32-6.36 (2H, m), 6.47 (1H, t, J = 2.0 Hz), 6.86 (1H, d, J = 4.8 Hz), 6.92-6.97 (2H, m), 7.04-7.09 (2H, m), 7.35-7.52 (5H, m), 7.67 (1H, dd, J = 8.4, 1.6 Hz), 8.06 (1H, d, J = 2.0 Hz), 8.09-8.16 (1H, m), 8.19 (1H, dd, J = 8.0, 1.6 Hz)

[0556]

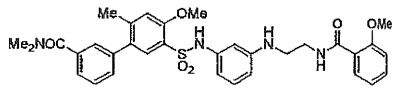
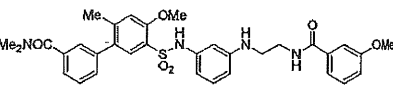
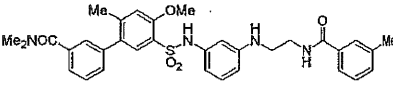
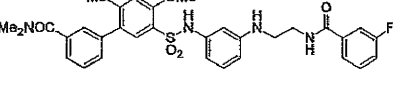
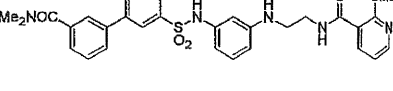
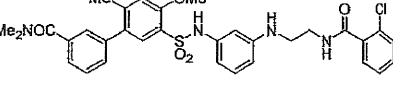
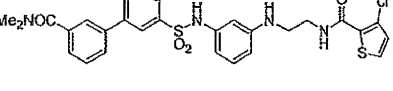
Table 20

78		657	δ 3.24 (2H, t, J = 6.0 Hz), 3.53-3.64 (2H, m), 3.80 (3H, s), 4.05 (3H, s), 4.08-4.17 (2H, m), 4.25-4.31 (1H, m), 6.29-6.34 (2H, m), 6.44-6.48 (1H, m), 6.85 (1H, brs), 6.91-7.07 (5H, m), 7.41-7.52 (3H, m), 7.63-7.65 (1H, m), 7.69 (1H, dd, J = 8.4, 2.4 Hz), 7.74 (1H, m), 8.06-8.13 (3H, m)
79		603	δ 1.55 (6H, s), 3.30 (2H, t, J = 6.0 Hz), 3.62-3.67 (2H, m), 3.83 (3H, s), 4.06 (3H, s), 4.38 (1H, brs), 6.30-6.36 (2H, m), 6.46 (1H, t, J = 2.4 Hz), 6.82 (1H, brs), 6.91-7.02 (4H, m), 7.08 (1H, t, J = 7.6 Hz), 7.36-7.50 (4H, m), 7.78-7.83 (2H, m), 8.07-8.13 (1H, m), 8.19 (1H, dd, J = 8.0, 2.0 Hz)
80		643	δ 1.52-1.68 (6H, m), 3.10 (1H, brs), 3.28-3.35 (4H, m), 3.63-3.72 (4H, m), 3.82 (3H, s), 4.07 (3H, s), 4.29 (1H, brs), 6.31-6.36 (2H, m), 6.46-6.49 (1H, m), 6.92-7.00 (2H, m), 7.05-7.09 (2H, m), 7.30-7.32 (1H, m), 7.38-7.47 (2H, m), 7.50-7.53 (2H, m), 7.68 (1H, dd, J = 7.6, 2.4 Hz), 8.06 (1H, d, J = 2.4 Hz), 8.11-8.16 (1H, m), 8.18 (1H, dd, J = 8.0, 2.0 Hz)
81		615	δ 2.32-3.36 (2H, m), 3.30 (2H, t, J = 6.0 Hz), 3.62-3.67 (2H, m), 3.81 (3H, s), 4.07 (3H, s), 4.21-4.31 (5H, m), 6.31 (1H, dd, J = 8.0, 2.0 Hz), 6.34 (1H, dd, J = 8.0, 2.0 Hz), 6.47 (1H, t, J = 2.0 Hz), 6.85 (1H, brs), 6.91-6.97 (2H, m), 7.05-7.09 (2H, m), 7.39-7.46 (2H, m), 7.51-7.58 (2H, m), 7.69 (1H, dd, J = 8.4, 2.0 Hz), 7.76 (1H, t, J = 1.6 Hz), 8.05 (1H, d, J = 2.4 Hz), 8.08-8.13 (1H, m), 8.18 (1H, dd, J = 8.0, 2.0 Hz)
82		619	δ 2.05-2.11 (2H, m), 3.31 (2H, t, J = 6.0 Hz), 3.54-3.64 (6H, m), 4.07 (3H, s), 4.17-4.24 (1H, m), 6.28 (1H, dd, J = 8.0, 1.6 Hz), 6.34 (1H, dd, J = 8.0, 2.0 Hz), 6.50-6.53 (1H, m), 6.57-6.63 (1H, m), 6.74-6.78 (1H, m), 6.88 (1H, brs), 6.96 (1H, t, J = 8.0 Hz), 7.07 (1H, d, J = 8.8 Hz), 7.21-7.30 (2H, m), 7.43-7.47 (2H, m), 7.60-7.63 (2H, m), 7.70 (1H, dd, J = 8.4, 2.4 Hz), 7.90 (1H, s), 8.10 (1H, d, J = 2.4 Hz)
83		616	δ 2.11-2.18 (2H, m), 3.31 (2H, t, J = 5.6 Hz), 3.59-3.67 (6H, m), 3.84 (3H, s), 4.06 (3H, s), 4.18-4.25 (1H, m), 6.29 (1H, dd, J = 8.0, 1.6 Hz), 6.33 (1H, dd, J = 8.4, 1.6 Hz), 6.48 (1H, t, J = 2.0 Hz), 6.66-6.72 (1H, m), 6.87-6.89 (2H, m), 6.95 (1H, t, J = 8.0 Hz), 7.04 (1H, d, J = 8.4 Hz), 7.46 (1H, t, J = 8.0 Hz), 7.60-7.63 (1H, m), 7.64-7.74 (3H, m), 7.90 (1H, t, J = 1.6 Hz), 8.01-8.07 (1H, m), 8.09-8.11 (1H, m)
84		603	δ 2.09-2.13 (2H, m), 3.26 (2H, t, J = 5.6 Hz), 3.59-3.65 (6H, m), 4.02 (3H, s), 4.35 (1H, brs), 6.22 (1H, dd, J = 8.0, 1.6 Hz), 6.30 (1H, dd, J = 8.0, 1.6 Hz), 6.54 (1H, t, J = 2.0 Hz), 6.68-6.76 (1H, m), 6.91-7.03 (3H, m), 7.13-7.31 (2H, m), 7.38-7.47 (3H, m), 7.58-7.66 (3H, m), 7.91 (1H, s), 8.10 (1H, d, J = 2.4 Hz)

