A fused benzamide compound represented by the following formula (the symbols in the formula have the same meanings as defined in the description), which inhibits VR1 activity, or a salt of the compound. The compound and salt have the excellent function of inhibiting VR1 activity and are effective in the treatment of diseases in which vanilloid receptor 1 (VR1) activity participates, such as pains, acute pains, chronic pains, neuropathic pains, pains of chronic articular rheumatism, and neuralgia.
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(54) Title: FUSED BENZAMIDE COMPOUND AND VANILLOID RECEPTOR 1 (VR1) ACTIVITY INHIBITOR

(54) 発明の名称: 綜合ベンザミド化合物及びパニリド受容体1型 (VR 1) 活性阻害剂

(57) Abstract: A fused benzamide compound represented by the following formula (the symbols in the formula have the same meanings as defined in the description), which inhibits VR1 activity; or a salt of the compound. The compound and salt have the excellent function of inhibiting VR1 activity and are effective in the treatment of diseases in which vanilloid receptor 1 (VR1) activity participates, such as pains, acute pains, chronic pains, neuropathic pains, pains of chronic articular rheumatism, and neuralgia.

(57) 要約: 下記式で表されるVR1活性阻害作用を有する総合ベンザミド化合物（式中の記号は明細書と同義）又はその塩は、これら化合物又はその塩が優れたVR1活性阻害作用を有し、疼痛、急性疼痛、慢性疼痛、神経障害性疼痛、慢性関節リウマチ症、神経痛などパニロイド受容体1型 (VR1) 活性が関与する疾患の治療に有効である。
添付公開書類：
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JUMBO APPLICATIONS / PATENTS

THIS SECTION OF THE APPLICATION / PATENT CONTAINS MORE THAN ONE VOLUME.

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SPECIFICATION

FUSED BENZAMIDE COMPOUND AND VANILLOID RECEPTOR 1 (VR1) ACTIVITY INHIBITOR

TECHNICAL FIELD

The present invention relates to a novel condensed benzamide compound having an inhibitory effect on vanilloid receptor subtype 1 (VR1) activity, and a pharmaceutical composition comprising the compound as an active ingredient, particularly a remedy of pain.

BACKGROUND ART

Capsaicin, which is the main ingredient of red pepper, is a pungency causing ingredient as well as a pain producing substance. It has been reported that many nociceptive nerves, particularly unmyelinated C fibers have capsaicin sensitivity and it is known that C fibers will selectively drop out when capsaicin is administered to an infant rodent. It has been also reported that there are many sites of action for capsaicin distributed in the skin, cornea, and oral mucosa, and the distribution thereof is also observed in the muscles, joints and internal organs, particularly in the cardiovascular system, respiratory system and bladder urinary tract system, and it is important for autonomic nerve reflex. In addition, capsaicin sensitivity is also observed in the nerves of the
preoptic area of the thalamus, and involvement in the regulation of body temperature is presumed. Depolarization by inflow of Na⁺ and Ca²⁺ by capsaicin administration is observed in the nociceptive nerves and discharge of glutamic acid and neuropeptides (mainly Substance P and calcitonin gene-related peptide) from the center side end of the primary afferent fiber of the spinal dorsal horn is resulted. Now that specific binding activity of resiniferatoxin (RTX) which brings about similar effects to that of capsaicin has been observed, and that capsazepine has been revealed as a competitive inhibitor, liposoluble capsaicin is considered to act on receptor protein (see Szallasi A, Blumberg PM. (1999) Pharmacol. Rev. 51, 159-212).

The capsaicin receptor gene was cloned in 1997 (see, for example, Caterina MJ, Schumacher MA Tominaga M, Posen TA, Levine JD, Julius D. (1997) Nature 389, 816-824). It was presumed from its amino acid sequence that it was an ion channel having a six-transmembrane domain. Since capsaicin has a vanillyl group in the structure, it is generically referred to as vanilloids along with its analogs such as RTX, and the cloned receptor was named vanilloid receptor subtype 1 (hereinafter referred to as VR1; This VR1 may be also referred to as TRPV1 (transient receptor potential vanilloid receptor 1)). Then, electrophysiological functional analysis using the patch clamping method has been performed by making oocytes of Xenopus laevis and human derived cultured cells to express VR1, and it has been revealed that VR1 is directly activated
by capsaicin, without mediated by an intracellular second
messenger (see, for example, Caterina MJ, Schumacher MA, Tominaga
and that VR1 is a non-selective cation ion channel having high
Ca\textsuperscript{2+} permeability with an outward rectification property (see,
545, 107-117).

Although capsaicin is a pain causing substance, it is used
as an analgesic agent to mitigate pain in diabetic neuropathy
or rheumatic neurosis (see, for example, Szallasi A, Blumberg
PM. (1999) Pharmacol. Rev. 51, 159-212). It is understood that
such mitigation is resulted from a phenomenon that the sensory
nerve end exposed to capsaicin stops answering to pain stimulus,
that is, desensitization. Although it is considered that the
desensitization mechanism of VR1 involves Ca\textsuperscript{2+}-mediated
regulation, regulation depending on potential, activity control
of VR1 by phosphorylation and dephosphorylation, etc., many
points remain unclear.

As well as capsaicin, heat and acid also cause pain and
it is known that the capsaicin sensitive nociceptive nerves
respond to two or more types of stimulation. It was found that
VR1 was directly activated by not only capsaicin but heat
stimulation of 43\textdegree C or more (see, for example, Yang D, Gereau
43\textdegree C is mostly in agreement with the temperature threshold which
causes a pain in humans and animals, suggesting that VR1
participates in nociceptive heat stimulation receptance.
Acidification occurs in an organ in the case of inflammation or ischemia and it is known to cause or enhance pain (see, for example, Bevan S, Geppetti P. (1994) Trends Neurosci. 17, 509-512). It has turned out that when the pH outside cells is reduced within the limits of the acidification which takes place in the case of an organ lesion, VR1 can be directly activated by the acidification (proton) alone, and it is surmised that VR1 is the actual molecule which receives stimulation by acidification in an organ which takes place in the case of inflammation or ischemia (see, for example, Yang D, Gereau RW 4th. (2002) J. Neurosci. 22, 6388-6393).

Immunohistological analysis using a specific antibody has confirmed that the number of unmyelinated C fibers expressing VR1 increases in an inflamed region as compared in a normal region (see, for example, Carlton SM, Coggeshall RE. (2001) Neurosci. Lett. 310, 53-56). The enhancement of VR1 expression in submucosal plexus has been actually observed in human inflammatory bowel disease (see, for example, Yianguo Y, Facer P, Dyer NH, Chan CL, Knowles C, Williams NS, Anand P. (2001) Lancet 357, 1338-1339). Such an increase in the amount of VR1 expression causes peripheral sensitization in an inflamed organ and presumably contributes to duration of inflammatory hyperalgesia.

It has been also reported that extracellular ATP, bradykinin and a neuro growth factor which are inflammation related substances increase VR1 activity (see, for example, Tominaga M, Wada M, Masu M. (2001) Proc. Natl. Acad. Sci. USA

The sensory nerve cells in a VR1-deficient mouse responded to none of capsaicin, proton and heat stimulation. It is also reported that in action analysis, VR1-deficient mouse does not show the pain reaction following capsaicin administration, and sensitivity to heat stimulation decreases and inflammatory hyperalgesia is not observed (see, for example, Caterina MJ, Leffler A, Malmberg AB, Martin WJ, Trafton J, Peterson-Zeitz KR, Koltzenburg M, Basbaum AI, Julius D. (2000) Science 288, 306-313 and Davis LB, Gray J, Gunthorpe MJ et al. (2000) Nature 405, 183-187). Thus, it has been confirmed also on an individual level from the analysis of VR1-deficient mouse that VR1 functions as a wide range pain stimulation receptor.

Moreover, as for the relation between vanilloid receptor subtype 1 (VR1) and a disease, it has been reported already that a substance which inhibits VR1 activity is useful as a therapeutical agent of various diseases.

Particularly with regard to a therapeutical agent of pain, there is a report that capsazepine which is known as a VR1 antagonist has exhibited a significant analgesic effect in
an animal model (see, for example, Ikeda Y, Ueno A, Naraba H, Oh-ishi S, (2001) Life Science 69, 2911-2919), and use is expected as a new therapeutical agent of pain having an inhibitory effect of VR1 activity.

It has been confirmed with regard to bladder hyperstrain type frequent urination and urinary incontinence that the bladder contraction function of VR1-deficient mouse decreases and there is a report that a compound having a capsaicin-like pharmacological mechanism or a compound having an inhibitory action on VR1, i.e., a compound inhibiting vanilloid receptor subtype 1 (VR1) activity is useful for improving bladder function, for example, as a therapeutical agent of frequent urination, urinary incontinence, etc (see, for example, (2002) Nat. Neurosci. 5, 856-860).

In addition, another reference reports that a substance having an inhibitory effect to the vanilloid receptor subtype 1 (VR1), particularly antagonist of VR1 receptor is useful for preventing and treating diseases related to VR1 activity, particularly urgent urinary incontinence, overactive bladder, chronic pain, neuropathic pain, postoperative pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, nerve damage, ischemic symptom, neurodegenerative, cerebral apoplexy, incontinence, inflammatory disease, urgent urinary incontinence (UUI) and/or conditions and diseases including overactive bladder (see, for example, JP 2003-192673).
Furthermore, it is also known that diseases relevant to the vanilloid receptor activity may include pain, acute pain, chronic pain, neuropathic pain, postoperative pain, migraine, joint pain, neuropathy, nerve damage, diabetic nervous disease, neurodegenerative disease, neurogenic skin disorder, cerebral apoplexy, bladder hypersensitivity, irritable bowel syndrome, abnormalities in respiratory organs such as asthma and chronic obstructive pulmonary disease, stimulation of skin, eye or mucosa, fever, stomach or duodenal ulcer, inflammatory bowel disease, inflammatory disease, etc (see, for example, JP 2004-506714 T2).

Accordingly, it can be said that substances having vanilloid receptor subtype 1 (VR1) antagonistic activity is useful as a therapeutic agent for conditions in which C fibers participates, for example, not to mention pruritus, allergic and allergic rhinitis, overactive bladder type frequent urination and urinary incontinence, apoplexy, irritable bowel syndrome, respiratory ailment such as asthma and chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer, inflammatory bowel disease, etc. but also pain, acute pain, chronic pain, neuropathic pain, postoperative pain, migraine, joint pain, neuropathy, nerve damage, diabetic nervous disease, neurodegenerative disease, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, neurogenic skin disorder, apoplexy, overweight, urgent urinary incontinence, ischemic symptom and an inflammatory disease, etc.
Next, compounds considered to relatively resemble the known vanilloid receptor subtype 1 (VR1) antagonist and the compound of present invention are described.

The amide-type compounds represented by the following general formula [A], [B] and [C] are disclosed in W003/068749 as compounds exhibiting antagonism to VR1.

![Chemical Structure A]

![Chemical Structure B]

![Chemical Structure C]

The urea-type compound represented by the following general formula [D] is disclosed in W003/080578 as a compound exhibiting antagonism to VR1.
Quinuclidine-3'-yl
1-phenyl-1,2,3,4-tetrahydroisoquinoline-2-carboxylate is disclosed as a compound exhibiting an inhibitory effect against capsaicin-induced extravasation of a plasma protein in the bladder is disclosed in WO03/006019.

The urea-type compound represented by the following general formula [E] is disclosed in WO03/053945 as a compound exhibiting antagonism to VR1.

The compound represented by the following general formula [F] is disclosed as a compound in WO03/099284 as a compound exhibiting binding activity to VR1.
However, these compounds are different from the compound of the present invention in the structure, and there can be found no description which suggests the compound of the present invention.

DISCLOSURE OF THE INVENTION

As an analgesic agent, narcotic analgesics (morphine etc.), nonnarcotic analgesics (NSAID (nonsteroidal anti-inflammatory drug)), etc. are mainly used now. However, use of narcotic analgesics is severely restricted due to development of resistance/dependency and other serious side effects. It is known well other that an upper gastrointestinal tract disorder and a liver disorder frequently occur during long-term administration of nonnarcotic analgesics, and analgesic agent with a few side effects with higher analgesic effect is eagerly desired. Furthermore, as for diabetes-induced neuropathic pain, postherpetic neuralgia, and neuropathic pain such as trigeminal neuralgia, no effective analgesic agent has been found yet and development of an effective analgesic agent thereof is also expected.

Capsaicin-like compounds which act on VR1 are considered to develop the analgesic effect based on a pharmacological
mechanism completely different from those of existing analgesic agents (desensitization of capsaicin-sensitive nerves), and the efficacy is greatly expected as a therapeutic agent for neuropathic pain and the pain which originates in various conditions such as rheumatic arthritis for which the existing analgesic agents are not effective.

The fact that the final target of various inflammation related substances is VR1 suggests possibility that an agent which acts on VR1 is effective for various inflammatory pains and interstitial cystitis and its efficacy is greatly expected as an analgesic agent which replaces the existing analgesic agents.

Therefore, the purpose of the present invention is to provide a new analgesic agent based on the pharmacological mechanism completely different from those of existing analgesic agents (desensitization of capsaicin-sensitive nerves), i.e., VR1 activity inhibitor.

As a result of intensive study for developing an analgesic agent based on new action mechanism which will replace conventional analgesic agents such as nonnarcotic analgesics, pyrazolone analgesics, non-pyrazolone analgesics and NSAIDs, the present inventors have found out a condensed benzamide compound which has excellent inhibitory effect on VR1, and completed the present invention. The present invention is described in more detail below.
1. A condensed benzamide compound represented by the following general formula [1] or a pharmaceutically acceptable salt thereof:

![Chemical Structure](image)

[wherein $Z$ is

(1) $\cdot O\cdot$,  
(2) $\cdot NR^5\cdot$ (wherein $R^5$ is a hydrogen atom or a C1-6 alkyl group),  
(3) $\cdot S\cdot$,  
(4) $\cdot SO\cdot$ or  
(5) $\cdot SO_2\cdot$;  

$l$ is 0, 1 or 2;  
$m$ is 0, 1 or 2;  

$R^1$ is

(1) a hydrogen atom or  
(2) a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the following Group A:

Group A:

(a) a halogen atom,  
(b) a hydroxyl group,  
(c) a C1-6 alkoxy group,  
(d) a carboxyl group,
(e) a C1-6 alkoxy carbonyl group,

(f) CONR^6R^7 (wherein R^6 and R^7 are the same or different and each represents a hydrogen atom, a C1-6 alkyl group or an acyl group and said alkyl group may be substituted with a hydroxyl group or an acyloxy group),

(g) NR^6R^7 (wherein R^6 and R^7 are the same as above),

(h) NR^6COR^7 (wherein R^6 and R^7 are the same as above),

(i) NR^6CONR^6R^7 (wherein R^6 and R^7 are the same as above, and R^8 is a hydrogen atom or a C1-6 alkyl group); and

(j) NR^6SO_2R^8 (wherein R^6 is the same as above, and R^8 is a C1-6 alkyl group);

R^2 is

(1) a hydrogen atom,

(2) a hydroxyl group,

(3) a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A (wherein the Group A is the same as above),

(4) a carboxyl group,

(5) a C1-6 alkoxy carbonyl group or

(6) CONR^{10}R^{11} (wherein R^{10} and R^{11} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group); or R^2 together with R^1 forms =O;

R^3

(1) a hydrogen atom, or

(2) a C1-6 alkyl group;

R^4 is

(1) a hydrogen atom,
(2) a halogen atom
(3) a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the following Group B,
(4) a C1-6 alkoxy group which may be substituted with 1 to 5 substituents selected from the following Group B,
(5) a cycloalkyl group which may be substituted with 1 to 5 substituents selected from the following Group B,
(6) an aralkyl group which may be substituted with 1 to 5 substituents selected from the following Group B,
(7) an aralkoxy group which may be substituted with 1 to 5 substituents selected from the following Group B, or
(8) a cycloalkylalkoxy group which may be substituted with 1 to 5 substituents selected from the following Group B,

Group B:
(a) a halogen atom,
(b) a halo C1-6 alkyl group,
(c) a hydroxyl group,
(d) a halo C1-6 alkoxy group,
(e) a C1-6 alkoxy carbonyl group,
(f) a C1-6 alkoxy group,
(g) a carboxyl group,
(h) -CONR^{12}R^{13} (wherein R^{12} and R^{13} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group);
(i) -NR^{12}R^{13} (wherein R^{12} and R^{13} are the same as above),
(j) -NR^{12}COR^{13} (wherein R^{12} and R^{13} are the same as above),
(k) -NR^{14}CONR^{12}R^{13} (wherein R^{12} and R^{13} are the same as above, and R^{14} is a hydrogen atom or a C1-6 alkyl group),
(1) \(-\text{SO}_2\text{R}^{15}\) (wherein \(\text{R}^{15}\) is a C1-6 alkyl group), and

(m) \(-\text{NR}^{12}\text{SO}_2\text{R}^{15}\) (wherein \(\text{R}^{12}\) and \(\text{R}^{15}\) are the same as above);

(9) a hydroxyl group,

(10) \(-\text{NR}^{16}\text{R}^{17}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{17}\) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group),

(11) \(-\text{COR}^{18}\) (wherein \(\text{R}^{18}\) is a C1-6 alkyl group, a C1-6 alkoxy group, a cycloalkyl group, an aralkyl group, an aralkoxy group, a cycloalkylalkoxy group or a hydroxyl group),

(12) \(-\text{CONR}^{16}\text{R}^{17}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{17}\) are the same as above),

(13) \(-\text{NR}^{19}\text{CONR}^{16}\text{R}^{17}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{17}\) are the same as above, and \(\text{R}^{19}\) is a hydrogen atom or a C1-6 alkyl group),

(14) \(-\text{NR}^{16}\text{COOR}^{20}\) (wherein \(\text{R}^{16}\) is the same as above, and \(\text{R}^{20}\) is a C1-6 alkyl group or a cycloalkyl group),

(15) \(-\text{SR}^{20}\) (wherein \(\text{R}^{20}\) is the same as above),

(16) \(-\text{SOR}^{20}\) (wherein \(\text{R}^{20}\) is the same as above),

(17) \(-\text{SO}_2\text{R}^{20}\) (wherein \(\text{R}^{20}\) is the same as above),

(18) \(-\text{SO}_2\text{NR}^{16}\text{R}^{17}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{17}\) are the same as above) or

(19) \(-\text{NR}^{16}\text{COR}^{18}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{18}\) are the same as above);

V is

(1) a single bond or

(2) \((-\text{CR}^{21}\text{R}^{22})_{n}\cdot\) (wherein \(\text{R}^{21}\) and \(\text{R}^{22}\) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, and \(n\) is 1 or 2);

P1 and P2 rings are the same or different and each represents

(1) a carbocyclic group which may be substituted with 1 to 5 substituents selected from the following group C or
(2) a heterocyclic group which may be substituted with 1 to 5 substituents selected from the following group, Group C:
(a) a halogen atom,
(b) a hydroxyl group,
(c) a C1-6 alkoxy group which may be substituted with 1 to 5 substituents selected from the Group A,
(d) an C1-6 alkylthio group,
(e) a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A,
(f) -CONR²³R²⁴ (wherein R²³ and R²⁴ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
(g) -NR¹²³₂⁻¹²⁴ (wherein R¹²³ and R¹²⁴ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
(h) -NR²²³COR²²⁴ (wherein R²²³ and R²²⁴ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
(i) -NR²⁵CONR³²³R³²⁴ (wherein R³²³ and R³²⁴ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
(j) -SR²⁶ (wherein R²⁶ is a C1-6 alkyl group),
(k) \(-\text{SOR}^{126}\) (wherein \(\text{R}^{126}\) is a C1-6 alkyl group),
(l) \(-\text{SO}_2\text{R}^{226}\) (wherein \(\text{R}^{226}\) is a C1-6 alkyl group),
(m) \(-\text{NR}^{423}\text{SO}_2\text{R}^{326}\) (wherein \(\text{R}^{423}\) is a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A, and \(\text{R}^{326}\) is a C1-6 alkyl group),
(n) \(-\text{SO}_2\text{NR}^{523}\text{R}^{524}\) (wherein \(\text{R}^{523}\) and \(\text{R}^{524}\) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
(o) \(-\text{COR}^{27}\) (wherein \(\text{R}^{27}\) is a C1-6 alkyl group, C1-6 alkoxy group, a cycloalkyl group, an aralkyl group, an aralkoxy group, a cycloalkylalkoxy group or a hydroxyl group),
(p) a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (o) of the Group C,
(q) a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (o) of the Group C,
(r) \(-\text{O-R}^{28}\) (wherein \(\text{R}^{28}\) is an acyl group, a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (o) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (o) of the Group C),
(s) \(-\text{O-}(\text{CR}^{21}\text{R}^{22})_n\text{R}^{28}\) (wherein \(\text{R}^{21}, \text{R}^{22}, n, \text{and} \text{R}^{28}\) are the same as above),
(t) a nitro group, and
(u) a cyano group.
2. The condensed benzamide compound or pharmaceutically acceptable salt thereof according to above 1 wherein Z is -O-, -NR\(^5\)-, -S- or -SO-.

3. The condensed benzamide compound or pharmaceutically acceptable salt thereof according to above 1 or 2 wherein R\(^3\) is a hydrogen atom or a C1-4 alkyl group.

4. The condensed benzamide compound or pharmaceutically acceptable salt thereof according to above 1 to 3 wherein the P1 ring is a saturated or unsaturated 5-membered or 6-membered heterocyclic ring having 1 to 4 hetero atoms selected from a nitrogen atom, an oxygen atom and a sulfur atom, or a condensed ring of these heterocyclic rings, or a condensed heterocyclic ring of said heterocyclic ring and a carbocyclic ring selected from benzene, cyclopentane and cyclohexane, or a phenyl group (wherein said heterocyclic ring may be substituted with a halogen atom, a hydroxyl group, a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A, and a C1-6 alkoxy group which may be substituted with 1 to 5 substituents selected from the Group A).

5. The condensed benzamide compound or pharmaceutically acceptable salt thereof according to above 4 wherein the P1 ring is a saturated or unsaturated 5-membered or 6-membered heterocyclic ring having at least 1 to 3 nitrogen atoms or a phenyl group (wherein said heterocyclic ring may be substituted with a C1-6 alkyl group which may be substituted with a hydroxyl group or a C1-6 alkyl group, a halogen atom, a hydroxyl group, and a C1-6 alkoxy group which may be
substituted with 1 to 5 substituents selected from the Group A).

6. The condensed benzamide compound or a pharmaceutically acceptable salt thereof according to above 5 wherein the P1 ring is a heterocyclic group selected from a pyridyl group, a pyrazinyl group and a thiazolyl group or a phenyl group (wherein these heterocyclic groups may be substituted with a C1-6 alkyl group which may be substituted with a hydroxyl group, a halogen atom, a hydroxyl group, and a C1-6 alkoxy group which may be substituted with 1 to 5 substituents selected from the Group A).