85		δ 2.04-2.11 (2H, m), 3.24-3.30 (2H, m), 3.51-3.62 (6H, m), 3.63 (3H, s), 3.64 (3H, s), 4.06 (3H, s), 4.28 (1H, bs), 6.14-6.18 (1H, m), 6.28 (2H, m), 6.43-6.51 (3H, m), 6.83-6.90 (1H, m), 6.93-6.97 (1H, m), 7.05-7.09 (1H, m), 7.20 (1H, t, J = 8.4 Hz), 7.42 (1H, t, J = 8.0 Hz), 7.56-7.59 (1H, m), 7.66-7.71 (2H, m), 7.90 (1H, s), 8.08 (1H, d, J = 2.4 Hz)
86		δ 1.90-2.01 (4H, m), 3.22 (2H, t, J = 5.6 Hz), 3.45-3.48 (2H, m), 3.61-3.67 (4H, m), 3.79 (3H, s), 4.04 (3H, s), 4.40-4.49 (1H, m), 6.19 (1H, dd, J = 8.0, 1.2 Hz), 6.31 (1H, dd, J = 8.0, 1.6 Hz), 6.49 (1H, t, J = 2.0 Hz), 6.82 (1H, bs), 6.92 (1H, t, J = 8.0 Hz), 6.98 (1H, ddd, J = 8.0, 1.2 Hz), 7.03 (1H, d, J = 8.0 Hz), 7.21-7.28 (2H, m), 7.36 (1H, t, J = 2.0 Hz), 7.41-7.43 (3H, m), 7.54-7.55 (1H, m), 7.66-7.70 (2H, m), 8.12-8.13 (1H, m)
87		δ 1.86-1.97 (4H, m), 3.28 (2H, t, J = 5.6 Hz), 3.40-3.44 (2H, m), 3.62-3.66 (4H, m), 4.06 (3H, s), 4.30 (1H, bs), 6.26 (1H, dd, J = 8.0, 1.2 Hz), 6.32 (1H, dd, J = 8.0, 2.0 Hz), 6.48 (1H, t, J = 2.0 Hz), 6.87 (1H, bs), 6.95 (1H, t, J = 8.0 Hz), 6.98-7.04 (1H, m), 7.05 (1H, d, J = 8.4 Hz), 7.22-7.32 (3H, m), 7.37-7.42 (2H, m), 7.49-7.56 (2H, m), 7.64 (1H, s), 7.69 (1H, dd, J = 8.8, 2.4 Hz), 8.09 (1H, d, J = 2.4 Hz)
88		δ 1.87-1.98 (4H, m), 3.24-3.27 (2H, m), 3.41-3.44 (2H, m), 3.55-3.63 (4H, m), 4.06 (3H, s), 4.20 (1H, bs), 6.27 (1H, dd, J = 8.0, 1.6 Hz), 6.33 (1H, dd, J = 8.0, 2.0 Hz), 6.46 (1H, t, J = 8.0 Hz), 6.85-7.00 (5H, m), 7.06 (1H, d, J = 8.4 Hz), 7.29-7.42 (2.0 Hz), 7.54 (1H, d, J = 7.2, 1.6 Hz), 7.62 (1H, s), 7.69 (1H, dd, J = 8.8, 2.4 Hz), 8.09 (1H, d, J = 2.0 Hz)
89		δ 1.87-1.98 (4H, m), 3.22-3.29 (2H, m), 3.41-3.45 (2H, m), 3.63-3.67 (4H, m), 4.07 (3H, s), 4.23 (1H, bs), 6.32-6.36 (2H, m), 6.47 (1H, t, J = 2.0 Hz), 6.90-6.98 (3H, m), 7.06 (1H, d, J = 8.8 Hz), 7.42-7.45 (4H, m), 7.52-7.55 (1H, m), 7.65 (1H, s), 7.69 (1H, dd, J = 8.4, 2.4 Hz), 8.08 (1H, d, J = 2.4 Hz)
90		δ 1.53-1.68 (6H, m), 3.28-3.36 (4H, m), 3.64-3.71 (4H, m), 4.07 (3H, s), 4.25 (1H, bs), 6.31-6.36 (2H, m), 6.46 (1H, t, J = 2.0 Hz), 6.84 (1H, bs), 6.96 (1H, t, J = 8.0 Hz), 7.05-7.11 (3H, m), 7.23-7.32 (2H, m), 7.39-7.47 (2H, m), 7.52-7.54 (2H, m), 7.69 (1H, dd, J = 8.4, 2.0 Hz), 8.02-8.08 (2H, m)
91		δ 1.52-1.68 (6H, m), 3.29-3.51 (4H, m), 3.64-3.72 (4H, m), 3.98 (3H, s), 4.07 (3H, s), 4.30 (1H, bs), 6.31 (1H, dd, J = 8.0, 1.6 Hz), 6.35 (1H, dd, J = 8.0, 2.4 Hz), 6.50 (1H, t, J = 2.0 Hz), 6.89 (1H, bs), 6.96 (1H, t, J = 8.0 Hz), 7.04-7.07 (2H, m), 7.30-7.32 (1H, m), 7.41 (1H, t, J = 8.0 Hz), 7.51-7.53 (2H, m), 7.70 (1H, dd, J = 8.8, 2.4 Hz), 8.07 (1H, d, J = 2.4 Hz), 8.14-8.19 (1H, m), 8.26 (1H, dd, J = 8.8, 2.0 Hz), 8.49 (1H, dd, J = 7.6, 2.0 Hz)

98		5 3.29-3.32 (2H, m), 3.61-3.65 (2H, s), 3.83 (3H, s), 6.50 (3H, s), 4.07 (3H, s), 4.18-4.25 (1H, m), 6.31-6.36 (2H, m), 7.01 (1H, t, J = 2.0 Hz), 6.56-6.64 (1H, m), 6.93-6.97 (2H, m), 7.22-7.33 (4H, m), 7.67 (1H, dd, J = 8.8, 2.4 Hz), 8.03 (1H, d, J = 2.4 Hz), 8.26 (1H, d, J = 2.8 Hz), 8.31 (1H, d, J = 2.0 Hz)
97		5 3.28-3.31 (2H, m), 3.63-3.67 (2H, m), 3.84 (3H, s), 3.89 (3H, s), 4.08 (3H, s), 4.29-4.37 (1H, m), 6.32-6.37 (2H, m), 6.46 (1H, t, J = 2.0 Hz), 6.85-6.90 (1H, m), 6.93-6.98 (2H, m), 7.06-7.10 (2H, m), 7.25-7.27 (1H, m), 7.43-7.47 (1H, m), 7.67 (1H, dd, J = 8.4, 2.4 Hz), 8.03 (1H, d, J = 2.4 Hz), 8.18 (1H, dd, J = 8.0, 2.0 Hz), 8.25 (1H, d, J = 2.8 Hz), 8.33 (1H, d, J = 2.0 Hz)
96		5 3.30 (2H, t, J = 6.0 Hz), 3.63-3.67 (2H, m), 3.85 (3H, s), 4.08 (3H, s), 4.29-4.37 (1H, m), 6.33-6.36 (2H, m), 6.47 (1H, s), 6.93-6.98 (3H, m), 7.07-7.11 (2H, m), 7.32 (1H, dd, J = 8.0, 4.8 Hz), 7.43-7.47 (1H, m), 7.67 (1H, dd, J = 8.8, 2.4 Hz), 7.77 (1H, dd, J = 8.0, 1.6 Hz), 8.04 (1H, d, J = 2.4 Hz), 8.18 (1H, dd, J = 8.0, 1.6 Hz), 8.55 (1H, d, J = 4.4 Hz), 8.74 (1H, s)
95		5 3.30 (2H, t, J = 6.0 Hz), 3.59-3.63 (2H, m), 3.83 (3H, s), 3.98 (3H, s), 4.07 (3H, s), 4.26-4.32 (1H, m), 6.32-6.34 (2H, m), 6.34-6.37 (1H, m), 6.50 (1H, t, J = 2.0 Hz), 6.64 (1H, dd, J = 8.0, 0.8 Hz), 6.91-6.96 (2H, m), 7.01 (1H, ddd, J = 8.0, 2.8, 0.8 Hz), 7.07 (1H, d, J = 8.8 Hz), 7.17-7.20 (1H, m), 7.24-7.31 (3H, m), 7.57 (1H, dd, J = 8.4, 3.6 Hz), 8.23 (1H, dd, J = 8.8, 2.4 Hz), 8.46 (1H, d, J = 2.4 Hz)
94		5 3.30 (2H, t, J = 6.0 Hz), 3.61-3.66 (2H, m), 3.79 (3H, s), 3.98 (3H, s), 4.08 (3H, s), 4.25-4.32 (1H, m), 6.33 (2H, dd, J = 8.0, 2.0 Hz), 6.49 (1H, t, J = 2.0 Hz), 6.64 (1H, d, J = 8.4 Hz), 6.89-6.96 (3H, m), 7.06-7.10 (2H, m), 7.24-7.26 (1H, m), 7.41-7.46 (1H, m), 7.57 (1H, t, J = 4.0 Hz), 8.05-8.11 (1H, m), 8.18 (1H, dd, J = 7.6, 2.0 Hz), 8.23 (1H, dd, J = 8.8, 2.4 Hz), 8.45 (1H, d, J = 2.0 Hz)
93		5 3.30 (2H, t, J = 6.0 Hz), 3.53-3.61 (2H, m), 3.83 (3H, s), 4.07 (3H, s), 6.28-6.33 (2H, m), 6.50-6.54 (2H, m), 6.91-6.96 (1H, m), 7.02 (1H, ddd, J = 8.4, 6.4, 1.2 Hz), 7.08 (1H, d, J = 8.8 Hz), 7.18-7.31 (6H, m), 7.40-7.45 (1H, m), 8.15 (1H, ddd, J = 8.8, 6.4, 1.2 Hz), 8.41 (1H, dt, J = 8.4, 1.6 Hz), 8.54 (1H, dd, J = 2.0, 1.2 Hz)
92		5 1.56-1.71 (6H, m), 3.10-3.19 (2H, m), 3.38-3.43 (2H, m), 3.55-3.60 (2H, m), 3.70-3.77 (2H, m), 4.03 (3H, s), 4.51 (1H, bs), 6.17 (1H, dd, J = 8.0, 1.2 Hz), 6.30 (1H, dd, J = 8.0, 2.4 Hz), 6.44 (1H, t, J = 2.0 Hz), 6.82 (1H, bs), 6.92 (1H, t, J = 8.0 Hz), 7.04 (1H, d, J = 8.8 Hz), 7.11-7.16 (1H, m), 7.26-7.33 (2H, m), 7.40-7.52 (3H, m), 7.55-7.57 (2H, m), 7.69 (1H, dd, J = 8.4, 2.0 Hz), 8.15-8.16 (1H, m)
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Table 23