7. The condensed benzamide compound or a pharmaceutically acceptable salt thereof according to above 1 to 3 wherein the P2 ring is a carbon cyclic group which may be substituted with a substituent group selected from

- a halogen atom,
- a hydroxyl group,
- a C1-6 alkoxy group (wherein said alkoxy group may be substituted with a halogen atom, -CONR^{623}R^{624} (wherein R^{623} and R^{624} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A), a C3-8 cycloalkyl group, a C1-6 alkoxy group, a carboxyl group or a C1-6 alkoxy carbonyl group),
- a C1-6 alkyl group (wherein said alkyl group may be substituted with a halogen atom, a hydroxyl group or a C1-6 alkoxy group),
- NR^{123}R^{124} (wherein R^{123} and R^{124} are the same as above),
- NR²²³COR²²⁴ (wherein R²²³ and R²²⁴ are the same as above),
- COR²⁷ (wherein R²⁷ is a C1-6 alkyl group, C1-6 alkoxy group, a cycloalkyl group, an aralkyl group, an aralkoxy group, a cycloalkylalkoxy group or a hydroxyl group),
- CONR²³R²⁴ (wherein R²³ and R²⁴ are the same as above),
- a heterocyclic group as a saturated or unsaturated substituent group which has 1 to 3 nitrogen atoms as hetero atoms (wherein said heterocyclic group may be substituted with a substituent group selected from a hydroxyl group, CONR³²³R³²⁴ (wherein R³²³ and R³²⁴ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A), a C1-6 alkoxy group, a carboxyl group, a C1-6 alkyl group which may be substituted with a C1-6 alkoxy group, a C1-6 alkoxy carbonyl group and an acyloxy group),
- O-R²⁸ (wherein R²⁸ is an acyl group, a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (i) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (l) of the Group C),
- O-(CR¹²¹R¹²²)ₙ-R¹²⁸ (wherein R¹²¹ and R¹²² are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, n is 1 or 2, and R²⁸ is an acyl group, a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (i) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (l) of the Group C),
- a nitro group and
- a cyano group, or

a heterocyclic group which may be substituted with a substituent as described for the carbocyclic group (wherein said heterocyclic group means a saturated or unsaturated 5-membered or 6-membered heterocyclic ring which has 1 to 4 hetero atoms selected from a nitrogen atom, an oxygen atom and a sulfur atom, or a condensed ring of these heterocyclic rings, or a condensed heterocyclic group of said heterocyclic ring and a carbocyclic ring selected from benzene, cyclopentane and cyclohexane).

8. The condensed benzamide compound or a pharmaceutically acceptable salt thereof according to above 7 wherein the P2 ring as a carbocyclic group is a phenyl group or a cyclohexyl group, or a heterocyclic group as the P2 ring is a thiazolyl group, a pyridyl group, a piperidyl group, a piperidino group, a quinolyl group, a benzo[1,3]dioxo group, a 2,3-dihydrobenzo[1,4]dioxo group or a 1,2,3,4-tetrahydroquinolyl group.

9. The condensed benzamide compound according to above 1 selected from the following group or a pharmaceutically acceptable salt thereof:

1) N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
2) 8-(4-tert-butylphenyl)carbamoyl-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylic acid,
3) N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-3-methyl-1,3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4) N-(4-tert-butylphenyl)-1-(3-chloropyridin-2-yl)-4-methyl-1,1,2,3,4-tetrahydroquinoxaline-5-carboxamide,
5) N-(4-tert-butylphenyl)-9-(3-chloropyridin-2-yl)-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxamide,
6) N-(4-chlorophenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
7) 4-(3-chloropyridin-2-yl)-N-(4-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
8) N-(1-tert-butylpiperidin-4-yl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
9) 4-(3-chloropyridin-2-yl)-N-(4-methylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
10) 4-(3-chloropyridin-2-yl)-N-(trans-4-methylcyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
11) 4-(3-chloropyridin-2-yl)-N-(4-isobutoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
12) N-benzy1-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
13) N-(4-chlorophenyl)methyl-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
14) N-(4-tert-butylphenyl)methyl-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
15) 4-(3-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
16) N-(4-tert-butylphenyl)-4-(pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
17) N-(4-tert-butylphenyl)-4-(3-trifluoromethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
18) N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-2,2-dimethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
19) N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-2-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
20) N-(4-tert-butylphenyl)-6-chloro-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
21) 4-(3-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
22) 4-(3-chloropyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
23) 4-(3-chloropyridin-2-yl)-N-(4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
24) N-(trans-4-tert-butylcyclohexyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
25) 4-(3-chloropyridin-2-yl)-N-(4-chloro-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
26) 4-(3-chloropyridin-2-yl)-N-(4-fluorophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
27) 4-(3-chloropyridin-2-yl)-N-(3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
28) 4-(3-chloropyridin-2-yl)-N-(4-trifluoromethylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
29) 4-(3-chloropyridin-2-yl)-N-(5-trifluoromethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
30) 4-(3-chloropyridin-2-yl)-N-(2-trifluoromethylpyridin-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
31) 4-(3-chloropyridin-2-yl)-N-(4-isopropylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
32) 4-(3-chloropyridin-2-yl)-N-(quinolin-7-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
33) 4-(3-chloropyridin-2-yl)-N-(1-methyl-1,2,3,4-tetrahydroquinolin-7-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
34) 4-(3-chloropyridin-2-yl)-N-(4-methoxycarbonylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
35) 4-[4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxyl]aminobenzoic acid,
36) N-(4-carbamoylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
37) 4-(3-chloropyridin-2-yl)-N-(4-methylcarbamoylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
38) 4-(3-chloropyridin-2-yl)-N-(4-dimethylcarbamoylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
39) N-(4-acetylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
40) 4-(3-chloropyridin-2-yl)-N-[4-(1-hydroxy-1-methyl)ethylphenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
41) 4-(3-chloropyridin-2-yl)-N-[4-(1-hydroxy-1-methyl)propyphenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
42) 4-(3-chloropyridin-2-yl)-N-(1-isobutyrylpyridin-4-yl)phenyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
43) N-(trans-4-tert-butoxycyclohexyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
44) 4-(3-chloropyridin-2-yl)-N-(3-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
45) 4-(3-chloropyridin-2-yl)-N-(3-isobutyloxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
46) N-(4-tert-butylphenyl)-4-(pyridin-4-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
47) N-(4-tert-butylphenyl)-4-(3-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
48) 4-(3-chloropyridin-2-yl)-N-(4-piperidinophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
49) 4-(3-chloropyridin-2-yl)-N-(4-dimethylaminophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
50) 4-(3-chloropyridin-2-yl)-N-(4-morpholinophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
51) 4-(3-chloropyridin-2-yl)-N-(3-fluoro-4-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
52) 4-(3-chloropyridin-2-yl)-N-(2-fluoro-4-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
53) 4-(3-chloropyridin-2-yl)-N-(4-fluoro-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
54) 4-(3-chloropyridin-2-yl)-N-(4-dimethylamino-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
55) 4-(3-chloropyridin-2-yl)-N-(4-isopropoxy-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
56) 4-(3-chloropyridin-2-yl)-N-(4-methoxy-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
57) 4-(3-chloropyridin-2-yl)-N-(4-isobutyloxy-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
58) N-(4-tert-butylphenyl)-4-(4-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
59) N-(4-tert-butylphenyl)-4-(6-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
60) N-(4-tert-butylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
61) N-(4-tert-butylphenyl)-4-(pyridin-3-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
62) N-(4-cyanophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
63) N-(3-amino-4-chlorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
64) N-(3-acetamido-4-chlorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
65) N-[4-(4-carbamoylpiperidin-1-yl)-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
66) N-[3-fluoro-4-(4-methylcarbamoylpiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, 
67) N-[4-(4-dimethylcarbamoylpiperidin-1-yl)-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, 
68) N-[4-(4-ethoxypiperidin-1-yl)-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, 
69) N-[3-fluoro-4-(4-isopropanoylpiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, 
70) N-[3-fluoro-4-(3-methoxypiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, 
71) N-[3-fluoro-4-(4-methoxymethylpiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, 
72) N-[3-fluoro-4-(3-methoxypyrrolidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, 
73) N-[4-isobutoxy-3-methoxycarbonylphenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, 
74) 2-isobutoxy-[[4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carbonyl]amino]benzoic acid,
75) N-(3-carbamoyl-4-isobutylxoyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
76) N-(4-isobutylox-3-methylcarbamoylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
77) N-(3-dimethylcarbamoyl-4-isobutylxoyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
78) N-(4-acetylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
79) N-[4-(1-hydroxy-1-methyl)ethylphenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
80) N-(3-acetyl-4-chlorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
81) N-[4-chloro-3-(1-hydroxy-1-methyl)ethylphenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
82) N-[4-(1-methoxy-1-methyl)ethylphenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
83) N-(3-chloro-4-methoxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
84) 4-(5-methylpyridin-2-yl)-N-(4-propoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
85) N-(3-fluoro-4-propoxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
86) N-(4-ethoxy-3-fluorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
87) N-(3-ethoxy-4-methylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
88) N-(3-carbamoylmethoxy-4-methylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
89) N-(3-methoxyethoxy-4-methylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
90) N-[3-fluoro-4-(2-methoxymethylpiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
91) N-(4-chlorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
92) N-(3-fluoro-4-methylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
93) N-(2-chloropyridin-5-yl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
94) 4-(3-chloropyridin-2-yl)-N-(4-ethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
95) 4-(3-chloropyridin-2-yl)-N-(3-fluoro-4-piperidinophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
96) 4-(3-chloropyridin-2-yl)-N-(trans-4-ethoxycyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
97) 4-(3-chloropyridin-2-yl)-N-(trans-4-isopropoxycyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
98) 4-(3-chloropyridin-2-yl)-N-(trans-4-cyclopent oxy cyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
99) 4\-(3\-chloropyridin-2-yl)\-N\-(trans\-4\-cyclohexyloxy)cyclohexyl\)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
100) 4\-(3\-chloropyridin-2-yl)\-N\-(4\-trifluoromethylphenyl)\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
101) 4\-(3\-chloropyridin-2-yl)\-N\-[3\-fluoro-4\-(4\-methoxypiperidin-1-yl)phenyl]\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
102) 4\-(3\-chloropyridin-2-yl)\-N\-[4\-(4\-ethoxypiperidin-1-yl)\-3\-fluorophenyl]\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
103) 4\-(3\-chloropyridin-2-yl)\-N\-[3\-fluoro-4\-(4\-isopropoxypiperidin-1-yl)phenyl]\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
104) 4\-(3\-chloropyridin-2-yl)\-N\-(3\-fluoro-4\-isobutyloxyphenyl)\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
105) N\-(trans\-4\-ethoxycyclohexyl)\-4\-(5\-methylpyridin-2-yl)\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
106) N\-(trans\-4\-isopropoxycyclohexyl)\-4\-(5\-methylpyridin-2-yl)\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
107) N\-(trans\-4\-cyclohexyloxy)cyclohexyl\)-4\-(5\-methylpyridin-2-yl)\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
108) N\-(trans\-4\-aminocyclohexyl)\-4\-(5\-methylpyridin-2-yl)\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
109) N\-(1,4\-dioxa\-spiro[4,5]deca-8-yl)\-4\-(5\-methylpyridin-2-yl)\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
110) 4\-(5\-methylpyridin-2-yl)\-N\-(4\-oxocyclohexyl)\-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
111) 4-(5-methylpyridin-2-yl)-N-(cis-4-morpholinocyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
112) N-(trans-4-dimethylaminocyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
113) N-(trans-4-diethylaminocyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
114) 4-(5-methylpyridin-2-yl)-N-[cis-4-(pyrrolidin-1-yl)cyclohexyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
115) 4-(5-methylpyridin-2-yl)-N-[trans-4-(pyrrolidin-1-yl)cyclohexyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
116) 4-(5-methylpyridin-2-yl)-N-(4-piperidinocyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
117) 4-(5-methylpyridin-2-yl)-N-(cis-4-morpholinocyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
118) N-(trans-4-acetamidocyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
119) N-(trans-4-cyclohexyloxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
120) 4-(5-chloropyridin-2-yl)-N-(4-ethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
121) N-(4-chlorophenyl)-4-(5-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
122) 4-(5-chloropyridin-2-yl)-N-[3-fluoro-4-(4-methoxypridin-1-yl)phenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
123) 4-[(5-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
124) 4-[(5-methylpyridin-2-yl)-N-(2-phenylethyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
125) N-[2-[(4-chlorophenyl)ethyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
126) N-[2-[(3-chlorophenyl)ethyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
127) N-[2-[(2-chlorophenyl)ethyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
128) 6-[(8-(4-trifluoromethylphenyl)carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
129) 4-[(3-chloro-5-[(1-hydroxy-1-methyl)ethyl]pyridin-2-yl]-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
130) 4-[(3-chloro-5-[(1-hydroxyethyl)pyridin-2-yl]-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
131) 5-chloro-6-[(8-[(3-fluoro-4-trifluoromethylphenyl)carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
132) N-(4-tert-butylphenyl)-4-(5-methoxycarbonylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
133) 6-[(8-(4-tert-butylphenyl)carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
134) 5-chloro-6-[(8-(4-trifluoromethylphenyl)carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
135) 5-chloro-6-[8-(4-tert-butylphenyl)carbamoyl-2,3-dihydrobenzo[1,4]oxazin-4-yl]nicotinic acid,
136) 4·(5-acetyl-3-chloropyridin-2-yl)-N·(4-trifluoromethylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
137) N·(4-tert-butylphenyl)·4·(5-methylcarbamoylpyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
138) N·(4-tert-butylphenyl)·4·(5-carbamoylpyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
139) N·(4-tert-butylphenyl)·4·(5-diethylaminopyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
140) N·(4-tert-butylphenyl)·4·(5-nitropyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
141) 4·(5-aminopyridin-2-yl)·N·(4-tert-butylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
142) 4·(5-acetamidopyridin-2-yl)·N·(4-tert-butylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
143) 4·(5-methoxymethylpyridin-2-yl)·N·(4-trifluoromethylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
144) 4·(5-ethoxypyridin-2-yl)·N·(4-trifluoromethylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
145) 4·(3-chloro-5-methoxypyridin-2-yl)·N·(4-trifluoromethylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
146) 4·(5-hydroxypyridin-2-yl)·N·(4-trifluoromethylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
147) 4·(5-methoxypyridin-2-yl)·N·(4-trifluoromethylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
148) N-(4-tert-butylphenyl)-4-(4-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
149) 4-(5-fluoropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
150) 4-(3,5-difluoropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
151) N-(4-tert-butylphenyl)-4-(3-methoxypyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
152) N-(4-tert-butylpiperidin-1-yl)-4-(5-methylpyridin-2-y1)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
153) 4-[5-(2-hydroxyethoxy)pyridin-2-yl]-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
154) 4-[3-chloro-5-(2-hydroxyethyl)pyridin-2-yl]-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
155) 4-(5-methylpyridin-2-yl)-N-(4-neopentyloxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
156) N-(3-fluoro-4-piperidinophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
157) N-(3-fluoro-4-morpholinophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
158) N-(3,4-difluorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
159) N-(3-fluoro-4-methoxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
160) N-(4-ethoxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
161) N-[4-(2-oxo-piperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
162) 4-(5-methylpyridin-2-yl)-N-(1-methyl-1,2,3,4-tetrahydropyridin-7-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
163) N-(4-dimethylamino-3-fluorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
164) N-{3-fluoro-4-[N-(2-methoxyethyl)-isopropylamino]phenyl}-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
165) N-{4-[N-(2-acetoxyethyl)-isopropylamino]-3-fluorophenyl}-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
166) N-{3-fluoro-4-[N-(2-hydroxyethyl)-isopropylamino]phenyl}-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
167) N-[4-(4-ethoxycarbonylpiperidin-1-yl)-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
168) 1-(4-{4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carbonyl}amino)phenyl)piperidine-4-carboxylic acid,
169) N-[3-fluoro-4-(4-methoxypiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
170) N-[4-(4-acetoxy)piperidin-1-yl]-3-fluorophenyl]-4-(5-methy1pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
171) N-[3-fluoro-4-(4-hydroxypiperidin-1-yl)phenyl]-4-(5-methy1pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
172) N-(3-methoxy-4-methylphenyl)-4-(5-methy1pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
173) N-[3-(2-hydroxyethoxy)-4-methylphenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
174) N-[4-(1-tert-butoxycarbonylpiperidin-4-yl)oxy-3-fluorophenyl]-4-(5-methy1pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
175) N-[4-(piperidin-4-yl)oxy-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
176) N-(trans-4-ethoxycarbonylmethoxy)cyclohexyl]-4-(5-methy1pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
177) (trans-4-{[4-(5-methy1pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carbonyl]amino}cyclohexyloxy)acetic acid,
178) N-(trans-4-carbamoylmethoxycyclohexyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
179) N-(trans-4-methylcarbamoylmethoxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

180) N-(trans-4-dimethylcarbamoylmethoxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

181) N-[trans-4-(2-hydroxyethyloxy)cyclohexyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

182) N-(trans-4-methoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

183) N-(trans-4-isopropoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

184) N-(cis-4-methoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

185) N-(cis-4-isopropoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

186) N-(trans-4-tert-butoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

187) N-(trans-4-isobutyloxyrcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

188) N-(4,4-dimethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
189) N-[(trans-4-(3-methylbutyloxy)cyclohexyl)-4-(5-methylpyridin-2-yl)\]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
190) N-(trans-4-benzyloxy cyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
191) N-(trans-4-isopropylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
192) N-(trans-4-propylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
193) N-(trans-4-neopentylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
194) N-(4-tert-butylphenyl)-4-(5-methoxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
195) 5-chloro-6-[8-(3-chloro-4-trifluoromethoxyphenyl)carbamoyle-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
196) 5-chloro-6-[8-(4-trifluoromethoxyphenyl)carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
197) 4-(5-methylaminopyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
198) 4-(5-ethoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
199) 4-(3-chloro-5-methoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
200) 4-[5-(2-hydroxyethoxy)pyridin-2-yl]-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
201) 4-(5-hydroxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
202) 4-(5-methoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
203) 4-(3-cyanopyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
204) 4-(3-carbamoylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
205) 4-(3-methylcarbamoylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
206) 4-(3-dimethylcarbamoylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
207) 4-(3-benzylxoycarbonylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
208) 2-[8-(4-trifluoromethoxyphenyl) carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl] nicotinic acid,
209) 4-(3-benzylxopyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
210) 4-(3-hydroxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
211) 4-[3-(2-hydroxyethoxy)pyridin-2-yl]-N-(4-trifluoromethoxyphenyl) -3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
212) 2-[8-(4-trifluoromethoxyphenyl) carbamoyl]-2,3-dihydro-benzo[1,4]oxazin-4-yl] pyridin-3-yl] oxyacetic acid,
213) 4-((3-carbamoylmethoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
214) 4-((3-methylcarbamoylmethoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
215) 4-((3-dimethylcarbamoylmethoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
216) 4-((3-(pyridin-2-yl)methoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
217) 4-((3-(pyridin-3-yl)methoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
218) 4-((5-cyanopyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
219) 4-((5-acetimidomethylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
220) 4-((3-chloro-5-methoxymethylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
221) 4-((5-(N-methylacetamido)methylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
222) 4-((5-aminomethylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
223) 4-\((5\text{-dimethylaminomethylpyridin-2-yl})\text{-N-(4-trifluoromethoxyphenyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide},
224) 4-\((3\text{-chloropyridin-2-yl})\text{-2-methylcarbamoyl-N-(4-trifluoromethoxyphenyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide},
225) 4-\((6\text{-hydroxymethylpyridin-2-yl})\text{-N-(4-trifluoromethylphenyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide},
226) 4-\((3\text{-chloropyridin-2-yl})\text{-2-dimethylcarbamoyl-N-(4-trifluoromethoxyphenyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide},
227) 4-\((5\text{-carbamoyl-3-chloropyridin-2-yl})\text{-N-(4-trifluoromethoxyphenyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide},
228) (S)\text{-3-acetamidomethyl-4-\((3\text{-chloropyridin-2-yl})\text{-N-(4-trifluoromethylphenyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide},
229) (R)\text{-4-\((3\text{-chloropyridin-2-yl})\text{-8-(4-trifluoromethylphenylcarbamoyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-3-carboxylic acid},
230) 4-\((3\text{-chloropyridin-2-yl})\text{-2-methoxycarbonyl-N-(4-trifluoromethoxyphenyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide},
231) 4-\((3\text{-chloropyridin-2-yl})\text{-2-hydroxymethyl-N-(4-trifluoromethoxyphenyl)}\text{-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide},
232) 4·(3-chloropyridin-2-yl)-8·(4-trifluoromethoxyphenylcarbamoyl)·3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylic acid,
233) (S)·3-carbamoyl·4·(3-chloropyridin-2-yl)·N·(4-trifluoromethylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
234) (S)·4·(3-chloropyridin-2-yl)·3-methylcarbamoyl·N·(4-trifluoromethylphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
235) 2-carbamoyl·4·(3-chloropyridin-2-yl)·N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
236) N·(4-chlorophenyl)·N-methyl·4·(5-methylpyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
237) N·(benzo[1,3]dioxol-5-yl)·4·(3-chloro-5-hydroxymethylpyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
238) N·(2,3-dihydrobenzo[1,4]dioxin-6-yl)·4·(3-chloro-5-hydroxymethylpyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
239) 4·(3-chloro-5-hydroxymethylpyridin-2-yl)·N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
240) 4·(3-chloro-5-hydroxymethylpyridin-2-yl)·N·(3-chloro-4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
41) \(4\cdot (3\text{-chboro}-5\text{-hydroxymethylpyridin}-2\text{-yl}) \cdot N\cdot (4\text{-trifluoromethylphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
42) \(4\cdot (3\text{-chloro}-5\text{-hydroxymethylpyridin}-2\text{-yl}) \cdot N\cdot (3\text{-fluoro}-4\text{-trifluoromethylphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
43) \(4\cdot (5\text{-hydroxymethylpyridin}-2\text{-yl}) \cdot N\cdot (4\text{-trifluoromethylphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
44) \(4\cdot (5\text{-hydroxymethylpyridin}-2\text{-yl}) \cdot N\cdot (4\text{-isobutyloxyphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
45) \(4\cdot (3\text{-chloro}-5\text{-methoxymethylpyridin}-2\text{-yl}) \cdot N\cdot (4\text{-trifluoromethylphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
46) \(4\cdot (4\text{-hydroxymethylpyridin}-2\text{-yl}) \cdot N\cdot (4\text{-trifluoromethylphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
47) \(4\cdot (3\text{-hydroxymethylpyridin}-2\text{-yl}) \cdot N\cdot (4\text{-trifluoromethoxyphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
48) (+) \cdot 4\cdot (3\text{-chloro}-5\cdot (1\text{-hydroxyethyl})pyridin-2\text{-yl}) \cdot N\cdot (4\text{-trifluoromethylphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
49) (-) \cdot 4\cdot (3\text{-chloro}-5\cdot (1\text{-hydroxyethyl})pyridin-2\text{-yl}) \cdot N\cdot (4\text{-trifluoromethylphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide, \\
50) (+) \cdot 4\cdot (3\text{-chloro}-5\cdot (1\text{-hydroxyethyl})pyridin-2\text{-yl}) \cdot N\cdot (4\text{-trifluoromethoxyphenyl}) \cdot 3,4\text{-dihydro}-2H\cdot benzophen[1,4]oxazine\cdot 8\cdot carboxamide,
251) (·) - 4 - (3-chloro-5-(1-hydroxyethyl)pyridin-2-yl) - N - (4-
trifluoromethoxyphenyl) - 3,4-dihydro-2H-benzo[1,4]oxazine-
8-carboxamide,
252) 4 - (3-chloro-5-hydroxymethylpyridin-2-yl) - N - (2-triflu-
romethylpyridin-5-yl) - 3,4-dihydro-2H-benzo[1,4]oxazine-8-
carboxamide,
253) N - (4-bromo-3-chlorophenyl) - 4 - (3-chloro-5-hydroxymethy-
lpypiridin-2-yl) - 3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxa-
mide,
254) 4 - [5-(1-hydroxy-1-methyl)ethylpyridin-2-yl] - N - (4-trif-
luoromethylphenyl) - 3,4-dihydro-2H-benzo[1,4]oxazine-8-car-
boxamide,
255) 4 - (3-chloro-5-hydroxymethylpyridin-2-yl) - N - (4-isoprop-
oxy-3-trifluoromethylphenyl) - 3,4-dihydro-2H-benzo[1,4]oxa-
ze-8-carboxamide,
256) 4 - (3-chloro-5-hydroxymethylpyridin-2-yl) - N - (2,3-dichl-
ropyrpidin-5-yl) - 3,4-dihydro-2H-benzo[1,4]oxazine-8-carbo-
oxamid,
257) 4 - (3-chloro-5-hydroxymethylpyridin-2-yl) - N - (3,4,5-tri-
chlorophenyl) - 3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxam-
ide,
258) 4 - (3-chloro-5-hydroxymethylpyridin-2-yl) - N - (6-fluorob-
enzothiazol-2-yl) - 3,4-dihydro-2H-benzo[1,4]oxazine-8-carb-
oxamide,
259) N - (4-bromo-3-trifluoromethylphenyl) - 4 - (3-chloro-5-hyd-
roxymethylpyridin-2-yl) - 3,4-dihydro-2H-benzo[1,4]oxazine-
8-carboxamide,
260) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(2-trifluoromethylbenzimidazol-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
261) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3,4,5-trifluorophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
262) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-fluoro-3-nitrophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
263) N-(3-amino-4-fluorophenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
264) N-(tert-butylphenyl)-4-(5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
265) N-(3,5-bistrifluoromethylphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
266) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-[(2,2,2-trifluoroethoxy)phenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
267) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-[3-chloro-(2,2,2-trifluoroethoxy)phenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
268) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-trifluoromethylmercaptophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
269) N-(trans-4-tert-butylcyclohexyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

270) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3-fluoro-4-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

271) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3-fluoro-4-piperidinophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

272) N-(4-tert-butylphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

273) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-[4-(1-hydroxy-2,2,2-trifluoro-1-trifluoromethyl)ethylphenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

274) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(2,2,3,3-tetrafluoro-2,3-dihydrobenzo[1,4]dioxin-6-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

275) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-isobutoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

276) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(2,3-dichlorophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

277) N-(4-bromo-3-fluorophenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
278) N-(4-bromophenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
279) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3,5-dichlorophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
280) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-difluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
281) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-chlorophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
282) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-isopropylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
283) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
284) 4-(3-chloro-5-(1-hydroxyethyl)pyridin-2-yl)-N-(3-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
285) 4-(3-chloro-pyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-3-oxo-2H-benzo[1,4]oxazine-8-carboxamide,
286) (R)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
287) (R)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
288) (S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
289) (S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
290) (S)-4-(5-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
291) (S)-4-(5-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
292) (S)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
293) (S)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
294) (S)-4-(3-chloropyridin-2-yl)-N-(3-fluoro-4-isopropoxyphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
295) (S)-4-(3-chloropyridin-2-yl)-N-(3-fluoro-4-trifluoromethoxyphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
296) (S)-4-(3-chloropyridin-2-yl)-N-(3-chloro-4-piperidino-phenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
297) (S)-4-(3-chloropyridin-2-yl)-N-(4-dimethylamino-3-fluoro-phenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
298) (S)-4-(3-chloropyridin-2-yl)-N-(3-fluoro-4-methylphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
299) (S)-N-(trans-4-tert-butylcyclohexyl)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
300) (S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(trans-4-neopentyloxy cyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
301) (S)-N-(4-chlorophenyl)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
302) (S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-isopropylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
303) (S)-4-(3-chloropyridin-2-yl)-N-(4-fluoro-3-trifluoromethylphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
304) (S)-N-(4-bromo-3-chlorophenyl)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
305) (S)-4-(3-chloropyridin-2-yl)-N-(4-dimethylamino-3-trifluoromethylphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
306) (S)-4-(3-chloropyridin-2-yl)-N-(4-isoproxy-3-trifluoromethylphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
307) (S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-piperidino-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
308) (S)-4-(3-chloropyridin-2-yl)-N-(4-ethoxy-3-fluorophenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
309) (S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-methoxy-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
310) (S)-4-(3-chloro-5-methoxypyridin-2-yl)-3-hydroxymethyl-1-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
311) (S)-4-(3-chloro-5-methoxypyridin-2-yl)-3-hydroxymethyl-1-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
312) (S)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-N-(2-trifluoromethylpyridin-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
313) (S)-3-hydroxymethyl-N-(4-methoxy-3-trifluoromethylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
314) (S)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-N-(4-piperidino-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
315) (S)·N·(3·fluoro·4·trifluoromethylphenyl)·3·hydroxymethyl·4·(5·methylpyridin-2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
316) (S)·N·(3·chloro·4·trifluoromethoxyphenyl)·3·hydroxymethyl·4·(5·methylpyridin-2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
317) (S)·N·(3,4·dichlorophenyl)·3·hydroxymethyl·4·(5·methylpyridin-2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
318) (S)·4·(3·chloropyridin-2·yl)·3·hydroxymethyl·N·{(4·isobutyloxy·3·trifluoromethylphenyl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
319) (S)·4·(3·chloropyridin-2·yl)·N·{(3,4·dichlorophenyl)·3·hydroxymethyl·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
320) (S)·4·(3·chloropyridin-2·yl)·N·{(4·chloro·3·trifluoromethylphenyl)·3·hydroxymethyl·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
321) (S)·N·{(4·bromo·3·fluorophenyl)·4·(3·chloropyridin-2·yl)·3·hydroxymethyl·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
322) (S)·N·{(4·chloro·3·trifluoromethylphenyl)·3·hydroxymethyl·4·(5·methylpyridin-2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
323) (S)·3·hydroxymethyl·N·{(4·isopropoxy·3·trifluoromethylphenyl)·4·(5·methylpyridin-2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
324) (S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-morpholino-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
325) (S)-N-(2-chloro-4-trifluoromethylphenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
326) (S)-3-hydroxymethyl-4-(3-methylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
327) (S)-3-hydroxymethyl-4-(3-methylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
328) (S)-N-(4-tert-butoxy-3-fluorophenyl)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
329) (S)-4-(3-chloropyridin-2-yl)-N-(3-fluoro-4-isobutylloxynphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
330) (S)-3-hydroxymethyl-N-(4-isobutyl oxy-3-trifluoromethylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
331) (S)-N-(3-fluoro-4-isopropoxyphenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
332) (S)-N-(3-fluoro-4-morpholinophenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
333) (S)·N·(4·chlorophenyl)·3·hydroxymethyl·4·(5·methylpyr
idin·2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·carboxamide,
334) (S)·3·hydroxymethyl·4·(5·methylpyridin·2·yl)·N·(2·morp
holino·4·trifluoromethylphenyl)·3,4·dihydro·2H·benzo[1,4]
oxazine·8·carboxamide,
335) (S)·N·(4·tert·butoxy·3·fluorophenyl)·3·hydroxymethyl·
4·(5·methylpyridin·2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine
·8·carboxamide,
336) (S)·N·(4·bromo·3·fluorophenyl)·3·hydroxymethyl·4·(5·me
thylpyridin·2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·car
boxamide,
337) 4·(3·chloropyridin·2·yl)·3·hydroxymethyl·N·(4·triflu
romethoxyphenyl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·carbo
xamide,
338) (S)·4·(3·chloropyridin·2·yl)·N·(3·fluoro·4·trifluorom
ethylphenyl)·3·hydroxymethyl·3,4·dihydro·2H·benzo[1,4]oxa
zine·8·carboxamide,
339) (S)·N·(3·fluoro·4·isobutyloxyphenyl)·3·hydroxymethyl·
4·(5·methylpyridin·2·yl)·3,4·dihydro·2H·benzo[1,4]oxazine
·8·carboxamide,
340) (S)·N·(4·fluoro·3·trifluoromethylphenyl)·3·hydroxymet
hyl·4·(5·methylpyridin·2·yl)·3,4·dihydro·2H·benzo[1,4]oxa
zine·8·carboxamide,
341) 4·(3·chloropyridin·2·yl)·3·(1·hydroxyethyl)·N·(4·trif
luoromethoxyphenyl)·3,4·dihydro·2H·benzo[1,4]oxazine·8·ca
boxamide,
342) (R) - 9 - (3 - chloropyridin - 2 - yl) - 7 - hydroxy - N - (4 - trifluoromethoxyphenyl) - 6, 7, 8, 9 - tetrahydro - 5 - oxa - 9 - azabenzocycloheptane - 4 - carboxamide,
343) (R) - 9 - (3 - chloropyridin - 2 - yl) - 7 - hydroxy - N - (4 - trifluoromethylphenyl) - 6, 7, 8, 9 - tetrahydro - 5 - oxa - 9 - azabenzocycloheptane - 4 - carboxamide,
344) (S) - 9 - (3 - chloropyridin - 2 - yl) - 7 - hydroxy - N - (4 - trifluoromethoxyphenyl) - 6, 7, 8, 9 - tetrahydro - 5 - oxa - 9 - azabenzocycloheptane - 4 - carboxamide,
345) N - (4 - tert - butylphenyl) - 4 - (pyrazin - 2 - yl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
346) N - (4 - chlorophenyl) - 4 - (5 - ethylpyrimidin - 2 - yl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
347) N - (4 - ethoxyphenyl) - 4 - (5 - ethylpyrimidin - 2 - yl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
348) 4 - (6 - chloropyridazin - 3 - yl) - N - (4 - trifluoromethylphenyl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
349) 4 - (4 - methylthiazol - 2 - yl) - N - (4 - trifluoromethoxyphenyl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
350) 4 - (5 - methylthiazol - 2 - yl) - N - (4 - trifluoromethoxyphenyl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
351) (S) - 3 - hydroxymethyl - 4 - (5 - methylthiazol - 2 - yl) - N - (4 - trifluoromethoxyphenyl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
352) (S) - 3 - hydroxymethyl - 4 - (5 - methylthiazol - 2 - yl) - N - (4 - trifluoromethylphenyl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
353) (S)-N-(3,4-dichlorophenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
354) (S)-N-(3-fluoro-4-isopropoxyphenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
355) (S)-N-(3-fluoro-4-tert-butoxyphenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
356) (S)-N-(3-fluoro-4-isobutoxyphenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
357) (S)-N-(4-bromo-3-fluorophenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
358) (S)-N-(4-fluoro-3-trifluoromethylphenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
359) (S)-N-(3-fluoro-4-morpholinophenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
360) (S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-N-(2-trifluoromethylpyridin-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
361) (S)-3-hydroxymethyl-4-(5-methyloxazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
(S)-3-hydroxymethyl-4-(5-methyloxazol-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

(S)-4-(4,5-dimethylthiazol-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

(S)-4-(4,5-dimethylthiazol-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

(S)-3-hydroxymethyl-4-(5-methyl[1,3,4]thiadiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

(S)-3-hydroxymethyl-4-(5-methyl[1,3,4]thiadiazol-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

4-(4,5-dimethylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxamide,

4-(4,5-dimethylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-1-oxo-3,4-tetrahydro-benzo[1,4]thiazine-8-carboxamide,

N-(4-tert-butylphenyl)-4-(4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

N-(4-isobutylxyloxyphenyl)-4-(4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

N-(4-chlorophenyl)-4-(4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
372) (S)-4-(2-chlorophenyl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
373) (S)-4-(2-chlorophenyl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
374) (S)-4-(2-chlorophenyl)-N-(3-fluoro-4-trifluoromethylphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
375) (S)-4-(4-chlorophenyl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
376) (S)-3-hydroxymethyl-4-(4-methoxyphenyl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
377) (S)-3-hydroxymethyl-4-(4-methoxyphenyl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
378) (S)-4-(4-chlorophenyl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide, and
379) (S)-4-(4-chlorophenyl)-N-(4-chlorophenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide.
10. A pharmaceutical composition comprising a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9 and a pharmaceutically acceptable carrier.
11. A pharmaceutical composition comprising a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9 and a pharmaceutically acceptable carrier for treating and/or preventing a disease selected from pain, acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer and inflammatory bowel disease, bladder hypersensitivity, and overactive bladder type frequent urination and urinary incontinence.

12. A pharmaceutical composition for treating and/or preventing pain comprising a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9 and a pharmaceutically acceptable carrier.

13. The pharmaceutical composition according to above 12 wherein the pain is acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic
neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy or neurodegenerative disease.

14. An inhibitor of vanilloid receptor subtype 1 (VR1) activity comprising a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9 and a pharmaceutically acceptable carrier.

15. A method for treating and/or preventing a disease selected from pain, acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer and inflammatory bowel disease, bladder hypersensitivity, and overactive bladder type frequent urination and urinary incontinence characterized in that the method comprises administering a pharmacologically effective amount of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9.

16. A method for treating and/or preventing pain characterized in that the method comprises administering a
pharmacologically effective amount of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9.

17. The treating and/or preventing method according to above 16 wherein the pain is acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy or neurodegenerative disease.

18. A commercial package comprising a pharmaceutical composition according to any of above 10 to 13 and written instructions concerning this pharmaceutical composition stating that said composition can be used or should be used for treating and/or preventing a disease selected from pain, acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and
duodenal ulcer and inflammatory bowel disease, bladder hypersensitivity, and overactive bladder type frequent urination and urinary incontinence.

19. Use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9 for preparing a pharmaceutical composition according to above 11 for treating and/or preventing a disease selected from pain, acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer and inflammatory bowel disease, bladder hypersensitivity, and overactive bladder type frequent urination and urinary incontinence.

20. Use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9 for preparing a pharmaceutical composition for treating and/or preventing pain according to above 12.

21. The use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to above
20 wherein the pain is acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy or neurodegenerative disease.

22. A drug comprising a combination of a pharmaceutical composition comprising a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9 and a pharmaceutically acceptable carrier with one or more agents selected from the group which consists of an anti-virus agent, an antidepressant, an anticonvulsant, an antiarrhythmic drug, a local anesthetic, an anesthetic drug, an N-methyl-D-aspartate receptor antagonist, adrenal cortical steroid, a nerve block, a nonsteroidal antiinflammatory analgesic, narcotics, an antagonist analgesic, α₂-adrenaline receptor agonist, a medicine for external application, a calcium channel antagonist, and a potassium channel opener.

23. Use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9 for preparing a drug according to above 22.

24. A method for treating and/or preventing a disease selected from pain, acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic
neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer and inflammatory bowel disease, bladder hypersensitivity, and overactive bladder type frequent urination and urinary incontinence characterized in that one or more agents selected from the group which consists of an anti-virus agent, an antidepressant, an anticonvulsant, an antiarrhythmic drug, a local anesthetic, an anesthetic drug, an N-methyl-D-aspartate receptor antagonist, adrenal cortical steroid, a nerve block, a nonsteroidal anti-inflammatory analgesic, narcotics, an antagonist analgesic, $\alpha_2$-adrenal receptor agonist, a medicine for external application, a calcium channel antagonist, and a potassium channel opener are used in combination with a pharmacologically effective amount of an inhibitor of vanilloid receptor subtype 1 (VR1) activity.

25. The treating and/or preventing method according to above 24 wherein the inhibitor of vanilloid receptor subtype 1 (VR1) activity is a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9.
26. A method for treating and/or preventing pain characterized in that the method uses administration of an inhibitor of vanilloid receptor subtype 1 (VR1) activity in combination with stimulation-produced analgesia selected from acupuncture, transcutaneous electroacupuncture stimulation therapy, transcutaneous electrical nerve stimulation therapy, silver spike point (SSP) therapy, peripheral nerve stimulation therapy, spinal cord electrical stimulation therapy, electroconvulsive therapy, laser therapy and low-frequency therapy.

27. The treating and/or preventing method according to above 26 wherein the inhibitor of vanilloid receptor subtype 1 (VR1) activity is a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any of above 1 to 9.

28. A method for treating and/or preventing postoperative neuralgia characterized in that an inhibitor of vanilloid receptor subtype 1 (VR1) activity is administered after performing a surgical operation selected from cicatrectomy, nerve freezing solidification, peripheral nerve excision, spinal cord dorsal root excision, sympathectomy, spinal cord dorsal root entry zone destruction, cordotomy, and frontal lobe excision.

29. The treating and/or preventing method according to above 28 wherein the inhibitor of vanilloid receptor subtype 1 (VR1) activity is a condensed benzamide compound or a
pharmaceutically acceptable salt thereof according to any of above 1 to 9.

The condensed benzamide compound of the present invention effectively inhibits vanilloid receptor subtype 1 (VR1) activity, and therefore it is effective in the medical treatment and/or prevention of diseases such as pain, acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer and inflammatory bowel disease, bladder hypersensitivity, and overactive bladder type frequent urination and urinary incontinence. Particularly, it is effective as a therapeutic agent and preventive agent of diseases accompanied with pain condition such as pain, acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic
neuropathy and neurodegenerative disease. In addition, effects by different mechanism from the conventional analgesics are also expected.

BEST MODE FOR CARRYING OUT THE INVENTION

The definition of each term used in this specification is as follows.

A "C1-6 alkyl group" represents a linear or branched alkyl group having 1 to 6 carbon atoms, and specifically includes a methyl group, an ethyl group, a propyl group, an isopropyl group, a butyl group, an isobutyl group, a sec-butyl group, a tert-butyl group, a pentyl group, an isopentyl group, a tert-pentyl group, a hexyl group, etc. A "C1-4 alkyl group" represents a linear or branched alkyl group having 1 to 4 carbon atoms, and specifically includes a methyl group, an ethyl group, a propyl group, an isopropyl group, a butyl group, an isobutyl group, a sec-butyl group, and a tert-butyl.

A preferred "C1-6 alkyl group" as \( R^1, R^2, R^3, R^4, R^5, R^6, R^7, R^8, R^9, R^{10}, R^{11}, R^{12}, R^{13}, R^{14}, R^{15}, R^{16}, R^{17}, R^{18}, R^{19}, R^{20}, R^{21}, R^{22}, R^{23}, R^{24}, R^{25}, R^{26} \) and \( R^{27} \) is a "C1-4 alkyl group," particularly a methyl group and an ethyl group. A preferred C1-6 alkyl group in a P1 ring and a P2 ring which may be substituted is a C1-4 alkyl group, particularly a methyl group, an ethyl group, a propyl group, an isopropyl group, a butyl group, an isobutyl group, a sec-butyl group and a tert-butyl group.
A "halogen atom" is a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, and a fluorine atom and a chlorine atom are preferred.

As R⁴, preferred is a chlorine atom, and as a halogen atom in a P₁ ring and a P₂ ring which may be substituted, preferred are a chlorine atom and a fluorine atom.

A "halo C₁-₆ alkyl group" is a "C₁-₆ alkyl group" of the above-mentioned definition substituted with "halogen atom" of the above-mentioned definition, and preferably a halogenated alkyl group in which the alkyl group thereof is a linear or branched alkyl group having 1 to 4 carbon atoms. Specifically, it includes a fluoromethyl group, a difluoromethyl group, a trifluoromethyl group, a bromomethyl group, a chloromethyl group, a 1,2-dichloromethyl group, a 2,2-dichloromethyl group, a 2,2,2-trifluoroethyl group, etc.

A "C₁-₆ alkoxy group" is an alkoxy group in which the alkyl part thereof is a "C₁-₆ alkyl group" of the above-mentioned definition. Specifically, it includes a methoxy group, an ethoxy group, a propoxy group, an isoproproxy group, a butoxy group, an isobutoxy group, a s-butoxy group, a tert-butyloxy group, a pentyloxy group, an isopentyloxy group, a 2-methylbutoxy group, an neopentyloxy group, a 1-ethylpropoxy group, a hexyloxy group, a 4-methylpentyloxy group, a 3-methylpentyloxy group, a 2-methylpentyloxy group, a 1-methylpentyloxy group, a 3,3-dimethylbutoxy group, a 2,2-dimethylbutoxy group, a 1,1-dimethylbutoxy group, a 1,2-dimethylbutoxy group, a 1,3-dimethylbutoxy group, a
2,3-dimethylbutoxy group, a 2-ethylbutoxy group, etc. Preferred is an alkoxy group in which the alkyl part thereof is a linear or branched alkyl group having 1 to 4 carbon atoms.

A "halo C1-6 alkoxy group" is a haloalkoxy group in which the "C1-6 alkyl group" which constitutes the C1-6 alkoxy group part thereof is substituted with one or more and the same or different halogen atoms, and, specifically includes a fluoromethoxy group, a difluoromethoxy group, a trifluoromethoxy group, a bromomethoxy group, a chloromethoxy group, a 1,2-dichloromethoxy group, a 2,2-dichloromethoxy group, a 2,2,2-trifluoroethoxy group, etc.

A "C1-6 alkylthio group" is a "C1-6 alkyl group" mentioned above attached to a sulfur atom, and for example, a linear or branched alkylthio group having 1 to 6 carbon atoms such as a methylthio group, ethylthio group, propylthio group, isopropylthio group, butylthio group, isobutylthio group, s-butylthio group, t-butylthio group, pentylthio group, isopentylthio group, 2-methylbutylthio group, neopentylthio group, hexylthio group, 4-methylpentylthio group, 3-methylpentylthio group, 2-methylpentylthio group, 3,3-dimethylbutylthio group, 2,2-dimethylbutylthio group, 1,1-dimethylbutylthio group, 1,2-dimethylbutylthio group, 1,3-dimethylbutylthio group, 2,3-dimethylbutylthio group, and preferably a C1-4 alkylthio group.