99		617	δ 2.27 (3H, s), 2.98 (3H, brs), 3.10 (3H, brs), 3.29-3.33 (2H, m), 3.63-3.68 (2H, m), 3.85 (3H, s), 4.04 (3H, s), 4.31 (1H, brs), 6.31 (1H, dd, $J = 7.6, 1.6$ Hz), 6.36 (1H, dd, $J = 7.6, 1.6$ Hz), 6.47 (1H, t, $J = 2.0$ Hz), 6.81 (1H, brs), 6.86 (1H, s), 6.93-6.99 (2H, m), 7.06-7.10 (1H, m), 7.23-7.26 (2H, m), 7.35-7.47 (3H, m), 7.68 (1H, s), 8.08-8.14 (1H, m), 8.18 (1H, dd, $J = 8.0, 1.6$ Hz)
100		617	δ 2.26 (3H, s), 3.04 (3H, brs), 3.10 (3H, brs), 3.20-3.23 (2H, m), 3.59-3.63 (2H, m), 3.81 (3H, s), 4.00 (3H, s), 4.52 (1H, brs), 6.20 (1H, dd, $J = 8.0, 1.6$ Hz), 6.33 (1H, dd, $J = 8.0, 1.6$ Hz), 6.46 (1H, t, $J = 2.0$ Hz), 6.81 (1H, brs), 6.84 (1H, s), 6.95 (1H, t, $J = 8.0$ Hz), 7.00 (1H, ddd, $J = 8.0, 2.4, 0.8$ Hz), 7.25-7.41 (8H, m), 7.76 (1H, s)
101		601	δ 2.26 (3H, s), 2.36 (3H, s), 3.03 (3H, brs), 3.10 (3H, brs), 3.20-3.23 (2H, m), 3.59-3.63 (2H, m), 4.01 (3H, s), 4.46 (1H, brs), 6.22 (1H, dd, $J = 7.6, 1.6$ Hz), 6.33 (1H, dd, $J = 8.0, 1.6$ Hz), 6.46 (1H, t, $J = 2.0$ Hz), 6.80 (1H, brs), 6.84 (1H, s), 6.95 (1H, t, $J = 8.0$ Hz), 7.22-7.30 (1H, m), 7.30-7.41 (6H, m), 7.54-7.58 (1H, m), 7.62 (1H, s), 7.76 (1H, s)
102		605	δ 2.26 (3H, s), 3.06 (3H, brs), 3.12 (3H, brs), 3.15-3.18 (2H, m), 3.56-3.62 (2H, m), 3.99 (3H, s), 4.52 (1H, brs), 6.16 (1H, dd, $J = 7.6, 1.6$ Hz), 6.31 (1H, dd, $J = 8.0, 1.6$ Hz), 6.45 (1H, t, $J = 2.0$ Hz), 6.80 (1H, brs), 6.83 (1H, s), 6.94 (1H, t, $J = 8.0$ Hz), 7.11-7.17 (1H, m), 7.29-7.44 (5H, m), 7.54-7.61 (2H, m), 7.62-7.70 (1H, m), 7.79 (1H, s)
103		618	δ 2.27 (3H, s), 2.99 (3H, brs), 3.10 (3H, brs), 3.29-3.33 (2H, m), 3.64-3.69 (2H, m), 3.99 (3H, s), 4.04 (3H, s), 4.27 (1H, brs), 6.30 (1H, dd, $J = 7.6, 1.6$ Hz), 6.36 (1H, dd, $J = 8.0, 1.6$ Hz), 6.49 (1H, t, $J = 2.0$ Hz), 6.82 (1H, brs), 6.86 (1H, s), 6.97 (1H, t, $J = 8.0$ Hz), 7.05 (1H, dd, $J = 7.6, 1.6$ Hz), 7.23-7.26 (2H, m), 7.35-7.40 (2H, m), 7.69 (1H, s), 8.14-8.22 (1H, m), 8.27 (1H, dd, $J = 8.8, 2.0$ Hz), 8.49 (1H, dd, $J = 7.6, 2.0$ Hz)
104		621	δ 2.27 (3H, s), 2.99 (3H, brs), 3.01 (3H, brs), 3.21-3.31 (2H, m), 3.62-3.67 (2H, m), 4.03 (3H, s), 4.29 (1H, brs), 6.25 (1H, dd, $J = 8.0, 1.6$ Hz), 6.34 (1H, dd, $J = 8.0, 2.0$ Hz), 6.48 (1H, t, $J = 2.0$ Hz), 6.83 (1H, brs), 6.86 (1H, s), 6.94-6.98 (2H, m), 7.25-7.42 (7H, m), 7.54-7.57 (1H, m), 7.72 (1H, s)
105		627	δ 2.27 (3H, s), 3.00 (3H, brs), 3.11 (3H, brs), 3.29-3.33 (2H, m), 3.62-3.67 (2H, m), 4.04 (3H, s), 4.20 (1H, brs), 6.32 (1H, dd, $J = 8.0, 2.0$ Hz), 6.36 (1H, dd, $J = 8.0, 2.0$ Hz), 6.47 (1H, t, $J = 2.0$ Hz), 6.83 (1H, brs), 6.87 (1H, s), 6.94-7.00 (2H, m), 7.25-7.27 (3H, m), 7.36-7.41 (2H, m), 7.45 (1H, d, $J = 5.6$ Hz), 7.70 (1H, s)