A "C1-6 alkoxy carbonyl group" is a linear or branched "C1-6 alkoxy group" attached to a carbonyl group, and specifically includes a methoxycarbonyl group, an
ethoxycarbonyl group, a propoxycarbonyl group, an isopropropoxycarbonyl group, a butoxycarbonyl group, an isobutoxycarbonyl group, a s-butoxycarbonyl group, a t-butoxycarbonyl group, a pentyloxycarbonyl group, an isopentyloxycarbonyl group, a 2-methylbutoxycarbonyl group, a neopentyloxycarbonyl group, a 1-ethyl propoxycarbonyl group, a hexyloxycarbonyl group, a 4-methylpentyloxycarbonyl group, a 3-methylpentyloxycarbonyl group, a 2-methylpentyloxycarbonyl group, a 1-methylpentyloxycarbonyl group, a 3,3-dimethylbutoxycarbonyl group, a 2,2-dimethylbutoxycarbonyl group, a 1,1-dimethylbutoxycarbonyl group, a 1,2-dimethylbutoxycarbonyl group, a 1,3-dimethylbutoxycarbonyl group, a 2,3-dimethylbutoxycarbonyl group, a 2-ethylbutoxycarbonyl group, etc. Preferred is an alkoxy carbonyl group in which the alkoxy group part thereof is a linear or branched alkoxy group having 1 to 4 carbon atoms.

Preferable examples of an "acyl group" include a formyl group; a carboxyl group; a carbamoyl group; a thiocarbamoyl group; a C1-6 alkyl-carbonyl group (e.g. an acetyl group, propionyl group, butyryl group, isobutyryl group, valeryl group, isovaleryl group, pivaloyl group and hexanoyl group); a C2-7 alkenyl-carbonyl group (e.g. a crotonyl group); a C3-8 cycloalkyl-carbonyl group (e.g. a cyclobutanecarbonyl group, cyclopentanecarbonyl group, cyclohexanecarbonyl group and
cycloheptanecarbonyl group); a C3-8 cycloalkenyl-carbonyl
group (e.g. a 2-cyclohexanecarbonyl group); a C6-14
aryl-carbonyl group (e.g. a benzoyl group, 1-naphthoyl group
and 2-naphthoyl group); a C7-14 aralkyl-carbonyl group (e.g.
a benzylcarbonyl group,
phenethylcarbonylphenylpropylcarbonyl group and
phenylbutylcarbonyl group); a C8-13 arylalkenyl-carbonyl
group (e.g. a styrylcarbonyl group); a C8-13
arylalkynyl-carbonyl group (e.g. a phenylethynylcarbonyl
group); an aromatic heterocyclic carbonyl group (e.g. a
nicotinoyl group, isonicotinoyl group, furylcarbonyl group,
thienylcarbonyl group, pyrimidinylcarbonyl group,
benzofuranylcarbonyl group, 1H-indazolylcarbonyl group and
quinolylcarbonyl group); a non-aromatic heterocyclic
carbonyl group (e.g. a pyrrolidinylcarbonyl group,
piperidinocarbonyl group, morpholinocarbonyl group,
thiomorpholinocarbonyl group, piperazinocarbonyl group,
thiazolidinylcarbonyl group, hexamethyleneiminylcarbonyl
group and tetrahydroisoquinolylcarbonyl group); a C1-6
alkoxy-carbonyl group (e.g. a methoxycarbonyl group,
ethoxycarbonyl group, propoxycarbonyl group, butoxycarbonyl
group and tert-butoxycarbonyl group); a C6-14
aryloxy-carbonyl group (e.g. a phenyloxycarbonyl group and
naphthoxy carbonyl group); a C7-14 aralkyloxy-carbonyl
group (e.g. a benzylloxy carbonyl group and
phenethyloxycarbonyl group); a mono or di-C1-6 alkylcarbamoyl
group (e.g. a methylcarbamoyl group, ethylcarbamoyl group,
dimethylcarbamoyl group, diethylcarbamoyl group, ethylmethylcarbamoyl group, propylcarbamoyl group, butylcarbamoyl group, tert-butylcarbamoyl group, pentylcarbamoyl group, hexylcarbamoyl group); a mono or di-C1-6 alkyl-thiocarbamoyl group (e.g. a methylthiocarbamoyl group and ethylthiocarbamoyl group); a C6-14 aryl-carbamoyl group (e.g. a phenylcarbamoyl group); a C3-10 cycloalkyl-carbamoyl group (e.g. a cyclopropylcarbamoyl group, cyclopentylcarbamoyl group and cyclohexylcarbamoyl group); a C7-14 aralkyl-carbamoyl group (e.g. a benzylcarbamoyl group, phenethylcarbamoyl group and diphenylethylcarbamoyl group); a C4-13 cycloalkylalkyl-carbamoyl group (e.g. a cyclohexylmethylcarbamoyl group); an aromatic heterocyclic carbamoyl group (e.g. an isoxazolylcarbamoyl group and benzothiazolylcarbamoyl group); a non-aromatic heterocyclic carbamoyl group (e.g. a pyrrolidinylcarbamoyl group); a C1-10 alkylsulfinyl group (e.g. a methylsulfinyl group and ethylsulfinyl group); a C1-10 alkylsulfonyl group (e.g. a methylsulfonyl group and ethylsulfonyl group); a C6-14 arylsulfonyl group (e.g. a phenylsulfonyl group); a (mono or di-C1-10 alkyl)phosphono group which may form a ring (e.g. a dimethylphosphono group; a diethylphosphono group; a diisopropylphosphono group; a dibutylphosphono group; a 2-oxide-1,3,2-dioxaphosphinanyl group; a mono or di-(C1-6 alkyl group which may be substituted with 1 to 3
halogen)-sulfamoyl group (e.g. a methylsulfamoyl group and ethylsulfamoyl group), etc.

Preferable examples of "acyloxy group" include an acyloxy group with 2 to 13 carbon atoms, for example, a C1-6 alkyl-carbonyloxy etc (e.g. an acetyloxy, propionyloxy, butyryloxy and isobutyryloxy).

Preferable examples of "heteroaryloxy group" include 5 to 7-membered single ring type heteroaryloxy group, for example, 2-pyridyloxy, 3-pyridyloxy, 2-imidazolyloxy, 2-pyrimidinyloxy, 1,2,4-triazol-5-yloxy etc.

A "carbocyclic group" is a saturated or unsaturated cyclic hydrocarbon group having 3 to 14 carbon atoms, and specifically means an aryl group, a cycloalkyl group, cycloalkenyl group described below or a condensed carbocyclic ring in which these rings are condensed.

Here, an "aryl group" is an aromatic hydrocarbon group having 6 to 14 carbon atoms, and specifically includes a phenyl group, a naphthyl group, a biphenyl group, an anthryl group, an indenyl group, a pentalenyl group, an azulenyloxy group, a fluorenyl group, a phenanthryl group, etc. Preferably, it is a phenyl group, a naphthyl group, and a biphenyl group. Particularly preferred is a phenyl group.

Here, a "cycloalkyl group" is a saturated cycloalkyl group having 3 to 8 carbon atoms, and specifically includes a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, and a cyclooctyl group.
A "cycloalkenyl group" is a cycloalkenyl group having 3 to 8 carbon atoms, and preferably contains at least one, preferably one or two double bonds. Specifically it includes a cyclopropenyl group, a cyclobutenyl group, a cyclopentenyl group, a cyclopentadienyl group, a cyclohexenyl group, cyclohexadienyl groups (2,4-cyclohexadien-1-yl group, 2,5-cyclohexadien-1-yl group, etc.), a cycloheptenyl group, a cyclooctenyl group, etc.

A "condensed carbocyclic group" in which the "aryl group," "cycloalkyl group," and "cycloalkenyl group" are condensed specifically includes an indenyl group, an indanyli group, a 1,4-dihydronaphthyl group, a 1,2,3,4-tetrahydronaphthyl group (a 1,2,3,4-tetrahydro-2-naphthyl group, a 5,6,7,8-tetrahydro-2-naphthyl group, etc.), a perhydronaphthyl group, etc. As a P2 ring, preferred is an aryl group and a cycloalkyl group, and a phenyl group, a biphenyl group and a cyclohexyl group are more preferably.

An "aralkyl group" is an arylalkyl group in which the aryl part thereof is an aryl group as mentioned above, particularly a phenyl group and the alkyl part thereof is a "C1-6 alkyl group" of the above-mentioned definition, and specifically it includes a benzyl group, a phenethyl group, a 3-phenyl propyl group, a 4-phenylbutyl group, a 6-phenyl hexyl group, etc.

A "aralkoxy group" is an arylalkoxy group in which the aryl part thereof is an aryl group as mentioned above, particularly a phenyl group and the alkoxy part thereof is
a "C1-6 alkoxy group" of the above-mentioned definition, and, specifically it includes a benzyloxy group, a 3-phenylpropyloxy group, a 4-phenylbutyloxy group, a 6-phenylhexyloxy group, etc.

A "cycloalkylalkoxy group" is a cycloalkylalkoxy group in which the cycloalkyl part thereof is a "cycloalkyl group" of the above-mentioned definition and the alkoxy part thereof is a "C1-6 alkoxy group" of the above-mentioned definition, and, specifically it includes a cyclopropylmethoxy group, a cyclobutylmethoxy group, a cyclopentylmethoxy group, a cyclohexylmethoxy group, etc.

A "heterocyclic group" means a saturated or unsaturated (including partial unsaturation and complete unsaturation) 5-membered or 6-membered monocyclic heterocyclic ring containing at least one, preferably 1 to 4 hetero atoms selected from a nitrogen atom, an oxygen atom and a sulfur atom besides the carbon atoms, or a condensed ring of these heterocyclic rings or a condensed ring of these heterocyclic rings and a carbocyclic ring selected from benzene, cyclopentane and cyclohexane.

A "saturated monocyclic heterocyclic group" includes a pyrrolidinyl group, a tetrahydrofuryl group, a tetrahydrothienyl group, an imidazolidinyl group, a pyrazolidinyl group, a 1,3-dioxolanyl group, a 1,3-oxathiolanyl group, an oxazolidinyl group, a thiazolidinyl group, a piperidinyl group, a piperidino group, a piperazinyl group, a piperazino group, a tetrahydropyranyl
group, a tetrahydrothiopyranyl group, a dioxanyl group, a morpholinyl group, a thiomorpholinyl group, a 2-oxopyrrolidinyl group, a 2-oxopiperidinyl group, a 4-oxopiperidinyl group, a 2,6-dioxo piperidinyl group etc.

A "unsaturated monocyclic heterocyclic group" includes a pyrrolyl group, a furyl group, a thieryl group, an imidazolyl group, a 1,2-dihydro-2-oxoimidazolyl group, a pyrazolyl group, a diazolyl group, an oxazolyl group, an isoxazolyl group, a thiazolyl group, an isothiazolyl group, a 1,2,4-triazolyl group, a 1,2,3-triazolyl group, a tetrazolyl group, a 1,3,4-oxadiazolyl group, a 1,2,4-oxadiazolyl group, a 1,3,4-thiadia zolyl group, a 1,2,4-thiadiazolyl group, a furazanyl group, a pyridyl group, a pyrimidinyl group, a 3,4-dihydro-4-oxopyrimidinyl group, a pyridazinyl group, a pyrazinyl group, a 1,3,5-triazinyl group, an imidazolinyl group, a pyrazolinyl group and an oxazolinyl group (a 2-oxazolinyl group, a 3-oxazolinyl group and a 4-oxazolinyl group), an isooxazolinyl group, a thiazolinyl group, an isothiazolinyl group, a pyranyl group, a 2-oxopyrananyl group, a 2-oxo-2,5-dihydrofuranyl group, a 1,1-dio xo-1H-isothiazolyl group.

A "condensed heterocyclic ring" includes an indolyl group (for example, a 4-indolyl group, a 7-indolyl group, etc.), an isoindolyl group, a 1,3-dihydro-1,3-dioxo isoindolyl group, a benzofuranyl group (for example, a 4-benzofuranyl group, a 7-benzofuranyl group, etc.), an indazolyl group, an isobenzofuranyl group, a benzothiophenyl group (for example,
a 4-benzothiophenyl group, a 7-benzothiophenyl group, etc.) a benzooxazolyl group (for example, a 4-benzooxazolyl group, a 7-benzooxazolyl group, etc.), a benzimidazolyl group (for example, a 4-benzimidazolyl group, a 7-benzimidazolyl group, etc.), a benzothiazolyl group (for example, a 4-benzothiazolyl group, a 7-benzothiazolyl group, etc.), an indolidinyl group, a quinolyl group, an iso quinolyl group, a 1,2-dihydro-2-oxoquinolyl group, a quinazolinyl group, a quinoxalinyl group, a cinnolinyl group, a phthalazinyl group, a quinolidinyl group, a puryl group, a pteridinyl group, an indolinyl group, an isoindolinyl group, a 5,6,7,8-tetrahydroquinolyl group, a 1,2,3,4-tetrahydroquinolyl group, a 2-oxo-1,2,3,4-tetrahydroquinolyl group, a benzo[1,3]dioxolyl group, a 2,3-dihydrobenzo[1,4]dioxosilyl group, a 3,4-methylene dioxydipyridyl group, a 4,5-ethylene dioxydipyrimidinyl group, a chromenyl group, a chromanyl group, an isochromanyl group, etc.

As for the group not specifically defined here, usual definition is to be followed.

In the general formula [1], preferable examples and particularly preferable examples of each symbol are as follows. However, the present invention is not limited thereto.

[Preferable Z]

Z is preferably -O-, -NR²-, -S- or -SO-, and R² is preferably a hydrogen atom or a C1-4 alkyl group. Preferable Z is -O-, -S- or -SO-.
[Preferable 1 and m]

Preferably 1 is 0 or 1 and preferably m is 0 or 1.
Preferably 1+m is 1 or 2 and it is more preferably 1.

[Preferable R¹]

R¹ is preferably a C1-6 alkyl group which may be preferably substituted with a hydroxyl group, or a hydrogen atom. Particularly preferable R¹ is a hydrogen atom.

[Preferable R²]

R² is preferably a hydrogen atom, a hydroxyl group, a C1-6 alkyl group (preferably a C1-4 alkyl group), a carboxyl group, a C1-6 alkoxy carbonyl group (preferably a C1-4 alkoxy carbonyl group), a C1-6 alkyl group substituted with a hydroxyl group (preferably a hydroxy substituted C1-4 alkyl group), a carbamoyl group (preferably a carbamoyl group, a mono C1-4 alkyl substituted carbamoyl group or a di-C1-4 alkyl substituted carbamoyl group), a C1-6 alkyl group substituted with an acylamino group (preferably an acylamino substituted C1-4 alkyl group) or R¹ and R² combined together form a carbonyl group.

R² is particularly preferably a hydrogen atom or a C1-4 alkyl group substituted with a hydroxy group.

[Preferable R³]

R³ is a hydrogen atom or a C1-6 alkyl group and preferably a hydrogen atom or a methyl group.

[Preferable R⁴]

R⁴ is preferably a hydrogen atom or a halogen atom.

[Preferable V]
V is preferably a single bond or -(CH\textsubscript{n})\textsuperscript{−}, wherein n is 1 or 2. It is particularly preferably a single bond.

[Preferable P1 and P2 rings]

P1 and P2 rings each is a heterocyclic group (preferably a 5-membered or 6-membered heterocyclic ring containing 1 to 4, more preferably 1 to 3 hetero atoms preferably selected from a nitrogen atom, an oxygen atom and a sulfur atom, or a condensed ring of these heterocyclic rings which may be the same or different, or a condensed ring of these heterocyclic rings and a carbocyclic ring selected from benzene, cyclopentane and cyclohexane (particularly preferably a pyridyl group, a thiazolyl group, a pyrazinyl group, a pyridazinyl group, a pyrimidinyl group, more preferably a pyridyl group), or a carbocyclic group selected from C3-8 cycloalkyl groups (for example, a cyclohexyl group, a cyclopentyl group, etc.), a C3-8 cycloalkenyl group (for example, a cyclohexenyl group), and an aryl group (for example, a phenyl group, etc.).

[Preferable P1 ring]

The P1 ring is preferably a 5-membered or 6-membered monocyclic aromatic heterocyclic group or aromatic carbocyclic group.

[Preferable aromatic heterocyclic group for P1 ring]

The preferable aromatic heterocyclic group for the P1 ring is a 5-membered or 6-membered monocyclic aromatic heterocyclic group having at least one nitrogen atom and preferably an aromatic heterocyclic group as follows.
It is more preferably an aromatic heterocyclic group as follows.

[Preferable aromatic carbocyclic group for P1 ring]

The preferable aromatic carbocyclic group for the P1 ring is a phenyl group, a biphenyl group or a naphthyl group and particularly preferably a phenyl group.

[Particularly preferable P1 group]

A pyridyl group or a phenyl group is particularly preferably as a P1 ring.

[Preferable P2 ring]

A ring preferable as P2 is an aromatic carbocyclic group, a 5-membered or 6-membered cycloalkyl group, a 5-membered or 6-membered saturated heterocyclic group or a 5-membered or
6-membered aromatic heterocyclic group and more details are as follows.

[Preferable aromatic carbocyclic group in P2 ring]

The preferable aromatic carbocyclic group in the P2 ring is a phenyl group, a biphenyl group or a naphthyl group and particularly preferably a phenyl group. These aromatic carbocyclic groups may be substituted with a methylenedioxy group or an ethylenedioxy group as shown below.

![Chemical structures](image1)

[Preferable cycloalkyl group for P2 ring]

Preferable 5-membered or 6-membered cycloalkyl group for the P2 ring is specifically a cyclopentyl group or a cyclohexyl group, and may be substituted with an oxo group as shown below, and may form an ethylenedioxy group and a spiro ring.

![Chemical structures](image2)

[Preferable saturated heterocyclic group for P2 ring]

The saturated 5-membered or 6-membered heterocyclic group which may form a preferable condensed ring in the P2 ring may be substituted with an oxo group as follows.
It is preferably a piperidyl group, a piperidino group and a tetrahydroquinolyl group as follows.

[Preferable aromatic heterocyclic group for P2 ring]

The 5-membered or 6-membered aromatic heterocyclic group which may form a preferable condensed ring for the P2 ring is 5-membered or 6-membered monocyclic aromatic heterocyclic group which preferably has at least one nitrogen atom, and is specifically an aromatic heterocyclic group as follows.

[Particularly preferable P2 group]

Particularly preferred as P2 ring is a ring group as follows.
[Preferable substituent group of P1 ring and P2 ring]

The heterocyclic group and carbocyclic group of these rings P1 and P2 may be substituted with 1 to 3 substituents selected from the following group, respectively:

- a halogen atom,
- a hydroxyl group,
- a C1-6 alkoxy group (wherein the C1-6 alkoxy group may be substituted with a carboxyl group, a hydroxyl group, and -CONR₆R₇ (wherein R₆ and R₇ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, and the alkyl group may be substituted with a hydroxyl group or an acyloxy group)),
- a C1-6 alkyl group (wherein the C1-6 alkyl group may be substituted with 1 to 5, preferably 1 to 3 substituents selected from a halogen atom, a hydroxyl group, a C1-6 alkoxy group, -NR₆R₇ (wherein R₆ and R₇ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, and the alkyl group may be substituted with a hydroxyl group or an acyloxy group), -NR₆COR₇ (wherein R₆ and R₇ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, and the alkyl group may be substituted with a hydroxyl group or an acyloxy group)).
-CONR²³R²⁴ (wherein R²³ and R²⁴ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from or the above-mentioned Group A),

-NR¹²³R¹²⁴ (wherein R¹²³ and R¹²⁴ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),

-NR²²³COR²²⁴ (wherein R²²³ and R²²⁴ are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),

-COR²⁷ (wherein R²⁷ is a C1-6 alkyl group, C1-6 alkoxy group, a C3-8 cycloalkyl group, an aralkyl group, an aralkoxy group, a cycloalkylalkoxy group or a hydroxyl group),

-O-(CR¹²¹R¹²²)ₙ-R¹²⁸ (wherein R¹²¹ and R¹²² are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, n is 1 or 2, and R¹²⁸ is an acyl group, a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (i) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (l) of the Group C, wherein R¹²⁸ is preferably a 5-membered or 6-membered heterocyclic group which preferably contains 1 to 3 nitrogen atoms and/or sulfur atoms, more preferably an unsaturated monocyclic heterocyclic group such as a pyridyl group, or an aryl group such as a phenyl group), and

-a cyano group.
The P2 ring is preferably a carbocyclic group (more preferably an aryl group such as a phenyl group or a C3-8 cycloalkyl group, particularly preferably a phenyl group or a C5-6 cycloalkyl group), or a heterocyclic group (wherein this heterocyclic group is a saturated or unsaturated 5-membered or 6-membered heterocyclic group which contains 1 to 3 hetero atoms, preferably nitrogen atoms and/or sulfur atoms, particularly preferably 1 to 3 nitrogen atoms, a condensed ring group of these heterocyclic rings or a condensed heterocyclic group of these heterocyclic rings which may be the same or different and a carbocyclic ring selected from benzene, cyclopentane and cyclohexane. Specific examples of the preferable heterocyclic group are a thiazolyl group, a pyridyl group, a piperidyl group, a piperidino group, a quinolyl group or a 1,2,3,4-tetrahydroquinolyl group, a benzothiazolyl group, a benzimidazolyl group, a pyrrolidino group, etc.), and the carbocyclic group and heterocyclic group may be substituted with a substituent group selected from the following:
- a halogen atom,
- a hydroxyl group,
- a C1-6 alkoxy group which may be substituted with a halogen atom, -CONR\(^{23}R^{24}\) (wherein R\(^{23}\) and R\(^{24}\) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group. The alkyl group may be further substituted with a hydroxyl group or an acyloxy group), a C3-8 cycloalkyl group, a C1-6 alkoxy group, a carboxyl group or a C1-6 alkoxy carbonyl group,
- a C1-6 alkyl group which may be substituted with a halogen atom, a hydroxyl group or a C1-6 alkoxy group,
- \( \cdot NR^{123}R^{124} \) (wherein \( R^{123} \) and \( R^{124} \) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group),
- \( \cdot NR^{223}COR^{224} \) (wherein \( R^{223} \) and \( R^{224} \) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
- \( \cdot COR^{27} \) (wherein \( R^{27} \) is preferably a C1-6 alkyl group, C1-6 alkoxy group, a C3-8 cycloalkyl group, an aralkyl group, an aralkoxy group, a cycloalkylalkoxy group or a hydroxyl group),
- a saturated or unsaturated heterocyclic group having 1 to 3 nitrogen atoms as hetero atom (preferably a piperidinyl group, a piperidino group, a 2-oxopiperidino group, a morpholino group or a piperazino group, a 1,3-dioxolanyl group (spiro), a pyrrolidino group, etc., and these heterocyclic rings group may be substituted with a substituent group selected from a hydroxyl group, \( \cdot CONR^{23}R^{24} \) (wherein \( R^{23} \) and \( R^{24} \) are the same as above), a C1-6 alkoxy group, a carboxyl group, a C1-6 alkyl group which may be substituted with a C1-6 alkoxy group, a C1-6 alkoxy carbonyl group and an acyloxy group),
- \( \cdot O-R^{28} \) (wherein \( R^{28} \) is an acyl group, a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (l) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (l) of the Group C),
- O- (CR^{121}R^{122})_n-R^{128} (wherein R^{121} and R^{122} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, n is 1 or 2, and R^{128} is an acyl group, a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (i) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (l) of the Group C),
- a nitro group, and
- a cyano group.

[Substituent group preferable as a substituent group in P1 ring]

More specifically, the following is preferable as a substituent group in P1 ring:
- a halogen atom, particularly preferably a fluorine atom, a chlorine atom,
- a C1-6 alkyl group (preferably a C1-4 alkyl group),
- a cyano group,
- a hydroxyl group,
- a carboxyl group,
- a C1-6 alkoxy carbonyl group (preferably a C1-4 alkoxy carbonyl group),
- NR^{123}R^{124} (wherein R^{123} and R^{124} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
- a C1-6 alkoxy group (preferably a C1-4 alkoxy group (wherein the alkoxy group may be substituted with 1 to 2 substituents
selected from a hydroxyl group, a carboxyl group, \( -\text{CONR}^{23}R^{24} \) (wherein \( R^{23} \) and \( R^{24} \) are the same as above) and a 5-membered or 6-membered monocyclic aromatic heterocyclic group (preferably a pyridyl group),
- an aralkoxy group (preferably a benzyloxy group),
- an aralkoxycarbonyl group (preferably a benzylooxycarbonyl group),
- a C1-6 alkyl group (preferably a C1-4 alkyl group) substituted with 1 to 2 substituents selected from a hydroxyl group, a C1-4 alkoxy group and \( -\text{NR}^{123}R^{124} \) (wherein \( R^{123} \) and \( R^{124} \) are the same as above), or
- \( -\text{CONR}^{23}R^{24} \) (wherein \( R^{23} \) and \( R^{24} \) are the same as above).

Preferable substituent group for the aromatic carbocyclic group which constitutes P2 ring is a substituent group as follows:
- a halogen atom,
- a C1-6 alkyl group (preferably a C1-4 alkyl group) which may be branched,
- a C1-6 alkoxy group (preferably a C1-4 alkoxy group) (wherein the C1-6 alkoxy group may be substituted with a hydroxyl group, a carbamoyl group or a C1-6 alkoxy group),
- a nitro group,
- a cyano group,
- a carboxyl group,
- a halo C1-6 alkoxy group (preferably a halo C1-4 alkoxy group),
- a halo C1-6 alkylthio group (preferably a halo C1-4 alkylthio group),
- a C1-6 alkoxy carbonyl group (preferably a C1-4 alkoxy carbonyl group),
- an acyl group,
- a C1-6 alkyl group substituted with 1 to 3 substituents selected a hydroxyl group, a halogen atom and a C1-6 alkoxy group (preferably a C1-4 alkoxy group),
- \(\text{NR}^{123}\text{R}^{124}\) (wherein \(\text{R}^{123}\) and \(\text{R}^{124}\) are the same as above),
- \(\text{CONR}^{23}\text{R}^{24}\) (wherein \(\text{R}^{23}\) and \(\text{R}^{24}\) are the same as above),
- a saturated 5-membered or 6-membered heterocyclic ring containing at least one nitrogen atom (preferably a piperidino group, a pyrrolidino group or a morpholino group which may be substituted with an oxo group) which may be substituted with a substituent group selected from a hydroxyl group, a C1-6 alkoxy group, a carboxyl group, a C1-6 alkoxy carbonyl group, a C1-6 alkyl group (wherein the C1-6 alkyl group may be substituted with a hydroxyl group or a C1-6 alkoxy group), a carbamoyl group, and an acyloxy group, or
- a piperidinyl group which may be substituted with a C1-6 alkoxy carbonyl group.

[Substituent group particularly preferable as a substituent group in P1 ring]

Particularly preferable substituent group as a substituent group in P1 ring is a halogen atom (particularly a fluorine atom and a chlorine atom), a C1-4 alkyl group, a C1-4 alkyl group substituted with a hydroxyl group, a C1-4 alkyl group substituted with a C1-4 alkyl group, a C1-4 alkoxy group, an amino group, an amino group substituted with a C1-4
alkyl group or an amino group substituted with two identical or different C1-4 alkyl groups.

[Substituent group for an aromatic carbocyclic group in P2 ring]
- a halogen atom,
- a C1-6 alkyl group (preferably a C1-4 alkyl group) which may be branched,
- a C1-6 alkoxy group (wherein the C1-6 alkoxy group may be substituted with a hydroxyl group, a carbamoyl group or a C1-6 alkoxy group),
- a nitro group,
- a cyano group,
- a carboxyl group,
- a halo C1-6 alkoxy group (preferably a halo C1-4 alkoxy group),
- a halo C1-6 alkylthio group (preferably a halo C1-4 alkylthio group),
- a C1-6 alkoxy carbonyl group (preferably a C1-4 alkoxy carbonyl group),
- an acyl group,
- a C1-6 alkyl group substituted with 1 to 3 substituents selected from a hydroxyl group, a halogen atom and a C1-6 alkoxy group (preferably a C1-4 alkoxy group),
- NR_{1}^{23} R_{124}^{24} (wherein R_{1}^{23} and R_{124}^{24} are the same as above),
- CONR_{2}^{23} R_{124}^{24} (wherein R_{2}^{23} and R_{124}^{24} are the same as above),
- a saturated 5-membered or 6-membered heterocyclic ring containing at least one nitrogen atom (preferably a piperidino group, a pyrroolidino group or a morpholino group which may
be substituted with an oxo group) which may be substituted with a substituent group selected from a hydroxyl group, a C1-6 alkoxy group, a carboxyl group, a C1-6 alkoxy carbonyl group, a C1-6 alkyl group (wherein the C1-6 alkyl group may be substituted with a hydroxyl group or a C1-6 alkoxy group), a carbamoyl group, and an acyloxy group, or a piperidyl oxy group which may be substituted with a C1-6 alkoxy carbonyl group.

[Particularly preferable substituent group for an aromatic carbocyclic group in P2 ring]

Particularly preferable substituents for an aromatic carbocyclic group in P2 ring are the following substituents:
- a halogen atom,
- a C1-6 alkyl group (preferably C1-4 alkyl group) which may be branched,
- a C1-6 alkoxy group (wherein the C1-6 alkoxy group may be substituted with a hydroxyl group, a carbamoyl group or a C1-6 alkoxy group),
- a halo C1-6 alkyl group (preferably a halo C1-4 alkyl group, particularly preferably a trifluoromethyl group),
- a halo C1-6 alkoxy group (preferably a halo C1-4 alkoxy group),
- \( \text{NR}^{123}\text{R}^{124} \) (wherein \( \text{R}^{123} \) and \( \text{R}^{124} \) are the same as above, and particularly, it may be a di-C1-4 alkylamino group),
- a saturated 5-membered or 6-membered heterocyclic ring containing at least one nitrogen atom (preferably a piperidino group, a pyrrolidino group or a morpholino group which may be substituted with an oxo group), which may be substituted
with a substituent group selected from a C1-6 alkoxy group, a C1-6 alkoxy carbonyl group, and a C1-6 alkyl group (wherein the C1-6 alkyl group may be substituted with a C1-6 alkoxy group), or
- a piperidyl oxy group which may be substituted with a C1-6 alkoxy carbonyl group.

[Substituent group preferable for a saturated hydrocarbocyclic group in P2 ring]
- a C1-6 alkyl group (wherein the C1-6 alkyl group may be substituted with a hydroxyl group or a C1-6 alkoxy group),
- a C1-6 alkoxy group (wherein the C1-6 alkoxy group may be substituted with a carboxyl group, a C1-6 alkoxy carbonyl group, a carboxamoyl group),
- an aralkoxy group,
- \( \cdot NR^{123}R^{124} \) (wherein \( R^{123} \) and \( R^{124} \) are the same as above),
- an oxo group,
- a C3-6 cycloalkyl C1-6 alkoxy group,
- a C3-6 cycloalkyloxy group or
- a saturated 5-membered or 6-membered heterocyclic ring containing at least one nitrogen atom (preferably a piperidino group, a pyrrolidino group, a morpholino group which may be substituted with an oxo group),

[Substituents particularly preferable for a saturated hydrocarbocyclic group in P2 ring]
- a C1-6 alkyl group (particularly a C1-4 alkyl group),
- a C1-6 alkoxy group (particularly a C1-4 alkoxy group), or
[Substituents preferable for a saturated heterocyclic group in P2 ring]
- a C1-6 alkyl group (wherein the C1-6 alkyl group may be substituted with a hydroxyl group or a C1-6 alkoxy group),
- a C1-6 alkoxy group,
- a carbamoyl group or
- an acyl group.

[Substituents preferable for an aromatic heterocyclic group in P2 ring]
- a halogen atom,
- a C1-6 alkyl group or
- a halo C1-6 alkyl group.

A "pharmacuetically acceptable salt" may be any kind of salt as long as it forms a nontoxic salt with a compound represented by the above-mentioned general formula [1], and can be obtained by reacting it with, for example, an inorganic acid such as hydrochloric acid, sulfuric acid, phosphoric acid, or hydrobromic acid; an organic acid such as oxalic acid, malonic acid, citric acid, fumaric acid, lactic acid, malic acid, succinic acid, tartaric acid, acetic acid, trifluoroacetic acid, gluconic acid, ascorbic acid, methylsulfonic acid, or benzenesulfonic acid; an inorganic base such as sodium hydroxide, potassium hydroxide, calcium hydroxide, magnesium hydroxide, or ammonium hydroxide; an organic base such as methylamine, diethylamine, triethylamine, triethanolamine, ethylenediamine, tris(hydroxymethyl)methylamine, guanidine, choline, or
cinchonine; or an amino acid such as lysine, arginine or alanine. A hydrated compound, hydrate and solvate of each compound are also included in the present invention.

In addition, various isomers exist for the compound represented by the above-mentioned general formula [1]. For example, E isomer and Z isomer exist as geometric isomers, and when an asymmetric carbon atom exists, enantiomers and diastereomers exist as stereoisomers based on these, and tautomers may exist. Therefore, all of these isomers and the mixtures thereof are included in the range of the present invention. In addition, the present invention also encompasses prodrug compounds of these compounds and metabolite compounds as equivalent compounds besides the compound represented by the above-mentioned general formula [1].

A "prodrug" is a derivative of the compound of the present invention having a group which may be decomposed chemically or metabolically and after administered to a living body, it goes through a chemical change to a compound which has an activity as a drug and exhibits original pharmacological effect, and complexes and salts not by a covalent bond are included.

A prodrug is used for improving absorption upon oral administration or targeting to a target site. Moieties to be modified for forming a prodrug include reactive functional groups such as a hydroxyl group, a carboxyl group, an amino group, and a thiol group in the compound of the present invention. Specific examples of the modifying group for a hydroxyl group
include an acetyl group, a propionyl group, an isobutyryl group, a pivaloyl group, a benzoyl group, a 4-methylbenzoyl group, a dimethylcarbamoyl group, a sulfo group, etc. Specific examples of the modifying group for a carboxyl group include an ethyl group, a pivaloyloxymethyl group, a 1-(acetoxy)ethyl group, a 1-(ethoxyacetyl)ethyl group, a 1-(cyclohexyloxyacetyl)ethyl group, a carboxymethyl group, a methyl (5-methyl-2-oxo-1,3-dioxol-4-yl) group, a phenyl group, an o-tolyl group, etc. Specific examples of the modifying group for an amino group include a hexylcarbamoyl group, a 3-methylthio-1-(acetylamino)propylcarbonyl group, a 1-sulfo-1-(3-ethoxy-4-hydroxyphenyl)methyl group, a methyl(5-methyl-2-oxo-1,3-dioxol-4-yl) group, etc.

A "pharmaceutical composition" encompasses a combination drug with another drugs, etc., besides the so-called "composition" which comprises an active ingredient as a drug and a combinational agent, etc. Needless to say, the pharmaceutical composition of the present invention may be used in combination with any kind of other drugs as long as it is permitted in the medical scene. Therefore, it can also be said that this pharmaceutical composition is a pharmaceutical composition for the combined use with other drugs.

A "pain" means every type of pain condition no matter what the condition is (for example, no matter whether it is a dull pain or a sharp pain, chronic or acute, etc.), no matter which disease causes the pain (for example, no matter whether
the pain is resulted from rheumatism, or the pain resulted from cancer, etc.). Therefore, the "pain" as used herein encompasses, in addition to the so-called "pain," acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy, and neurodegenerative disease.

An "inhibitor of vanilloid receptor subtype 1 (VR1) activity" means a substance which inhibits the function of the vanilloid receptor subtype 1 as an ion channel, and eliminates or attenuates the activity. Specifically, it includes vanilloid receptor subtype 1 antagonist, etc. The vanilloid receptor subtype 1 antagonist means a substance which inhibits the effect of the agonist which acts on the vanilloid receptor subtype 1, thereby inhibiting the function of the vanilloid receptor subtype 1 as an ion channel. The inhibitor of the present invention has not to compete with the agonist but may also inhibit the function as a VR1 ion channel. Specifically, agonists which act on the vanilloid receptor subtype 1 include capsaicin, capsaicin derivatives, acid stimulation (proton), heat stimulation, etc., the inhibitor of vanilloid receptor subtype 1 (VR1) activity may be a substance which inhibits the Ca²⁺ inflow into the cell caused
by agonist stimulation of capsaicin, acid stimulation (proton) or heat stimulation.

The pharmaceutical composition of the present invention can be administered to human as well as other mammals (mouse, rat, hamster, rabbit, cat, dog, cow, horse, sheep, monkey, etc.). Therefore, the pharmaceutical composition of the present invention is useful also as a drug for animal not to mention for human.

When the compound of the present invention is used as a pharmaceutical preparation, it can be mixed with a pharmacologically acceptable carrier usually known in itself, excipient, diluent, extender, disintegrating agent, stabilizer, preservative, buffer, emulsifier, flavor, colorant, sweetener, thickener, corrigent, dissolution auxiliary agent, and other additive agents, specifically water, plant oil, alcohol such as ethanol or benzyl alcohol, carbohydrates such as polyethylene glycol, glycerol triacetate, gelatin, lactose and starch, magnesium stearate, talc, lanolin, vaseline, etc. to prepare a drug in the form such as tablet, pill, powder, granule, suppository, injection agent, eye-drops, liquid medicine, capsule agent, troche, aerosol agent, elixir agent, suspension, emulsion and syrup for systemic or local administration by oral or parenteral route.

Although the dosage varies depending on age, weight, condition, therapeutical effect, administration methods, etc., it is usually administered at a dose in the range of
0.01 mg to 1 g per dose, 1 time to several times per day, to adults, in the form of an oral preparate or injection preparation such as an intravenous injection, etc.

"Preventing" is the so-called prevention, and means, for example, suppressing the onset of neuralgia or chronicity of neuralgia prophylactically. As for pain, specifically included is prophylactically suppressing the onset of acute postherpetic neuralgia, onset of postherpetic neuralgia, transition to postherpetic neuralgia from acute herpetic pain, chronicity of postherpetic neuralgia, onset of postoperative pain, chronicity of postoperative pain, onset of symptoms of cancer pain, chronicity of cancer pain, onset of symptoms of inflammatory pain, onset of interstitial cystitis, chronicity of inflammatory pain, onset of posttraumatic neuralgia or chronicity of posttraumatic neuralgia.

A "drug comprising a combination" means a drug characterized in that it is a formulation containing a pharmaceutical composition or an agent to be combined, a drug characterized in that it is a kit comprising a pharmaceutical composition or an agent to be combined, a drug characterized in that a pharmaceutical composition or an agent to be combined is administered via the same or different administration routes, respectively.

The compound and pharmaceutical composition of the present invention can be used in combination with one or more other agents following a general method currently performed in the usual medical site. When used in combination, the drug
to be used with may be administered simultaneously or separately with a time lag. Although there are various compounds which can be used in combination with the compound of the present invention, particularly preferred are an anti-virus agent, an antidepressant, an anticonvulsant, an antiarrhythmic drug, a local anesthetic, an anesthetic drug, a N-methyl-D-aspartate receptor antagonist, adrenal cortical steroid, a nerve block, a nonsteroidal antiinflammatory analgesic, narcotics, an antagonist analgesic, an \( \beta_2 \)-adrenaline receptor agonist, a stimulation analgesic method, drugs for external application, a calcium channel antagonist, and a potassium channel opener.

The anti-virus agent specifically includes vidarabine, acyclovir, ganciclovir, zidovudine, didanosine, amantadine, and idoxuridine, interferon, etc.

The antidepressant specifically includes amitriptyline, imipramine, clomipramine, trimipramine, lofepramine, doxulepin, desipramine, amoxapine, nortriptyline, fluoxetine, fluvoxamine, maprotiline, mianserin, setiptiline, trazodone, etc.

The anticonvulsant specifically includes gabapentin, pregabalin, phenobarbital, primidone, phenytoin, mephenytoin, nirvanol, ethotoin, trimethadione, ethosuximide, acetylpheneturide, carbamazepine, zonisamide, acetazolamide, diazepam, clonazepam, nitrazepam, diphenylhydantoin, valproic acid, baclofen, etc.
The antiarrhythmic drug specifically includes quinidine, disopyramide, procainamide, ajmaline, prajmalium, cibenzoline, lidocaine, mexiletine, aprindine, tonicaid, phenytoin, flecainide, pilcicainide, propafenone, propranolol, amiodarone, verapamil, bepridil, etc.

The local anesthetic specifically includes lidocaine, mexiletine, cocaine, procaine, bupivacaine, mepivacaine, prilocaine, tetracaine, dibucaine, ethyl aminobenzoate, etc.

The anesthetic drug specifically includes benzodiapine, diazepam, midazolam, thiopental, thiamylal, propofol, baclofen, droperidol, sufentanil, etc. are mentioned. The N-methyl-D-aspartate receptor antagonist specifically includes ketamine, dextromethorphan, memantine, amantadine, etc. are included.

The adrenal cortical steroid specifically includes cortisol, cortisone, prednisolone, triamcinolone, dexamethasone, betamethasone, paramethasone, fluocinolone acetonide, fluocinonide, beclomethasone, fludrocortisone, etc.

The nerve block specifically includes stellate ganglion block, epidural ganglion block, brachial plexus ganglion block, nerve root block, thoracic/lumbar sympathetic ganglion, trigger point block, subarachnoid ganglion block, trigeminal nerve block, sympathetic nerve block, local infiltration block, peripheral nerve block, etc.

The nonsteroidal antiinflammatory analgesic specifically includes celecoxib, rofecoxib, etodolac,
meloxicam, nimesulid, sodium diclofenac, mefenamic acid, zaltoprofen, sodium loxoprofen, sulindac, nabumetone, diflunisal, piroxicam, ibuprofen, naproxen, fenoprofen, acetylsalicylic acid, tolmetin, indomethacin, flurbiprofen, oxaprozin, ketoprofen, mofezolac, acetaminophen, ketorolac, zomepirac, nitroaspirin, tiaprofen, ampiroxicam, tiaramide, epirizole, etc.

The narcotics specifically include morphine, fentanyl, oxycodone, methadon, codeine, cocaine, pethidine, opium, ipecac, etc.

The antagonist analgesic specifically includes pentagyn, buprenorphine, nalorphine, cyclazocine, butorphanol, etc.

The $\alpha_2$-adrenaline receptor agonist specifically includes clonidine, dexmedetomidine, tizanidine, guanfacine, guanabenz, etc.

The medicine for external application specifically includes capsaicin cream etc.

The stimulation analgesic method specifically includes acupuncture, a percutaneous electricity needle stimulation therapy, a percutaneous electricity nerve stimulation therapy, a silver spike point (SSP) treatment, a peripheral nerve stimulus, a spine electricity stimulus, an electric spasm treatment, laser surgery, a low-frequency therapy, etc.

In addition, the compound of the present invention can be used following the general method usually performed in the art by administration after performing a surgical operation to prevent or treat pain. Although various surgical
operations can be performed in combination with the compound of the present invention, cicatrectomy, nerve freezing, peripheral nerve excision, spinal dorsal root excision, sympathectomy, spinal cord dorsal root entry zone destruction, cordotomy, and frontal lobe excision are particularly preferable.

Although application of the compound of the present invention has been described mainly as a use for preventing or treating pain, the compound of the present invention can be applied to the conditions in which C fibers participates, for example, pruritus, allergic and allergic rhinitis, overactive bladder type frequent urination and urinary incontinence, apoplexy, irritable bowel syndrome, respiratory ailment such as asthma and a chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer, inflammatory bowel disease, etc.

Next, a preparation method of the compound represented by the general formula [1] of the present invention is described specifically but, needless to say, the present invention is not limited to these preparation methods. Therefore, the compound of the present invention may be synthesized according to the following manufacture methods A, B or C, but it can be prepared according to the below-mentioned examples, or referring to these processes. In preparation of the compound of the present invention, the order of reaction operation can be changed suitably. It can be performed starting from the reaction step or substitution part considered to be rational.
For example, the P2 ring may be introduced before the P1 ring is introduced, and this order may be reversed. As for the formation of the hetero ring condensed to the benzene ring (of benzamide group), a closed ring reaction may be performed to form this hetero ring before introducing the P1 ring and/or P2 ring or alternatively, a closed ring reaction may be performed to form this hetero ring after introducing the P1 ring and/or P2 ring. Protection and deprotection may be suitably conducted when there is a reactant functional group. In order to enhance development of the reaction, reagents other than those illustrated can be used suitably.

The following production process flow is an example of the typical preparation method, but preparation of the compound of the present invention is not particularly limited to the following method. Each compound obtained at each step can be isolated and purified by a usual method, but depending on the case the compound can be used in the next step without being isolated and purified.

1. Preparation method A;
(wherein, R represents a carboxyl protecting group (the carboxyl protecting group here includes, for example, a methyl group, an ethyl group, a propyl group, a tert-butyl group, a benzyl group, a paramethoxy benzyl group, etc.), and forms an ester which is easily led to a carboxylic acid by hydrolysis or catalytic hydrogenation reaction. X represents a halogen atom such as chloro and bromo or a sulfonyloxy group such as a 3-nitrobenzene sulfonyloxy group, a p-toluenesulfonyloxy group, a benzene sulfonyloxy group, a p-bromobenzenesulfonyloxy group, a methanesulfonyloxy group or a trifluoromethanesulfonyloxy group, and each other symbol is the same as above.)

First Step

This is the reaction for obtaining a compound (IIIA) by the palladium catalyzed Buchwald/Hartwig type amination reaction from a compound (IA) and a compound (IIA).
The compound (IIIA) can be obtained by reacting the (IA) with the compound (IIA) in toluene, 1,4-dioxane, tetrahydrofuran or the like or a mixed solvent of these, using a palladium catalyst such as a mixture of palladium acetate and 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl, bis(diphenylphosphino)ferrocene palladium chloride (II) or tris(dibenzylideneaceto) dipalladium together with a base such as sodium carbonate, tripotassium phosphate (K₃PO₄), potassium carbonate, cesium carbonate, sodium bicarbonate, potassium bicarbonate or potassium tert butoxide, at a temperature of 20°C to reflux temperature, preferably 60°C to reflux temperature for 5 hours to 96 hours preferably for 8 hours to 48 hours.

When the P1 ring is an aromatic hetero ring such as oxazole, thiazole or 1,3,4-thiadiazole, the compound (IIA) is not used and the reaction builds up the aromatic hetero ring by a known method and proceeds to the Second Step.