106		δ 2.68 (3H, s), 2.99 (3H, bs), 3.13 (3H, bs), 3.23-3.27 (2H, m), 3.59-3.65 (2H, m), 3.84 (3H, s), 4.38 (1H, bs), 6.31-6.35 (2H, m), 6.36 (1H, t, J = 2.0 Hz), 6.61 (1H, bs), 6.93-6.99 (2H, m), 7.05-7.09 (1H, m), 7.32-7.38 (2H, m), 7.41-7.47 (2H, m), 7.53-7.56 (2H, m), 7.62 (1H, dd, J = 8.0, 2.0 Hz), 8.08-8.16 (1H, m), 8.16-8.19 (2H, m)
107		δ 2.99 (3H, bs), 3.13 (3H, bs), 3.28-3.32 (2H, m), 3.63-3.68 (2H, m), 3.85 (3H, s), 4.39 (1H, bs), 6.37-6.41 (2H, m), 6.46 (1H, t, J = 2.0 Hz), 6.78 (1H, bs), 6.93-7.01 (2H, m), 7.05-7.10 (1H, t, J = 2.0 Hz), 7.21-7.24 (1H, m), 7.38 (1H, dt, J = 7.6, 1.2 Hz), 7.42-7.47 (2H, m), 7.50-7.54 (2H, m), 7.70 (1H, ddd, J = 7.6, 4.4, 2.4 Hz), 8.02 (1H, dd, J = 6.8, 2.4 Hz), 8.08-8.16 (1H, m), 8.18 (1H, dd, J = 8.0, 2.0 Hz)
108		δ 1.60 (3H, t, J = 6.8 Hz), 2.98 (3H, bs), 3.12 (3H, bs), 3.28-3.32 (2H, m), 3.63-3.68 (2H, m), 3.82 (3H, s), 4.30 (2H, q, J = 6.8 Hz), 4.33 (1H, bs), 6.32 (1H, dd, J = 8.0, 1.6 Hz), 6.35 (1H, dd, J = 8.0, 1.6 Hz), 6.49 (1H, t, J = 2.0 Hz), 6.80 (1H, bs), 6.92-6.98 (2H, m), 7.03-7.09 (2H, m), 7.33 (1H, dt, J = 7.6, 1.2 Hz), 7.39-7.46 (2H, m), 7.51-7.54 (2H, m), 7.66 (1H, dd, J = 8.4, 2.4 Hz), 8.05 (1H, d, J = 2.4 Hz), 8.08-8.16 (1H, m), 8.18 (1H, dd, J = 8.0, 1.6 Hz)
109		δ 3.28-3.32 (2H, m), 3.62-3.67 (2H, m), 3.83 (3H, s), 4.02 (3H, s), 4.25 (1H, bs), 6.31-6.36 (2H, m), 6.45-6.47 (1H, m), 6.83-7.10 (6H, m), 7.17 (1H, dt, J = 10.0, 2.4 Hz), 7.25-7.27 (1H, m), 7.32-7.37 (1H, m), 7.42-7.47 (1H, m), 7.65 (1H, dd, J = 8.8, 2.4 Hz), 8.04 (1H, d, J = 2.4 Hz), 8.18-8.22 (1H, m), 8.18-8.22 (1H, m)
573		δ 2.98 (3H, bs), 3.14 (3H, bs), 3.26-3.31 (2H, m), 3.62-3.68 (2H, m), 3.86 (3H, d, J = 7.6 Hz), 6.47 (1H, s), 6.50-6.57 (1H, m), 6.91-6.95 (2H, m), 7.00 (1H, t, J = 8.0 Hz), 7.05-7.09 (1H, m), 7.36-7.51 (6H, m), 7.69 (1H, d, J = 7.6 Hz), 7.77 (1H, d, J = 8.0 Hz), 7.94 (1H, s), 8.10-8.17 (1H, m), 8.18 (1H, dd, J = 8.0, 2.0 Hz)
617		δ 2.78 (3H, s), 2.99 (3H, bs), 3.06-3.11 (5H, m), 3.30-3.40 (2H, m), 3.76 (3H, s), 4.07 (3H, s), 6.31-6.37 (3H, m), 6.47 (1H, t, J = 2.0 Hz), 6.85-6.99 (4H, m), 7.07 (1H, d, J = 8.8 Hz), 7.16 (1H, dd, J = 7.6, 1.6 Hz), 7.32-7.36 (2H, m), 7.40-7.44 (1H, m), 7.51-7.54 (2H, m), 7.69-7.72 (1H, m), 8.04-8.06 (1H, m)
617		δ 2.92 (3H, s), 2.97 (3H, bs), 3.11 (3H, bs), 3.48-3.51 (2H, m), 3.56-3.59 (2H, m), 3.70 (3H, s), 4.08 (3H, s), 6.36 (1H, dd, J = 8.0, 1.6 Hz), 6.50 (1H, dd, J = 8.0, 2.0 Hz), 6.58 (1H, t, J = 2.0 Hz), 6.88 (1H, bs), 6.89 (1H, d, J = 8.0 Hz), 7.00-7.10 (3H, m), 7.33 (1H, dt, J = 7.6, 1.6 Hz), 7.37-7.45 (2H, m), 7.50-7.54 (2H, m), 7.68 (1H, dd, J = 8.4, 2.0 Hz), 7.99-8.04 (1H, m), 8.05-8.06 (1H, m), 8.20 (1H, dd, J = 8.0, 2.0 Hz)

[0561]

Experimental Example 1

Evaluation of agonist activity against OX2R

NAFT-luciferase gene and OX2R gene were each
5 constitutively expressed in CHO cell, which is a cell line
derived from Chinese hamster ovary to establish a cell line
(CHOOX2R). The cells thereof were sown on a 96 well plate at
10000 cells/well, and cultured in a 5% FBS-added DMEM medium
for 48 hr. The cells were dissolved in Fura2AM (final
10 concentration 40 μ M), HEPE buffer (HEPES Buffer containing 20
mM HEPES, 115 mM NaCl, 5.4 mM KCl, 0.8 mM MgCl₂, 1.8 mM CaCl₂,
13.8 mM D-glucose, 2.5 mM probenecid acid and 0.1% BSA), and
fluorescence labeled at 37°C for 60 min. After washing, HEPES
buffer (75 μ L) was added, a test compound solution (25 μ L) was
15 added, and changes in the fluorescence intensity were measured
by FDSS3000 (Hamamatsu Photonics K.K.). The test compound was
dissolved in DMSO (final concentration of DMSO was 1%), and
diluted with HEPES buffer to a final concentration of 0.1 μ M,
1.0 μ M or 10 μ M. The agonist activity value of each compound
20 is shown in Table 25 to Table 29.