For example, when the P1 ring is 5-methylloxazole, the compound (IA) is reacted with thiophosgene or 1,1-thiocarbonyldiimidazole etc., in tetrahydrofuran, ethyl acetate, toluene, water or a mixed solvent of these, in the presence or absence of a base such as triethylamine, sodium bicarbonate, potassium carbonate or pyridine to form an isothiocyanate compound which is then allowed to react with 1-azidoacetone and triphenylphosphine in a solvent such as dichloromethane, dichloroethane and chloroform at a temperature of -20°C to reflux temperature, preferably 0°C.
to 50°C for 0.5 hour to 24 hours, preferably 2 hours to 8 hours to obtain the compound (IIIA).

When the P1 ring is 4-methylthiazole, the compound (IA) is reacted with thiophosgene or 1,1-thiocarbonyldiimidazole, etc., in tetrahydrofuran, ethyl acetate, toluene, water or a mixed solvent of these, in the presence or absence of a base such as triethylamine, sodium bicarbonate, potassium carbonate or pyridine to form an isothiocyanate compound which is then allowed to react with ammonia water and the like to obtain a thiourea compound, and this compound is allowed to react with 1-chloroacetone, 1-bromoacetone, etc. in a solvent such as methanol, ethanol, tetrahydrofuran or acetone at a temperature of -20°C to reflux temperature, preferably 0°C to reflux temperature for 0.5 hour to 24 hours, preferably 1 hour to 12 hours to obtain the compound (IIIA).

Second Step

This is a step to remove R from the compound (IIIA) and obtain a carboxylic acid (IVA).

For example, when R is a methyl group, ethyl group, propyl group, etc., the compound (IIIA) can be hydrolyzed in water, methanol, ethanol, propanol, tetrahydrofuran, or a mixed solvent of these using a base such as sodium hydroxide, potassium hydroxide, lithium hydroxide, potassium carbonate or sodium carbonate, at a temperature of -20°C to reflux temperature, preferably 20°C to reflux temperature for 0.5 hour to 24 hours, preferably 0.5 hour to 8 hours to obtain the compound (IVA).
For example, when R is a tert-butyl group, the compound (IVA) can be obtained by the reaction of the compound (IIIA) without a solvent or in water, methanol, ethanol, propanol, tetrahydrofuran or a mixed solvent of these using an acid such as hydrochloric acid or trifluoroacetic acid at a temperature of 0°C to reflux temperature, preferably 0°C to 50°C for 0.5 hour to 24 hours, preferably 0.5 hour to 8 hours.

When R is a benzyl group, paramethoxybenzyl group, etc., the compound (IVA) can be obtained by the reaction in methanol, ethanol, propanol, tetrahydrofuran or a mixed solvent of these in the presence of palladium carbon catalyst, etc. using hydrogen or ammonium formate at a temperature of about 0°C to reflux temperature, preferably about 20°C to reflux temperature for 0.5 hour to 96 hours, preferably 1 hour to 48 hours.

Third Step

This is a reaction to obtain a compound (1) by condensation reaction of a compound (IVA) and a compound (VA).

The condensation reaction can be performed using a condensing agent, or via an acid chloride etc.

When a direct condensation reaction is performed using a condensing agent, a compound (IVA) is reacted with a compound (VA) in N,N-dimethylformamide, methylene chloride, chloroform etc. or a mixed solvent of these using a condensing agent such as dicyclohexylcarbodiimide or 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide at a temperature of -20°C to reflux temperature, preferably about
0°C to 50°C for 1 hour to 48 hours, preferably about 1 hour to 24 hours. In this case, it is preferable to add additives such as hydroxybenzotriazole or N-hydroxsuccinimide.

When the process goes via an acid chloride, the compound (IVA) is reacted with thionyl chloride, oxalyl chloride, etc. in chloroform, methylene chloride, tetrahydrofuran, etc. or a mixed solvent of these to obtain an acid chloride of (IVA) and this is reacted with a compound (VA) in toluene, chloroform, tetrahydrofuran or a mixed solvent of these in the presence of a base such as triethylamine or pyridine at a temperature of -20°C to reflux temperature, preferably about 0°C to 40°C for 0.5 hour to 24 hours, preferably about 0.5 hour to 12 hours to obtain the compound (I).

2. Preparation method B;

![Chemical diagram]
This is an alternative method of preparing a compound (IIIA) in which Z is an oxygen atom in the Preparation method A. (wherein, Z is an oxygen atom, R is a C1-6 alkyl group and forms an ester which is easily led to a carboxylic acid by hydrolysis or catalytic hydrogenation reaction. R' represents a protecting group of a phenolic hydroxyl group easily removable by hydrolysis or catalytic hydrogenation reaction. Each other symbol is the same as above.)

First Step

This is the reaction for obtaining a compound (IIB) by the palladium catalyzed Buchwald/Hartwig type amination reaction from a compound (IB) and a compound (IIA).

The compound (IIB) can be obtained by reacting the (IB) with the compound (IIA) in toluene, 1,4-dioxane, tetrahydrofuran or the like or a mixed solvent of these, using a palladium catalyst such as a mixture of palladium acetate and 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl, bis(diphenylphosphino)ferrocene palladium chloride (II) or tris(dibenzylideneacetone)dipalladium together with a base such as sodium carbonate, tripotassium phosphate (K₃PO₄), potassium carbonate, cesium carbonate, sodium bicarbonate, potassium bicarbonate or potassium tert butoxide, at a temperature of 20°C to reflux temperature, preferably 60°C to reflux temperature for 5 hours to 96 hours preferably for 8 hours to 48 hours.
When P1 ring is an aromatic hetero ring such as oxazole, thiazole or 1,3,4-thiadiazole, the compound (IIA) is not used and the reaction builds up the aromatic hetero ring by a known method and proceeds to the Second Step.

For example, when the P1 ring is 5-methylloxazole, the compound (IB) is reacted with thiophosgene or 1,1-thiocarbonyldiimidazole etc., in tetrahydrofuran, ethyl acetate, toluene, water or a mixed solvent of these, in the presence or absence of a base such as triethylamine, sodium bicarbonate, potassium carbonate or pyridine to form an isothiocyanate compound which is then allowed to react with 1-azidoacetone and triphenylphosphine in a solvent such as dichloromethane, dichloroethane or chloroform at a temperature of -20°C to reflux temperature, preferably 0°C to 50°C for 0.5 hour to 24 hours, preferably 2 hours to 8 hours to obtain the compound (IIB).

When the P1 ring is 4-methylthiazole, the compound (IB) is reacted with thiophosgene or 1,1-thiocarbonyldiimidazole etc., in tetrahydrofuran, ethyl acetate, toluene, water or a mixed solvent of these, in the presence or absence of a base such as triethylamine, sodium bicarbonate, potassium carbonate or pyridine to form an isothiocyanate compound which is then allowed to react with ammonia water or the like to obtain a thiourea compound, and this compound is allowed to react with 1-chloroacetone, 1-bromoacetone, etc. in a solvent such as methanol, ethanol, tetrahydrofuran or acetone at a temperature of -20°C to reflux temperature, preferably 0°C
to reflux temperature for 0.5 hour to 24 hours, preferably 1 hour to 12 hours to obtain the compound (IIB).

Second Step

This is a step to remove R' which is a protecting group of the phenolic hydroxyl group of the compound (IIB).

For example, when R' is a methoxymethyl group, a tetrahydropyranyl group, etc., the compound (IIIB) can be obtained by reacting the compound (IIB) without solvent or in water, methanol, ethanol, isopropanol, tetrahydrofuran, chloroform or a mixed solvent of these using an acid such as sulfuric acid, hydrochloric acid or trifluoroacetic acid at a temperature of 0°C to reflux temperature, preferably 0°C to 50°C for 0.5 hour to 24 hours, preferably 0.5 hour to 8 hours.

Third Step

This is a step to obtain a cyclic compound (IIIA) from the compound (IIIB).

For example, when l is 1, R1 and R2 are hydrogen atoms and m is 0, the compound (IIIB) is reacted with a dihalo reagent such as 1,2-dibromoethane, 1,2-dichloroethane or 1-bromo-2-chloroethane, in chloroform, tetrahydrofuran, N,N-dimethylformamide, dimethylsulfoxide, ethyl acetate, methanol, water or a mixed solvent of these in the presence of a base such as sodium carbonate, potassium carbonate, sodium hydroxide, potassium hydroxide or triethylamine at a temperature of 0°C to reflux temperature, preferably 0°C to 60°C for 0.5 hour to 24 hours to obtain a compound (IIIA).
Alternatively, after making it react with a haloacetic acid chloride such as acetyl chloride, and forming a cyclic lactam, the compound (IIIA) can also obtained by performing a reduction reaction in tetrahydrofuran using a borane-tetrahydrofuran complex reagent etc at 0°C to reflux temperature for 0.5 hour to 8 hours.

For example, when 1 is 1; R1 is a hydroxymethyl group; R2 is a hydrogen atom and m is 0, the similar reaction as a dihalo reagent can be performed using glycidyl chloride, glycidyl tosylate or glycidyl nosylate to obtain a compound (IIIA). In this case, the produced hydroxyl group may be subjected to a step to introduce a widely used protecting group such as an acetyl group, a methoxymethyl group, a tetrahydropyranyl group or a benzyl group, and used in the following steps, and can be suitably deprotected to the compound (1).

3. Preparation method C;

This is an alternative method of preparing a compound (1) in which Z is an oxygen atom in the Preparation method A.
(wherein, R is a C1-6 alkyl group and forms an ester which is easily led to a carboxylic acid by hydrolysis or catalytic hydrogenation reaction. R'' represents a protecting group of an amino group easily removable by hydrolysis or catalytic hydrogenation reaction. Each other symbol is the same as above.)

First step

This is a step to obtain a cyclic compound (II C) from a compound (I C).

For example, when 1 is 1, R1 and R2 are hydrogen atoms, and m is 0, the compounds (I C) is reacted with a dihalo reagent such as 1,2-dibromoethane, 1,2-dichloroethane or 1-bromo-2-chloroethane in chloroform, tetrahydrofuran, N,N-dimethylformamide, dimethylsulfoxide, ethyl acetate, methanol, water or a mixed solvent of these in the presence of a base such as sodium carbonate, potassium carbonate, sodium hydroxide, potassium hydroxide or triethylamine at a
temperature of 0°C to reflux temperature for 0.5 hour to 8 hours to obtain a compound (IIC).

For example, when \( l \) is 1, \( R_1 \) is a hydroxymethyl group, \( R_2 \) is a hydrogen atom, and \( m \) is 0, the compound (IIC) can be obtained using the compound (IC) along with glycidyl chloride, glycidyl tosylate or glycidyl nosylate in the similar reaction as with a dihalo reagent. In this case, the produced hydroxyl group may be subjected to a step to introduce a widely used protection group such as an acetyl group, a methoxymethyl group, a tetrahydropyranyl group or a benzyl group, and used in the following steps, and can be suitably deprotected to the compound (I).

Second step

This is a step to remove \( R \) of the compound (IIC) to obtain a carboxylic acid compound (IIIC).

For example, when \( R \) is a methyl group, ethyl group, propyl group, etc., the compound (IIIC) can be obtained by hydrolyzing the compound (IIC) in water, methanol, ethanol, propanol, tetrahydrofuran, etc. or a mixed solvent of these using a base such as sodium hydroxide, potassium hydroxide, lithium hydroxide, potassium carbonate and sodium carbonate or an aqueous solution of these at a temperature of -20°C to reflux temperature preferably 20°C to reflux temperature for 0.5 hour to 24 hours, preferably for 0.5 hour to 8 hours.

For example, when \( R \) is a tert-butyl group, the compound (IIIC) can be obtained by reacting the compound (IIC) without solvent or in water, methanol, ethanol, propanol, or
tetrahydrofuran, etc. or a mixed solvent of these using an acid such as hydrochloric acid or trifluoroacetic acid, at a temperature of 0°C to reflux temperature preferably 0°C to 50°C for 0.5 hour to 24 hours, preferably 0.5 hour to 8 hours.

When R is a benzyl or paramethoxybenzyl group, etc., the compound (IIIC) can be obtained by reacting in methanol, ethanol, propanol, tetrahydrofuran, etc. or a mixed solvent of these using hydrogen or ammonium formate in the presence of a palladium carbon catalyst, etc. at a temperature of about 0°C to reflux temperature, preferably about 20°C to 50°C for 0.5 hour to 96 hours, preferably 1 hour to 48 hours.

Third step

This is a reaction which obtains a compound (IVC) by the condensation reaction of a compound (IIIC) and a compound (VA). The condensation reaction can be performed either by a method using a condensing agent or by a method via an acid chloride, etc.

When a direct condensation reaction using a condensing agent is performed, the compound (IVC) can be obtained by reacting the compound (IIIC) with the compound (VA) in N,N-dimethylformamide, methylene chloride, chloroform, etc. or a mixed solvent of these using a condensing agent such as dicyclohexylcarbodiimide or 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide at a temperature of -20°C to reflux temperature, preferably about 0°C to 50°C for 1 hour to 48 hours, preferably 1 hour to 24
hours. In this case, it is preferable to add additives such as hydroxybenzotriazole and N-hydroxysuccinic acid imide.

When a reaction via an acid chloride is performed, the compound (IIIC) is reacted with thionyl chloride, oxalyl chloride etc. in chloroform, methylene chloride, tetrahydrofuran, etc. or a mixed solvent of these to obtain an acid chloride of (IIIC), and this compound is reacted with the compound (VA) in toluene, chloroform, or tetrahydrofuran, etc. or a mixed solvent of these in the presence of a base such as triethylamine or pyridine at a temperature of -20°C to reflux temperature, preferably about 0°C to 40°C for 0.5 hour to 24 hours, preferably 0.5 hour to 12 hours and thereby the compound (IVC) can be obtained.

Fourth step

This is a step to remove the amino protecting group R'' of compound (IVC) and obtain a compound (VC).

For example, when R'' is an acetyl group or a formyl group, the compound (VC) can be obtained by reacting the compound (IVC) in tetrahydrofuran, methanol, ethanol, isopropanol, water or a mixed solvent of these in the presence of a base such as sodium hydroxide, potassium hydroxide, lithium hydroxide or sodium methoxide or in the presence of an acid such as sulfuric acid, hydrochloric acid or trifluoroacetic acid at a temperature of -20°C to reflux temperature, preferably about 0°C to reflux temperature for 0.5 hour to 24 hours, preferably 0.5 hour to 8 hours.
For example, when R'' is a tert-butoxycarbonyl group, the compound (VC) can be obtained by reacting the compound (IVC) without solvent or in tetrahydrofuran, methanol, ethanol, isopropanol, chloroform, water or a mixed solvent of these in the presence of an acid such as hydrochloric acid or trifluoroacetic acid at a temperature of 0°C to reflux temperature, preferably about 0°C to 50°C for 0.5 hour to 24 hours, preferably 0.5 hour to 8 hours.

When R' is a benzyloxycarbonyl group, a benzyl group, etc., the compound (VC) can be obtained by reacting the compound (IVC) in methanol, ethanol, propanol, tetrahydrofuran, etc. or a mixed solvent of these using hydrogen or ammonium formate in the presence of a palladium carbon catalyst, etc. at a temperature of about 0°C to reflux temperature, preferably about 20°C to 50°C for 0.5 hour to 96 hours, preferably 1 hour to 48 hours.

Fifth step

This is a reaction for obtaining a compound (1) by the palladium catalyzed Buchwald/Hartwig type amination reaction from a compound (VC) and a compound (IIA).

The compound (1) can be obtained by reacting the compound (VC) with the compound (IIA) in toluene, 1,4-dioxane, tetrahydrofuran or the like or a mixed solvent of these, using a palladium catalyst such as a mixture of palladium acetate and 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl, bis(diphenylphosphino)ferrocene palladium chloride (II) or tris(dibenzylideneacetone)dipalladium together with a base
such as sodium carbonate, tripotassium phosphate (K₃PO₄), potassium carbonate, cesium carbonate, sodium bicarbonate, potassium bicarbonate, or potassium tert-butoxide, at a temperature of 20°C to reflux temperature, preferably 60°C to reflux temperature for 5 hours to 96 hours preferably for 8 hours to 48 hours.

When the P1 ring is an aromatic hetero ring such as oxazole, thiazole or 1,3,4-thiadiazole, the compound (IIA) is not used and the reaction builds up the aromatic hetero ring by a known method and proceeds to the Second Step.

When the P1 ring is 5-methylloxazole, the compound (VC) is reacted with thiophosgene or 1,1-thiocarbonyldiimidazole etc., in tetrahydrofuran, ethyl acetate, toluene, water, etc. or a mixed solvent of these in the presence or absence of a base such as triethylamine, sodium bicarbonate, potassium carbonate or pyridine to form an isothiocyanate compound which is then allowed to react with 1-azidoacetone and triphenylphosphine in a solvent such as dichloromethane, dichloroethane or chloroform at a temperature of -20°C to reflux temperature, preferably 0°C to 50°C for 0.5 hour to 24 hours, preferably 2 hours to 8 hours to obtain the compound (1).

When the P1 ring is 4-methylthiazole, the compound (VC) is reacted with thiophosgene or 1,1-thiocarbonyldiimidazole etc., in tetrahydrofuran, ethyl acetate, toluene, water, etc. or a mixed solvent of these in the presence or absence of a base such as triethylamine, sodium bicarbonate, potassium carbonate or pyridine to form an isothiocyanate compound which
is then allowed to react with ammonia water and the like to obtain a thiourea compound, and this compound is allowed to react with 1-chloroacetone, 1-bromoacetone, etc. in a solvent such as methanol, ethanol, tetrahydrofuran or acetone at a temperature of -20°C to reflux temperature, preferably 0°C to reflux temperature for 0.5 hour to 24 hours, preferably 1 hour to 12 hours to obtain the compound (1).

The following examples refer to compounds shown in Tables 1 through 50.

Example 1
Example 1 - 001:
Production of N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of methyl 3-nitrosalicylate:
3-Nitrosalicylic acid (7.5 g) was dissolved in methanol (75 ml), concentrated sulfuric acid (2 ml) was added, and the mixture was refluxed for 24 hours. Reaction mixture was left cool to and the precipitated pale yellowish solid was collected by filtration, washed with water and dried in vacuo to obtain the title compound (7.19 g).

Second Step
Production of methyl 3-aminosalicylate:
Methyl 3-nitrosalicylate (7.19 g) obtained in the First Step was dissolved in tetrahydrofuran (100 ml) and ethyl acetate (50 ml), 5% palladium-carbon (water content 50%) (0.70 g) was added, and the mixture was stirred at room temperature for
2 hours under hydrogen atmosphere. Palladium-carbon was filtered off from the reaction suspension, the filtrate was concentrated, and n-hexane was added. The precipitated white solid was filtered off and dried to obtain the title compound (5.59 g).

Third Step
Production of methyl 3-chloroacetylaminosalicylate
Methyl 3-aminosalicylate (5.59 g) obtained in the Second Step was dissolved in chloroform (100 ml), aqueous saturated sodium hydrogen carbonate (50 ml) was added, further chloroacetyl chloride (3.2 ml) was added with vigorous stirring under ice-cooling, then the mixture was stirred for 1.5 hours under ice-cooling. The reaction mixture was partitioned, and the chloroform layer was dried over anhydrous magnesium sulfate and concentrated to obtain the title compound (white solid 8.07 g).

Fourth Step
Production of methyl-3,4-dihydro-3-oxo-2H-benzo[1,4]oxazine-8-carboxylate
Methyl 3-chloroacetylaminosalicylate (8.07 g) obtained in the Third Step was dissolved in N,N-dimethylformamide (80 ml), potassium carbonate (9.15 g) was added, and the mixture was stirred at 80°C for 1 hour. The reaction suspension was concentrated, and white solid precipitate obtained by adding water was filtered, washed with water and dried to obtain the title compound (6.37 g).

Fifth Step
Production of methyl 3,4-dihydro-3-oxo-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3,4-dihydro-3-oxo-2H-benzo[1,4]oxazine-8-carboxylate (2.07 g) obtained in the Fourth Step was suspended in tetrahydrofuran (20 ml), 1 M solution of borane-tetrahydrofuran complex in tetrahydrofuran (12 ml) was added under ice-cooling with stirring, and the mixture was refluxed for 1 hour in a stream of argon. The reaction mixture was cooled on ice, water (10 ml) and acetone (5 ml) were added in this order to terminate the reaction, further 6 N hydrochloric acid (10 ml) was added, and the mixture was stirred for 1 hour. The reaction mixture was adjusted to pH 8-9 by adding 4 N sodium hydroxide and aqueous saturated sodium hydrogen carbonate, and extracted with ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride solution in this order, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (1.82 g) as colorless oil.

Sixth Step

Production of methyl 4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3,4-dihydro-3-oxo-2H-benzo[1,4]oxazine-8-carboxylate (0.58 g) obtained in the Fifth Step was dissolved in toluene (7.5 ml), 2,3-dichloropyridine (0.44 g), tris(dibenzylideneacetone)dipalladium (69 mg), 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl (93 mg) and
cesium carbonate (1.47 g) were added in this order, and the mixture was stirred at 80°C for 24 hours. 2,3-Dichloropyridine (0.44 g), tris(dibenzylideneacetone)dipalladium (69 mg) and 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl (93 mg) were added again and stirring was further continued for 15 hours under excessive heating. After cooling, the reaction mixture was partitioned between the ethyl acetate and water. The obtained ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The thus obtained residue was purified by the use of silica gel chromatography (hexane - tetrahydrofuran = 3 : 1) to obtain the orange colored oily compound in the title (0.53 g).

Seventh Step

Production of 4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl 4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (0.53 g) obtained in the Sixth Step was dissolved in methanol (5 ml), 4 M sodium hydroxide solution (1 ml) was added, and the mixture was stirred at 60°C for 1 hour under heating. After cooling, the reaction mixture was neutralized by adding 1 M hydrochloric acid (4 ml), water (20 ml) was added, the precipitated solid was filtered off and dried to obtain the pale orange solid compound in the title (0.43 g).
Eighth Step

Production of

N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (0.43 g) obtained in the Seventh Step was dissolved in N,N-dimethylformamide (5 ml), 4-tert-butylaniline (0.24 g), 1-hydroxybenzotriazole (0.25 g) and 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride (0.31 g) were added in this order, and the mixture was stirred at room temperature for 1 hour. After adding water and saturated sodium hydrogen carbonate solution, the mixture was extracted with ethyl acetate. The ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The thus obtained residue was purified by the use of silica gel chromatography (hexane - ethyl acetate = 3 : 2) to obtain the white amorphous solid compound in the title (0.42 g).

$^1$H-NMR 400 MHz (CDCl$_3$) δ=1.33 (s, 9 H) 3.88-3.97 (m, 2H) 4.57-4.65 (m, 2H) 6.66 (dd, J=8.00, 1.28 Hz, 1H) 6.90 (t, J=7.88 Hz, 1H) 7.11 (dd, J=7.88, 4.87 Hz, 1H) 7.38 (d, J=8.58 Hz, II H) 7.60 (d, J=8.58 Hz, 2H) 7.77-7.82 (m, 2H) 8.37 (dd, J=4.75, 1.28 Hz, 1H) 9.57 (s, 1H).