[0562]

Table 25

Example No.	Response (10 μ M (%))	Response (1.0 μ M (%))	Response (0.1 μ M (%))
1	57	21	<5
2	67	27	<5
3	50	11	<5
4	17	<5	<5
5	23	<5	<5
6	7	<5	<5
7	145	62	10
8	46	9	<5
9	<5	<5	<5
10	41	<5	<5
11	23	17	<5
12	73	44	<5
13	99	64	<5
14	8	<5	<5
15	6	<5	<5
16	<5	<5	<5
17	182	18	<5
18	151	13	<5
19	143	8	<5
20	180	13	<5
21	11	<5	<5
22	21	15	<5
23	<5	<5	<5
24	<5	<5	<5
25	38	7	<5
26	29	5	<5
27	<5	<5	<5
28	<5	<5	<5
29	73	16	<5
31	103	67	6
32	75	26	<5
33	93	31	<5

[0563]

Table 26

Example No.	Response (10 μ M (%))	Response (1.0 μ M (%))	Response (0.1 μ M (%))
34	123	64	13
35	127	68	13
36	68	11	<5
37	163	102	16
38	113	37	8
39	102	27	<5
40	74	74	35
41	115	107	70
42	114	103	65
43	94	55	<5
44	94	28	<5
45	98	80	<5
46	91	89	18
47	82	89	14
48	84	91	44
49	88	84	9
50	76	71	<5
51	87	9	<5
52	64	79	40
53	66	89	46
54	126	168	33
55	122	144	23
56	122	22	<5
57	87	9	<5
58	119	130	<5
59	64	<5	<5
60	75	43	<5
61	69	18	<5
62	94	92	71
63	82	81	<5
64	72	68	<5
65	74	96	74

[0564]

Table 27

Example No.	Response (10 μ M (%))	Response (1.0 μ M (%))	Response (0.1 μ M (%))
66	67	88	97
67	76	96	55
68	74	86	42
69	111	123	67
70	85	105	34
71	80	39	<5
72	88	16	<5
73	98	90	45
74	92	51	<5
75	76	87	49
76	98	95	24
77	82	91	42
78	49	33	<5
79	34	18	<5
80	91	91	37
81	98	98	75
82	74	51	<5
83	71	10	<5
84	70	58	7
85	90	90	19
86	81	89	10
87	80	90	42
88	86	100	76
89	97	96	93
90	88	68	51
91	80	90	42
92	94	92	27
93	73	33	<5
94	22	<5	<5
95	30	<5	<5
96	88	61	<5
97	37	12	<5

[0565]

Table 28

Example No.	Response (10 μ M (%))	Response (1.0 μ M (%))	Response (0.1 μ M (%))
98	91	22	<5
102	85	83	61
103	80	93	77
104	86	88	67
105	79	84	78
106	39	27	<5
107	82	79	65
108	82	81	44
109	47	46	13
110	28	<5	<5
112	139	131	<5

[0566]

- 5 (As used herein, Response in Table 25 to Table 28 is a value obtained by dividing the agonist activity value, when the test compound is evaluated with orexin-A as a full-agonist (maximum value of agonist activity: 100%), at 10 μ M, 1.0 μ M, 0.1 μ M by the agonist activity value of orexin-A.)

[0567]

Table 29

Example No.	EC ₅₀ (μM)	E _{max} (%)
29	0.8	54
40	0.05	74
41	0.033	57
42	0.031	90
43	0.3	94
44	1.047	94
45	0.115	100
46	0.1	94
48	0.36	87
51	3.277	87
52	0.783	84
53	1.149	87
54	0.63	94
62	0.102	99
65	0.8	85
66	0.023	98
73	0.182	98
75	0.077	87
81	0.365	85
96	0.268	88
orexin-A	0.001	100

[0568]

5 (As used herein, E_{max} in Table 29 is a value by conversion with orexin-A as a full agonist (maximum value of agonist activity: 100%)).

As is clear from the results of Table 25 to Table 29, the compound of the present invention has an OX2R agonist activity.

10 Industrial Applicability

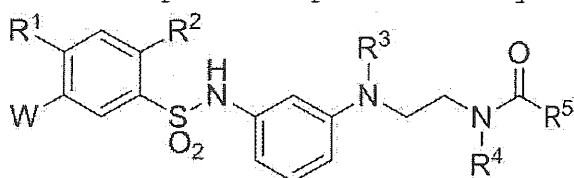
[0569]

The compound of the present invention shows an orexin agonist activity, and is useful as an agent for the treatment or prophylaxis of narcolepsy.