Example 1-002

Production of

8-(4-tert-butylphenyl)carbamoyl-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylic acid:
First Step

Production of
N-(4-tert-butylphenyl)-3-nitro-2-hydroxybenzamide:

3-Nitrosalysilic acid (10.0 g) was dissolved in methylene chloride (100 ml), oxaly chloride (6.2 ml) and N,N-dimethylformamide (0.1 ml) were added, and the mixture was stirred at room temperature for 2 hours. The reaction mixture was concentrated, and the concentrate was added to a solution of 4-tert-butylaniline (5.2 g) and triethylamine (5.1 ml) in acetonitrile (100 ml), then the mixture was stirred at room temperature for 3 hours. After concentrating the reaction mixture, the reaction mixture was partitioned between chloroform and water. The obtained chloroform layer was washed with water, dried over anhydrous sodium sulfate and then concentrated. The thus obtained residue was purified by the use of silica gel chromatography (hexane - ethyl acetate = 1 : 1) to obtain the title compound (3.0 g).

Second Step

Production of 3-amino-N-(4-tert-butylphenyl)-2-hydroxybenzamide:

N-(4-tert-butylphenyl)-3-nitro-2-hydroxybenzamide (3.0 g) obtained in the First Step was dissolved in methanol (100 ml), iron(III) chloride hexahydrate (0.27 g) and active charcoal (0.5 g) were added, and the mixture was stirred at 80°C, and hydrazine hydrate (2.6 ml) was added dropwise. After stirring under heating for 1 hour, the reaction mixture was cooled and insoluble substance was filtered off, then the
solvent was evaporated in vacuo. The residue was partitioned between ethyl acetate and water. The obtained ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous magnesium sulfate and then concentrated. The thus obtained residue was dissolved in ethyl acetate, n-hexane was added, the precipitated solid was filtered off and dried to obtain the title compound (1.61 g).

Third Step

Production of ethyl

8-(4-tert-butylphenyl)carbamoyl-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylate:

3-Amino-N-(4-tert-butylphenyl)-2-hydroxybenzamide (1.6 g) obtained in the Second Step was dissolved in acetone (20 ml), ethyl 2,3-dibromopropionate (2.2 g) and potassium carbonate (2.3 g) were added, and the mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated, water was added, and the precipitated solid substance was filtered off. The thus obtained solid substance was suspended in and washed with a mixed solvent of ethyl acetate and n-hexane, filtered off and dried to obtain the title compound (1.67 g).

Fourth Step

Production of ethyl

8-(4-tert-butylphenyl)carbamoyl-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylate:

8-(4-tert-butylphenyl)carbamoyl-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylate (1.66 g) obtained in the Third Step
was dissolved in toluene (10 ml), 2,3-dichloropyridine (0.64 g), tris(dibenzylideneacetone)dipalladium (394 mg), 2,2'-bis(diphenylphosphino)-1,1'-binaphthyl (270 mg) and caesium carbonate (3.5 g) were added in this order, and the mixture was stirred at 80°C for 24 hours. The reaction mixture was diluted with ethyl acetate (10 ml), active charcoal (1 g) was added, and the mixture was stirred and filtered with Celite. The filtrate was concentrated and purified by the use of silica gel chromatography (gradient elution with n-hexane - ethyl acetate) to obtain the pale yellowish solid in the title (1.03 g).

Fifth Step
Production of
8-(4-tert-butylphenyl)carbamoyl-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylic acid:

8-(4-tert-butylphenyl)carbamoyl-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylate (1.03 g) obtained in the Fourth Step was dissolved in tetrahydrofuran (10 ml) and ethanol (10 ml), 2 M sodium hydroxide (2 ml) was added, and the mixture was stirred at room temperature for 3 hours. The reaction mixture was concentrated, water (20 ml) was added, the mixture was neutralized by adding 1N potassium hydrogen sulfate, and the precipitated solid substance was filtered off. The thus obtained solid substance was suspended and washed with a mixed solvent of ethyl acetate and n-hexane, filtered off and dried to obtain the title compound (0.60 g).
$^1$H-NMR 400 MHz (DMSO-d$_6$)  δ=1.29 (m, 9H) 3.62 (m, 1H) 4.18 (m, 1H) 4.70 (m, 1H) 6.47 (dd, J=8.07, 1.47 Hz, 1H) 6.79 (t, J=7.89 Hz, 1H) 7.21-7.35 (m, 4H) 7.84-7.92 (m, 2H) 7.98 (dd, J=7.70, 1.47 Hz, 1H) 8.43 (dd, J=4.99, 1.51 Hz, 1H) 12.34 (s, 1H).

Example 1-003

Production of

N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-3-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl 3-nitro-2-(2-oxopropoxy)benzoate:

Methyl 3-nitrosalicylate (1.95 g) obtained in the First Step of Example 1-001 was dissolved in N,N-dimethylformamide (20 ml), potassium carbonate (2.73 g) and bromoacetone (1.1 ml) were added, and the mixture was stirred at room temperature overnight. The reaction mixture was partitioned between water and ethyl acetate, and the ethyl acetate layer was washed twice with saturated sodium chloride solution, dried over anhydrous sodium sulfate and concentrated to obtain brownish oily compound in the title (2.46 g).

Second Step

Production of methyl 3-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3-nitro-2-(2-oxopropoxy)benzoate (2.46 g) obtained in the First Step was dissolved in tetrahydrofuran (25 ml), 10% palladium carbon (containing 50% water) (0.25 g) was added, and the mixture was stirred at room temperature for 24 hours under hydrogen atmosphere. The catalyst was
filtered off from the reaction mixture and the filtrate was concentrated. The concentrate was purified by the use of silica gel chromatography (n-hexane - ethyl acetate = 1 : 1) to obtain pale yellowish oily compound in the title (0.45 g).

Third Step
Production of methyl
4-(3-chloropyridin-2-yl)-3-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:
methyl
2-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (0.62 g) obtained in the method of Second Step was subjected to the reaction in a similar manner as in the Fourth Step in Example 2, and purified by the use of silica gel chromatography (n-hexane - acetone = 5 : 1) to obtain pale yellowish oily compound in the title (0.10 g).

Fourth Step
Production of
4-(3-chloropyridin-2-yl)-3-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:
Methyl
4-(3-chloropyridin-2-yl)-3-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (0.09 g) obtained in the Third Step was dissolved in tetrahydrofuran (1 ml) and methanol (1 ml), 4 M sodium hydroxide solution (0.5 ml) was added, and the mixture was stirred at 60°C for 1 hour. The reaction mixture was neutralized with 1 M hydrochloric acid and the solvent was evaporated in vacuo. The residue was partitioned between
water and ethyl acetate, and the ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and concentrated to obtain pale yellowish oily compound in the title (75 mg).

Fifth Step
Production of
N-(4-tert-butyphenyl)-4-(3-chloropyridin-2-yl)-3-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(3-Chloropyridin-2-yl)-3-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (75 mg) obtained in the Fourth Step was subjected to the reaction in a similar manner as in the Eighth Step in Example 1-001, and water and saturated aqueous sodium hydrogen carbonate solution were added in this order. The precipitated pale brownish solid substance was filtered off and dried to obtain the title compound (95 mg).

$^1$H-NMR 400 MHz (DMSO-d$_6$) δ=1.23 (d, J=6.49 Hz, 3H) 1.28 (s, 9H) 4.07-4.15 (m, 1H) 4.22-4.32 (m, 2H) 6.38 (dd, J=8.12, 1.62 Hz, 1H) 6.80 (t, J=7.88 Hz, 1H) 7.04 (dd, J=7.54, 1.51 Hz, 1H) 7.31-7.39 (m, 3H) 7.64-7.71 (m, 2H) 8.07 (dd, J=7.88, 1.62 Hz, 1H) 8.45 (dd, J=4.64, 1.62 Hz, 1H) 10.10 (s, 1H).

Example 1-004
Production of
N-(4-tert-butyphenyl)-1-(3-chloropyridin-2-yl)-4-methyl-1,2,3,4-tetrahydroquinoxaline-5-carboxamide:

First Step
Production of methyl 2-chloro-3-nitrobenzoate:
2-Chloro-3-nitrobenzoic acid (15.0 g) was subjected to the reaction in a similar manner as in the First Step in Example 1-001 to obtain pale yellowish compound in the title (9.0 g).

Second Step

Production of methyl 2-[(ethoxycarbonylmethyl)·(N-methyl)amino]-3-nitrobenzoate:

Methyl 2-chloro-3-nitrobenzoate (2.0 g) obtained in the First Step was dissolved in n-butanol (20 ml), sodium carbonate (2.46 g) and sarcosine ethyl ester hydrochloride (2.14 g) were added, and the mixture was refluxed for 4.5 hours with stirring. The cooled reaction mixture was poured into a mixture of 1 M hydrochloric acid (50 ml)/ethyl acetate (50 ml) under ice-cooling with stirring. The ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate, and concentrated to obtain red oily substance containing the title compound which was supplied to the subsequent step without purification.

Third Step

Production of methyl 4-methyl-2-oxo-1,2,3,4-tetrahydroquinoxaline-5-carboxylate:

The oily substance containing methyl 2-[(ethoxycarbonylmethyl)·(N-methyl)amino]-3-nitrobenzoate obtained in the Second Step was dissolved in methanol (20 ml), 5% palladium carbon (0.2 g) was added, and the mixture was stirred at room temperature for 2 hours under hydrogen
atmosphere. The catalyst was filtered off from the reaction suspension. The filtrate was concentrated and then purified by the use of silica gel chromatography (n-hexane - ethyl acetate = 1:3), diethyl ether was added, and the precipitated pale brownish solid substance was filtered off and dried to obtain the title compound (0.41 g).

Fourth Step

Production of methyl 4-methyl-1,2,3,4-tetrahydroquinoxaline-5-carboxylate:

Methyl 4-methyl-2-oxo-1,2,3,4-tetrahydroquinoxaline-5-carboxylate (0.41 g) obtained in the Third Step was subjected to the reaction in a similar manner as in the Fifth Step in Example 1 to obtain yellowish oily compound in the title (203 mg).

Fifth Step

Production of methyl 1-(3-chloropyridin-2-yl)-4-methyl-1,2,3,4-tetrahydroquinoxaline-5-carboxylate:

Methyl 4-methyl-1,2,3,4-tetrahydroquinoxaline-5-carboxylate (206 mg) obtained in the Fourth Step was subjected to the reaction in a similar manner as in the Fourth Step in Example 1-002 to obtain pale yellowish brown solid compound in the title (179 mg).

Sixth Step

Production of methyl 1-(3-chloropyridin-2-yl)-4-methyl-1,2,3,4-tetrahydroquinoxaline-5-carboxylic acid:
Methyl 1-(3-chloropyridin-2-yl)-
4-methyl-1,2,3,4-tetrahydroquinoxaline-5-carboxylate (176 mg)
obtained in the Fifth Step was subjected to the reaction in a
similar manner as in the Fourth Step in Example 1-003 to obtain
pale yellowish brown solid compound in the title (130 mg).

Seventh Step

Production of
N-(4-tert-butylphenyl)-1-(3-chloropyridin-2-yl)-4-methyl-1,
2,3,4-tetrahydroquinoxaline-5-carboxamide:

Methyl 1-(3-chloropyridin-2-yl)-
4-methyl-1,2,3,4-tetrahydroquinoxaline-5-carboxylic acid (130
mg) obtained in the Sixth Step was subjected to the reaction
in a similar manner as in the Eighth Step in Example 1-001, and
the reaction mixture was partitioned between aqueous saturated
sodium hydrogencarbonate solution and ethyl acetate. The ethyl
acetate layer was washed with saturated sodium chloride
solution, washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated.
The residue was purified by the use of silica gel
chromatography (n-hexane - ethyl acetate = 3:2) to obtain
pale yellowish amorphous compound in the title (100 mg).

$^1$H-NMR 400 MHz (DMSO-$d_6$) δ=1.28 (s, 9H) 2.83 (s, 3H) 3.34-3.39
(m, 2H) 3.69-3.74 (m, II H) 6.33 (dd, J=8.12, 1.39, 1 Hz, 1H)
6.72 (t, J=7.77 Hz, 1H) 7.03 (dd, J=7.65, 1.39 Hz, 1H) 7.30
(dd, J=7.88, 4.64 HzH) 7.33-7.38 (m, 2H) 7.63-7.68 (m, 2H)
8.02 dd, J=7.88, 1.62 Hz, 1H) 8.44 (dd, J=4.75, 1.74 Hz, 1H) 10.62 (s, 1H).

Example 1-005

Production of N-(4-tert-butylphenyl)-9-(3-chloropyridin-2-yl)-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxamide:

First Step

Production of methyl 5-chloro-3-nitrosalicylate

Methyl 5-chlorosalicylate (5.0 g) was dissolved in concentrated sulfuric acid (15 ml), a mixture of concentrated nitric acid (1.2 ml) and concentrated sulfuric acid (1.2 ml) was added dropwise with stirring under ice-cooling, and further stirred after finishing the dropwise addition. The reaction mixture was poured into ice-water, the precipitated solid substance was filtered off, washed with water, then and dried to obtain the title compound (5.51 g).

Second Step

Production of methyl 3-amino-5-chlorosalicylate

Methyl 5-chloro-3-nitrosalicylate (9.16 g) obtained in the method of First Step was subjected to the reaction in a similar manner as in the Second Step in Example 2, and purified by the use of silica gel chromatography (n-hexane - ethyl acetate = 1 : 1) to obtain white solid compound in the title (4.46 g).

Third Step

Production of methyl 3-acetamide-5-chlorosalicylate
Methyl 3-amino-5-chlorosalicylate (4.46 g) obtained in the Second Step was dissolved in tetrahydrofuran (50 ml), and pyridine (2 ml) and acetyl chloride (1.6 ml) were added with stirring under ice-cooling. After 30 minutes, the reaction mixture was concentrated and partitioned between water and ethyl acetate. The ethyl acetate layer was washed with 1 M hydrochloric acid, saturated sodium chloride solution, saturated aqueous sodium hydrogen carbonate and saturated sodium chloride solution, in this order, dried over anhydrous sodium sulfate and concentrated to obtain the white solid compound in the title (4.86 g).

Fourth Step

Production of methyl 9-acetyl-2-chloro-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylate:

Methyl 3-acetamide-5-chlorosalicylate obtained in the Third Step was dissolved in N,N-dimethylformamide (10 ml), potassium carbonate (1.38 g) and 1-bromo-3-chloropropane (2.0 ml) were added, and the mixture was stirred at room temperature for 1 hour. Then the temperature to 120°C was increased and the mixture was stirred for 3 hours. After cooling the reaction mixture, water was added, and the mixture was extracted with ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride solution, in this order, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel chromatography.
(n-hexane - ethyl acetate = 1 : 1) to obtain the white solid compound in the title (630 mg).

Fifth Step

Production of methyl 9-acetyl-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylate:

Methyl 9-acetyl-2-chloro-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylate (1.9 g) obtained by the same method of the Fourth Step was dissolved in tetrahydrofuran (20 ml) and methanol (20 ml), 5% palladium carbon (0.2 g) and triethylamine (1.2 ml) were added, and the mixture was stirred at room temperature for 45 hours under hydrogen atmosphere. The catalyst was filtered off from the reaction suspension, the filtrate was concentrated, and the concentrate was partitioned between water and ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride solution in this order, dried over anhydrous sodium sulfate and concentrated to obtain the colorless oily compound in the title (1.45 g).

Sixth Step

Production of 6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylic acid:

Water (25 ml) and concentrated sulfuric acid (5 ml) were added to methyl 9-acetyl-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene
-4-carboxylate (1.43 g) obtained in the Fifth Step and the mixture was refluxed with stirring overnight. The reaction mixture was left to cool, neutralized by adding 2 M sodium hydroxide solution to slightly acidic and extracted twice with ethyl acetate. The ethyl acetate layer was washed twice with saturated sodium chloride solution, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (530 mg). The aqueous layer was extracted with tetrahydrofuran, dried over anhydrous sulfate and concentrated to further obtain the compound (320 mg).

Seventh Step

Production of ethyl

6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylate:

6,7,8,9-Tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxyl acid (850 mg) was dissolved in ethanol (30 ml), concentrated sulfuric acid (2 ml) was added, and the mixture was refluxed with stirring for 2 hours. The reaction mixture was cooled, neutralized with saturated sodium hydrogen carbonate solution and extracted with ethyl acetate. The ethyl acetate layer was concentrated to obtain the oily compound in the title (721 mg).

Eighth Step

Production of ethyl 9-(3-chloropyridin-2-yl)-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylate:
Ethyl
6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylate (720 mg) obtained in the Seventh Step was subjected to the reaction in a similar manner as in the Fourth Step in Example 1-002, and the crude product was purified by the use of silica gel chromatography (n-hexane - ethyl acetate = 3 : 1) to obtain the oily compound in the title (154 mg).

Ninth Step

Production of 9-(3-chloropyridin-2-yl)-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylic acid:

Ethyl 9-(3-chloropyridin-2-yl)-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylate (150 mg) was dissolved in tetrahydrofuran (2 ml) and ethanol (2 ml), 2 M sodium hydroxide solution (2 ml) was added, and the mixture was stirred at 60°C for 1.5 hours. The reaction mixture was concentrated, acidified with 1 M aqueous potassium hydrogen sulfate and extracted twice with ethyl acetate. The ethyl acetate layer was washed with saturated sodium chloride solution, washed over anhydrous sodium sulfate and concentrated to obtain the oily compound in the title (102 mg).

Tenth Step

Production of
N-(4-tert-butylphenyl)-9-(3-chloropyridin-2-yl)-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxamide:
9-(3-Chloropyridin-2-yl)-
6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxylic acid (100 mg) obtained in the Ninth Step was subjected to
the reaction in a similar manner as in the Eighth Step in Example
1-001, and the crude product was purified by the use of silica
gel chromatography (n-hexane - ethyl acetate = 2 : 1), n-hexane
was added, precipitated white solid substance was filtered off
and dried to obtain the title compound (35 mg).

^1^H-NMR 400 MHz (DMSO-d_6) δ=1.25 (s, 9H) 1.95 (m, 2H) 3.97 (m,
2H) 4.27 (m, 2H) 6.61 (dd, J=8.00, 1.51 Hz, 1H) 6.93 (t, J=7.77
Hz, 1H) 7.06 (dd, J=7.77, 4.75 Hz, 1H) 7.26 (dd, J=7.54, 1.51
Hz, 1H) 7.30-7.38 (m, 2H) 7.63 (ddd, J=9.04, 2.55, 2.32 Hz, 2H)
7.76 (dd, J=7.65, 1.62 Hz, 1H) 8.31 (dd, J=4.64, 1.62 Hz, 1H)
10.15 (s, 1H).

Example 1-006

In the method similar to Example 1-001, corresponding
carboxylic acid and amine were used to obtain the compound of
Example 1-006 shown in the following table.

Example 1-007

In the method similar to Example 1-001, corresponding
carboxylic acid and amine were used to obtain the compound of
Example 1-007 shown in the following table.

Example 1-008

In the method similar to Example 1-001, corresponding
carboxylic acid and amine were used to obtain the compound of
Example 1-008 shown in the following table.
Example 1 - 009
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 009.

Example 1 - 010
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 010.

Example 1 - 011
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 011.

Example 1 - 012
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 012.

Example 1 - 013
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 013.

Example 1 - 014
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 014.
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 015.

Example 1 - 016

In the Sixth Step in Example 1 - 001, 2-bromopyridine was used in the place of 2,3-dichloropyridine, and others were treated by the same manner as in the example to obtain the compound of Example 1 - 016.

Example 1 - 017

In the Sixth Step in Example 1 - 001, 2-chloro-3-trifluoromethylpyridine was used in the place of 2,3-dichloropyridine, and others were treated by the same manner as in the example to obtain the compound of Example 1 - 017.

Example 1 - 018

In the Third Step in Example 1 - 001, 2-bromoisobutyl bromide was used in the place chloroacetyl chloride, and others were treated by the same manner as in the example to obtain the compound of Example 1 - 018.

Example 1 - 019

In the Third Step in Example 1 - 001, 2-chloropropionyl chloride was used in the place chloroacetyl chloride, and others were treated by the same manner as in the example to obtain the compound of Example 1 - 019.
3-amino-5-chlorosalicylic acid produced by the method of the Second Step in Example 1 005 was used in the method subsequent to the Third Step in Example 1 - 001 to obtain the compound of Example 1 - 020.

Example 1 - 021

Production of
4 - (3-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(3-chloropyridine-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (100 mg) obtained in the Seventh Step in Example 1 - 001 was dissolved in tetrahydrofuran (1 ml), oxalyl chloride (0.05 ml) and N,N-dimethylformamide (0.01 ml) were added, the mixture was stirred at room temperature for 30 minutes, and the reaction mixture was concentrated to obtain acid chloride as yellowish solid. Tetrahydrofuran (1 ml) and 1 M sodium hydroxide solution (1 ml) were added to 4-trifluoroaniline (48 mg), and a solution of the acid chloride in tetrahydrofuran (1 ml) was added with stirring under ice-cooling, and the reaction mixture was stirred at room temperature for 30 minutes. The reaction mixture was concentrated, partitioned between ethyl acetate and water, and the ethyl acetate layer was washed with 1 M sodium hydroxide and saturated sodium chloride solution in this order, dried over anhydrous magnesium sulfate and then concentrated. The residue was suspended in n-hexane, filtered off and dried to obtain the title compound (72 mg).

Example 1 - 022
In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 022.

Example 1 - 023

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 023.

Example 1 - 024

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 024.

Example 1 - 025

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 025.

Example 1 - 026

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 026.

Example 1 - 027

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 027.

Example 1 - 028

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 028.
Example 1 - 029

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 029.

Example 1 - 030

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 030.

Example 1 - 031

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 031.

Example 1 - 032

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 032.

Example 1 - 033

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 033.

Example 1 - 034

Production of
4-(3-chloropyridine-2-yl)-N-(methoxycarbonylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

Methyl 4-aminobenzoate (260 mg) was used in the method similar to Example 1 - 021 to obtain the title compound (515 mg).
Example 1 - 035

Production of

N-(4-carboxyphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(3-Chloropyridin-2-yl)-N-(methoxycarbonylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (500 mg) produced in Example 1 - 034 was dissolved in tetrahydrofuran (5 ml) and methanol (5 ml), 4 M sodium hydroxide solution (1 ml) was added, and the mixture was stirred at 60°C for 5 hours. The reaction mixture was neutralized by adding 1 M hydrochloric acid and then concentrated. The concentrate was diluted with water and stirred, and white solid substance was collected by filtration and dried to obtain the title compound (433 mg).

Example 1 - 036

Production of

N-(4-carbamoylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4] oxazine-8-carboxamide:

N-(4-carboxyphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4] oxazine-8-carboxamide (100 mg) obtained in Example 1 - 035 was dissolved in N,N-dimethylformamide (2 ml), ammonium chloride (65 mg), 1-hydroxybenzotriazole (56 mg), 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride (70 mg) and triethylamine (0.20 ml) were added in this order and the mixture was stirred at room temperature for 10 hours. After adding water and saturated sodium hydrogencarbonate solution to the reaction mixture, the mixture was partitioned by adding ethyl acetate. After ethyl
acetate layer was washed with water, concentrated and dried, the concentrate was suspended in and washed with water, the white solid substance was collected by filtration and dried to obtain the title compound (80 mg).