15 This application is based on a patent application No. 2013-257523 filed in Japan (filing date: December 12, 2013), the contents of which are incorporated in full herein.

Claims

1. A compound represented by the formula (I)



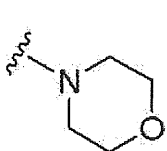
(I)

5 wherein

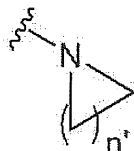
R^1 is a hydrogen atom, alkyl having 1 to 4 carbon atoms or halogen,

R^2 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, $-OH$, $-NR^{2a}R^{2b}$ (wherein R^{2a}

10 is a hydrogen atom, alkyl having 1 to 6 carbon atoms or cycloalkyl having 3 to 8 carbon atoms, R^{2b} is a hydrogen atom, alkyl having 1 to 6 carbon atoms or cycloalkyl having 3 to 8 carbon atoms), or the formula (a) or (b)



(a)



(b)

15 (wherein n' is an integer of 1 to 4)

R^3 is a hydrogen atom or alkyl having 1 to 4 carbon atoms,

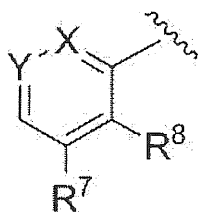
R^4 is a hydrogen atom or alkyl having 1 to 4 carbon atoms,

R^5 is aryl or heteroaryl (wherein aryl or heteroaryl is optionally substituted by optionally selected 1 to 4 R^6),

20 R^6 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, $-CN$, $-CF_3$, $-CH_2F$, $-CHF_2$, $-OCF_3$, $-OH$, $-NO_2$ or $-NR^{6a}R^{6b}$ (wherein R^{6a} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-CH_2CH_2OMe$ or $-CH_2CH_2OEt$, R^{6b} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, -

25 CH_2CH_2OMe or $-CH_2CH_2OEt$),

W is the formula (II):



(II)

(wherein

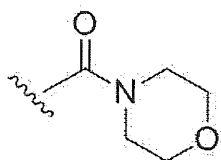
R^7 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms or halogen,

R^8 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms or halogen,

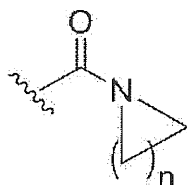
X is $-N=$ or $-CR^9=$,

Y is $-N=$ or $-CR^9=$,

R^9 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, $-OH$, $-NR^{10}R^{11}$, $-CH_2OR^{10}$, $-CF_3$, $-OCF_3$, $-CN$, $-C(O)OR^{10}$, $-C(O)NR^{10}R^{11}$, the formula (III) or (IV),



(III)



(IV)

(wherein n is an integer of 1 to 4)

X and Y are $-CR^9=$, each R^9 may be the same or different,

R^{10} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-CH_2CF_3$, $-CH_2CH_2OH$ or $-CH_2CH_2OMe$,

R^{11} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-CH_2CF_3$, $-CH_2CH_2OH$ or $-CH_2CH_2OMe$,

or a pharmaceutically acceptable acid addition salt thereof.

20

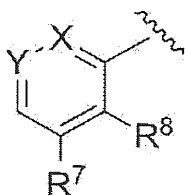
2. The compound according to claim 1, wherein

R^1 is a hydrogen atom or alkyl having 1 to 4 carbon atoms,

R^2 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms or halogen,

R^6 is alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, $-CN$, $-CF_3$, $-OH$ or $-NR^{6a}R^{6b}$ (wherein R^{6a} is alkyl having 1 to 4 carbon atoms, R^{6b} is alkyl having 1 to 4 carbon atoms),

W is a compound of the formula (II):



(II)

30

(wherein

R^7 is a hydrogen atom, alkoxy having 1 to 4 carbon atoms or

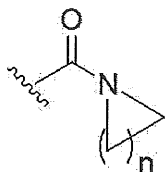
halogen,

R^8 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms or halogen,

X is $-CR^9=$ or $-N=$,

5 Y is $-CR^9=$ or $-N=$,

R^9 is a hydrogen atom, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, $-OH$, $-NR^{10}R^{11}$, $-CH_2OH$, $-C(O)OR^{10}$, $-C(O)NR^{10}R^{11}$ or the formula (IV):



(IV)

10 (wherein n is an integer of 1 to 4),

R^{10} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-CH_2CF_3$, $-CH_2CH_2OH$ or $-CH_2CH_2OMe$, and

R^{11} is a hydrogen atom, alkyl having 1 to 4 carbon atoms, $-CH_2CF_3$, $-CH_2CH_2OH$ or $-CH_2CH_2OMe$,

15 or a pharmaceutically acceptable acid addition salt thereof.