Example 1 - 037

In the method similar to Example 1 - 036, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 037.

Example 1 - 038

In the method similar to Example 1 - 036, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 038.

Example 1 - 039

Production of
N-(4-acetylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4'-aminoacetophenone (279 mg) was used in the similar method of Example 1 - 021 to obtain the title compound (673 mg).

Example 1 - 040

Production of
4-(3-chloropyridin-2-yl)-N-[4-((1-hydroxy-1-methyl)ethyl)phenyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

N-4(4-acetylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (150 mg) obtained in Example 1 - 039 was dissolved in tetrahydrofuran (15 ml), 1 M methyl magnesium bromide-tetrahydrofuran solution (1.1 ml) was added under ice-cooling, and then the mixture was stirred
at room temperature for 30 minutes. Then, 1 M methyl magnesium bromide-tetrahydrofuran solution (0.37 ml) was further added and stirred and the mixture was at room temperature for 30 minutes. The reaction mixture was cooled on ice, the reaction was terminated by adding saturated aqueous ammonium chloride, and the mixture was extracted with ethyl acetate. The ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel chromatography (n-hexane-ethyl acetate=1:4) to obtain the white amorphous compound in the title (79 mg).

Example 1 - 041
Production of 4-(3-chloropyridin-2-yl)-N-[4-(1-hydroxy-1-methyl)propyl]phenyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

In the reaction similar to Example 1 - 040, 2 M ethyl magnesium bromide was used to obtain the white amorphous compound in the title (100 mg) from N-(4-acetylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (150 mg).

Example 1 - 042
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 042.

Example 1 - 043
In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 043.
Example 1 - 044

In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 044.
Example 1 - 045

In the method similar to Example 1 - 001, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 045.
Example 1 - 046

In the Sixth Step in Example 1 - 001, 4-bromopyridine was used in the place of 2,3-dichloropyridine and others were treated by the same manner as in the example to obtain the compound of Example 1 - 046.
Example 1 - 047

In the Sixth Step in Example 1 - 001, 2-bromo-3-picoline was used in the place of 2,3-dichloropyridine and others were treated by the same manner as in the example to obtain the compound of Example 1 - 047.
Example 1 - 048

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 048.
Example 1 - 049
In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 049.

Example 1 - 050

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 050.

Example 1 - 051

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 051.

Example 1 - 052

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 052.

Example 1 - 053

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 053.

Example 1 - 054

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 054.

Example 1 - 055

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 055.
Example 1 - 056

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 056.

Example 1 - 057

In the method similar to Example 1 - 021, corresponding carboxylic acid and amine were used to obtain the compound of Example 1 - 057.

Example 1 - 058

Production of N-(4-tert-butylphenyl)-4-(4-chloropyridine-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of 4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

In the Sixth Step in Example 1 - 001, 2-chloro-4-picoline (890 mg) was used in the place of 2,3-dichloropyridine and others were treated by the same manner as in the Seventh Step of Example 1 - 001 to obtain the title compound (690 mg).

Second Step

Production of N-(4-tert-butylphenyl)-4-(4-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

N-(4-tert-butylphenyl)-4-(4-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (150 mg) obtained
in the process 1-001 was used in the similar method in Example 21 to obtain the title compound (130 mg).

Example 1-059

Example 1-059 compound shown in the following table was obtained from the corresponding bromopicoline by applying the similar method as of Example 21.

Example 1-060

Example 1-060 compound shown in the following table was obtained from the corresponding bromopicoline by applying the similar method as of Example 1-021.

Example 1-061 to Example 1-236

A compound of Examples 1-061 to 1-236 hereinbelow were obtained by similarly performing any methods described in the general processes A to C for producing the compound and/or a method described in Examples 1-001 to 1-060 hereinbefore.

Example 2

Example 2-01

Production of
N-(benzo[1,3]dioxol-5-yl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl
4-{3-chloro-5-\{(tetrahydro pyran-2-yl)oxymethylpyridin-2-yl\}}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

3,4-Dihydro-2H-benzo[1,4]oxazine-8-carboxylate (2.0 g) obtained by the method of Fifth Step in Example 1-001 was dissolved in toluene (30 ml),
2,3-dichloro-5-(tetrahydropyran-2-yl)oxymethylpyridine (2.76 g), palladium acetate (0.25 g),
2,2'-bis(diphenylphosphino)-1,1'-binaphthyl (0.7 g) and
caesium carbonate (7.4 g) were added, and the mixture was
stirred at 90°C overnight. The reaction mixture was filtered
and concentrated, and the thus obtained residue was purified
by the use of silica gel column chromatography (hexane -
tetrahydrofuran = 2 : 1) to obtain oily compound in the title
(2.48 g).

Second Step
Production of
4-{3-chloro-5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-
3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl
4-{3-chloro-5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-
3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (2.48 g)
obtained in the First Step was subjected to the reaction similar
to that of the Fourth Step in Example 1-003 to obtain the title
compound (1.33 g).

Third Step
Production of
N-(benzo[1,3]dioxol-5-yl)-4-{3-chloro-5-(tetrahydropyran-2-
yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-
8-carboxamide:

4-{3-chloro-5-(tetrahydropyran-2-yl)oxymethylpyridin-2-
yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (0.20
g) obtained in the Second Step and
5-amino-benzo[1,3]dioxolane (0.10 g) were subjected to the reaction similar to that of the Eighth Step in Example 1-001 to obtain the title compound (0.21 g).

Fourth Step

Production of

\[ N\cdot(benzo[1,3]dioxol-5-yl)\cdot4\cdot(3\cdot\text{chloro}-5\cdot\text{hydroxymethylpyridin}-2\cdot\text{yl})\cdot3,4\cdot\text{dihydro}-2\text{H}-\text{benzo}[1,4]\text{oxazine}-8\cdot\text{carboxamide} \]

\[ N\cdot(benzo[1,3]dioxol-5-yl)\cdot4\cdot(3\cdot\text{chloro}-5\cdot(\text{tetrahydropyran}-2\cdot\text{yl})\text{oxymethylpyridin}-2\cdot\text{yl})\cdot3,4\cdot\text{dihydro}-2\text{H}-\text{benzo}[1,4]\text{oxazine}-8\cdot\text{carboxamide} \]

(0.21 g) obtained in the Third Step was dissolved in tetrahydrofuran (3 ml), 6 M hydrochloric acid (3 ml) was added, and the mixture was stirred at room temperature overnight. The reaction mixture was concentrated, and the residue was purified by the use of silica gel column chromatography to obtain the title compound (0.12 g).

Example 2-02

Production of

\[ N\cdot(2,3\cdot\text{dihydrobenzo}[1,4]\text{dioxin}-6\cdot\text{yl})\cdot4\cdot(3\cdot\text{chloro}-5\cdot\text{hydroxymethylpyridin}-2\cdot\text{yl})\cdot3,4\cdot\text{dihydro}-2\text{H}-\text{benzo}[1,4]\text{oxazine}-8\cdot\text{carboxamide} \]

In the Third Step of Example 2-01, 6-amino-2,3-dihydrobenzo[1,4]dioxin (111 mg) was used in the place of 5-amino-benzo[1,3]dioxolane, and others were treated by the same manner as in the process to obtain the title compound (109 mg).

Example 2-03
Production of
N-(4-trifluoromethoxyphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

In the Third Step of Example 2-01, 4-trifluoromethoxyaniline (87 mg) was used in the place of 5-amino-benzo[1,3]dioxolane, and others were treated by the same manner as in the process to obtain the title compound (101 mg).

Example 2-04

Production of
N-(3-chloro-4-trifluoromethoxyphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

In the Third Step of Example 2-01, 3-chloro-4-trifluoromethoxyaniline (87 mg) was used in the place of 5-amino-benzo[1,3]dioxolane, and others were treated by the same manner as in the process to obtain the title compound (58 mg).

Example 2-05

Production of
N-(4-trifluoromethylphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of
N-(4-trifluoromethylphenyl)-4-(3-chloro-5-(tetrahydropyra
n-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-{3-Chloro-5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (300 mg) obtained in the Second Step of Example 2-01 was dissolved in pyridine (5 ml), thionyl chloride (88 mg) was added, and the mixture was stirred at 80°C for 1 hour. The reaction mixture was cooled to room temperature, 4-trifluoromethylaniline (80 mg) was added, and the mixture was stirred for 1 hour. The reaction mixture was partitioned between water and ethyl acetate, and the ethyl acetate layer was washed with water, diluted aqueous potassium hydrogen sulfate, diluted aqueous sodium hydroxide and saturated sodium chloride solution, dried over magnesium sulfate and then concentrated to obtain residue containing the title compound, which was supplied to the subsequent process without purification.

Second Step

Production of

N-(4-trifluoromethylphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

The concentrated residue obtained in the First Step was dissolved in tetrahydrofuran (3 ml), 6 M hydrochloric acid (3 ml) was added, and the mixture was stirred at room temperature for 1 hour. The reaction mixture was partitioned between water and ethyl acetate, and the ethyl acetate layer was washed with
water and saturated sodium chloride solution in this order, dried over anhydrous magnesium sulfate and then concentrated. The thus obtained concentrated residue was purified by the use of silica gel chromatography to obtain the title compound (71 mg).

Example 2-06

Production of
N-(3-fluoro-4-trifluoromethylphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

In the First Step of Example 2-05, 3-fluoro-4-trifluoromethylaniline (110 mg) was used in the place of 4-trifluoromethylaniline, and others were treated by the same manner as in the process to obtain the title compound (50 mg).

Example 2-07

Production of
N-(4-trifluoromethylphenyl)-4-(5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of
4-(5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl
4-(5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (560 mg), which was obtained by using
2-chloro-5-(tetrahydropyran-2-yl)oxymethylpyridine in the place of
2,3-dichloro-5-(tetrahydropyran-2-yl)oxymethylpyridine in the
First Step of Example 2-01, was dissolved in methanol (8 ml),
4 M sodium hydroxide solution (1.1 ml) was added, and the
mixture was stirred at 60°C for 2 hours. The reaction mixture
was concentrated, and the concentrate was partitioned by
adding water and ethyl acetate. The aqueous layer was adjusted
to pH 4 by adding hydrochloric acid and was extracted with ethyl
acetate. The ethyl acetate layer was dried over anhydrous
sodium sulfate and then concentrated. Precipitated solid
obtained by adding diisopropyl ether was collected by
filtration and dried to obtain the title compound (420 mg).

Second Step

Production of
N-(4-trifluoromethylphenyl)-4-{5-(tetrahydropyran-2-yl)oxym
ethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carbo
xamide:

A reaction similar to that of the First Step of Example
2-05 was performed by using
4-{5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihyd
ro-2H-benzo[1,4]oxazine-8-carboxylic acid (200 g) obtained in
the First Step of Example 2-05 to obtain a concentrated residue
containing the title compound.

Third Step

Production of
N-(4-trifluoromethylphenyl)-4-(5-hydroxymethylpyridin-2-y
1)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide hydrochloride:

The concentrated residue containing N-(4-trifluoromethylphenyl)-4-{5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide was dissolved in methanol (2 ml) and tetrahydrofuran (2 ml), 6 M hydrochloric acid (1 ml) was added, and the mixture was stirred at room temperature for 1 hour. The reaction mixture was concentrated, water and diethyl ether were added, then precipitated yellow solid substance was collected by filtration and dried to obtain the title compound (170 mg).

Example 2.08

Production of 4-(5-hydroxymethylpyridin-2-yl)-N-(4-isobutoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-{5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (200 mg) obtained in the First Step of Example 2.07 and 4-isobutoxyaniline (110 mg) were used and the others were subjected to the reaction similar to Example 2.05 to obtain the title compound (140 mg).

Example 2.09

Production of 4-(3-chloro-5-methoxymethylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of methyl
4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl
4-(3-chloro-5-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (1.3 g) obtained in the First Step of Example 2-01 was dissolved in tetrahydrofuran (5 ml), 6 M hydrochloric acid was (5 ml) added, and the mixture was stirred at room temperature for 2 hours. The reaction mixture was neutralized and extracted with ethyl acetate. The extract was concentrated to obtain a concentrated residue containing the title compound.

Second Step

Production of methyl
4-(3-chloro-5-methoxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

The concentrated residue obtained in the First Step was dissolved in N,N-dimethylformamide (5 ml), sodium hydride (50%) (0.32 g) was added, and the mixture was stirred at room temperature until foaming was terminated. Subsequently, sodium iodide (1 ml) was added and the mixture was further stirred for 1 hour. The reaction mixture was partitioned between water and ethyl acetate, and the ethyl acetate layer was washed with water, dried over anhydrous sodium sulfate and then concentrated. The concentrate was purified by the use of silica gel chromatography to obtain the oily compound in the title (1.20 g).
Third Step

Production of
4-(3-chloro-5-methoxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl
4-(3-chloro-5-methoxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (1.20 g) obtained in the Second Step was dissolved in tetrahydrofuran (10 ml) and methanol (10 ml), 2 M sodium hydroxide (5 ml) was added, and the mixture was stirred at 60°C for 1 hour. The reaction mixture was concentrated and neutralized to obtain black colored solid substance by filtration and dried to obtain the title compound (915 mg).

Fourth Step

Production of
4-(3-chloro-5-methoxymethylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(3-chloro-5-methoxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (200 mg) obtained in the Third Step was dissolved in chloroform (2 ml), oxalyl chloride (60 μl) and N,N-dimethylformamide (one drop) were added, and the mixture was stirred at room temperature for 1 hour. The reaction mixture was concentrated, pyridine (2 ml) and 4-trifluoromethylaniline (75 μl) were added, and the mixture was stirred at room temperature for 1 hour. The reaction mixture was partitioned between water and ethyl
acetate, and the ethyl acetate layer was washed with water, dried over anhydrous magnesium sulfate and then concentrated. Precipitated solid obtained by adding n-hexane and diethyl ether to the concentrate was collected by filtration and dried to obtain the title compound (64 mg).

Example 2-10

Production of

N-(4-trifluoromethylphenyl)-4-(4-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of

4-{4-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl

4-{4-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate, which was obtained by the process, in which, in the First Step of Example 2-01, the reaction was performed by the similar manner using 2-chloro-4-(tetrahydropyran-2-yl)oxymethylpyridine (1.52g) in the place of 2,3-dichloro-5-(tetrahydropyran-2-yl)oxymethylpyridine, and the reaction mixture was filtered, concentrated and purified by the use of silica gel column chromatography, was dissolved in methanol (10 ml) and tetrahydrofuran (10 ml), 4 M sodium hydroxide solution (5 ml) was added, and the mixture was stirred at 60°C for 1.5 hour. The reaction mixture was neutralized and then concentrated. The concentrate was
partitioned by adding water and ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride solution in this order, dried over anhydrous magnesium sulfate and concentrated to obtain the title compound (913 mg).

Second Step

Production of

N-(4-trifluoromethylphenyl)-4-(4-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-{4-[(tetrahydrofuran-2-yl)oxymethylpyridin-2-yl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (300 mg) obtained in the First Step was dissolved in pyridine (5 ml), thionyl chloride (89 µl) was added, and the mixture was stirred at room temperature for 2 hours. Subsequently, 4-trifluoromethylaniline (102 µl) was added and stirred at the same temperature overnight. The reaction mixture was partitioned between water and ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride in this order, dried over anhydrous magnesium sulfate and then concentrated. The concentrate was purified by the use of silica gel chromatography. To the purified concentrated substance was added tetrahydrofuran (5 ml) and 6 M hydrochloric acid (2 ml) and stirred at room temperature for 2 hours. The reaction mixture was partitioned between water and ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride solution in this order, dried over anhydrous magnesium sulfate and then concentrated.
The concentrate was purified by the use of silica gel chromatography to obtain the title compound (12 mg).

Example 2-11

Production of

N-(4-trifluoromethoxyphenyl)-4-(3-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl

4-{3-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

In the First Step of Example 2-01, the similar reaction was performed by using

2-chloro-3-(tetrahydropyran-2-yl)oxymethylpyridine (712 mg) in the place of

2,3-dichloro-5-(tetrahydropyran-2-yl)oxymethylpyridine to obtain the title compound (536 mg).

Second Step

Production of

4-{3-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl

4-{3-(tetrahydropyran-2-yl)oxymethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (536 mg) obtained in the First Step was dissolved in ethanol (10 ml), 2 M sodium hydroxide solution (2 ml) was added, and the mixture was stirred at 60°C for 1 hour. The reaction mixture was neutralized by
adding hydrochloric acid and concentrated, partitioned between water and ethyl acetate. The thus obtained ethyl acetate layer was dried over anhydrous sodium sulfate and concentrated to obtain the title compound (493 mg).

Third Step

Production of
N-(4-trifluoromethoxyphenyl)-4-{3-(tetrahydro-2-yl)oxy methylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-{3-(Tetrahydro-2-yl)oxyethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (266 mg) obtained in the Second Step and 4-trifluoromethoxyaniline (97 µl) was subjected to the reaction similar to that of the Eighth Step of Example 1-001 to obtain the title compound (222 mg).

Fourth Step

Production of
N-(4-trifluoromethoxyphenyl)-4-(3-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

N-(4-trifluoromethoxyphenyl)-4-{3-(tetrahydro-2-yl)oxyethylpyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (217 mg) obtained in the Third Step was dissolved in methanol (5 ml), 6 M hydrochloric acid (0.3 ml) added, and the mixture was stirred at room temperature for 1 hour. The reaction mixture was diluted with water, neutralized by adding sodium hydroxide solution and extracted with ethyl acetate. The ethyl acetate layer was dried over anhydrous sodium sulfate, concentrated and purified by the
use of silica gel chromatography to obtain the title compound (164 mg).

Example 2-12

Production of

\((+)-N-(4\text{-}{\text{trifluoromethylphenyl}})-4-\{3\text{-}{\text{chloro-5-(1\text{-}{\text{hydroxyethyl}}}}\text{-}pyridin-2\text{-}{\text{yl}}\}\text{-}3,4\text{-}{\text{dihydro-2H-benzo[1,4]oxazine-8-carboxamide}}:\)

First Step

Production of methyl 3-acetamide-2-hydroxybenzoate:

Methyl 3-aminosalicylate (10 g) obtained in the Second Step of Example 1-001 was dissolved in ethyl acetate (30 ml), water (30 ml) and sodium hydrogen carbonate (5.54 g) were added, and acetyl chloride (5 ml) was added with stirring under ice-cooling. After stirring at room temperature for 30 minutes, the ethyl acetate layer was dried over anhydrous sodium sulfate and concentrated to obtain the title compound (11.47 g).

Second Step

Production of methyl 4-acetyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3-acetamide-2-hydroxybenzoate obtained in the First Step was dissolved in N,N-dimethylformamide (42 ml), potassium carbonate (9.67 g) and 1-bromo-2-chloroethane (4.99 ml) were added, and the mixture was stirred at 50°C overnight. The reaction mixture was partitioned between ethyl acetate and water, and ethyl acetate layer was washed with water, dried
over anhydrous sodium sulfate and concentrated to obtain the
title compound (4.32 g).

Third Step

Production of

4-acetyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

To methyl

4-acetyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate
(4.32 g) obtained in the Second Step, 2 M sodium hydroxide
solution, was added and the mixture was refluxed with stirring
for 1.5 hour. The reaction mixture was cooled on ice, and
tetrahydrofuran (10 ml) solution containing acetyl chloride
(2.62 ml) was added dropwise. 2 M sodium hydroxide and acetyl
chloride were further added until the reaction finished. The
thus obtained reaction mixture was acidified with aqueous
citric acid solution and extracted with ethyl acetate. The
ethyl acetate layer was washed with saturated sodium chloride,
dried over anhydrous sodium sulfate and then concentrated.
Isopropanol was added to the concentrate, and the precipitated
solid substance was collected by filtration and dried to obtain
the title compound (2.25 g).

Fourth Step

Production of

4-acetyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo
[1,4]oxazine-8-carboxamide:

4-Acetyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxyl
ic acid (2.75 g) obtained by the similar method of the Third
Step was dissolved in tetrahydrofuran (30 ml), oxalyl chloride (1.3 ml) and N,N-dimethylformamide (one drop) were added, and the mixture was stirred at room temperature for 1 hour. To the reaction mixture was pyridine (30 ml) and 4-trifluoromethylaniline (1.9 ml) were added, and stirred at room temperature for 1 hour. The reaction mixture was partitioned between water and ethyl acetate, and the ethyl acetate layer was washed with water, dried over anhydrous sodium sulfate and then concentrated. Precipitated solid obtained by adding n-hexane and diethyl ether to the concentrate was collected by filtration and dried to obtain the title compound (2.70 g).

Fifth Step

Production of N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

Tetrahydrofuran (15 ml) was added to 4-acetyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (2.70 g) obtained in the Fourth Step, water (15 ml) and concentrated sulfuric acid (5 ml) were added, and the mixture was refluxed for 4 hours with stirring. The reaction mixture was cooled, neutralized with sodium hydroxide solution and extracted with ethyl acetate. The ethyl acetate layer was concentrated and purified by the use of silica gel column chromatography to obtain the title compound (1.94 g).

Sixth Step
Production of (2R)-2-phenylpronionate
1-(5,6-dichloropyridine-3-yl)ethyl ester:

2,3-dichloro-5-(1-hydroxy)ethylpyridine (7.58 g) was dissolved in tetrahydrofuran (65 ml), (R)-2-phenylpropion acid (5.5 ml), diisopropyl azodicarboxylate (9.5 ml) and triphenylphosphine (12.54 g) were added, and the mixture was stirred for 1 hour under ice-cooling. A mixed solution of hexane-ethyl acetate (9:1) was added to the reaction mixture, the precipitated solid was filtered off, the filtrate was concentrated and purified by the use of silica gel chromatography to obtain more polar isomer (4.54 g) and less polar isomer (5.38 g) of the title compound.

Seventh Step
Production of
4-{3-chloro-5-[1-(2R)-2-phenylpropionyl]oxyethyl}pyridine-2-yl)-N-[4-trifluoromethylphenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

The more polar isomer of (2R)-2-phenylpropionate
1-(5,6-dichloropyridine-3-yl)ethyl ester (324 mg) obtained in the Sixth Step and
N-[4-trifluoromethylphenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (322 mg) were subjected to the reaction similar to that of the First Step of Example 2.01 to obtain the title compound (80 mg).

Eighth Step
Production of
(+)-N-[4-trifluoromethylphenyl]-4-{3-chloro-5-(1-hydroxye
thyl)pyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-{3-chloro-5-[1-((2R)-2-phenylpropionyl)oxyethyl]pyridin-2-yl}-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (80 mg) obtained in the Seventh Step was dissolved in methanol (0.3 ml) and tetrahydrofuran (0.3 ml), 4 M sodium hydroxide (0.3 ml) was added, and the mixture was stirred at 60°C for 1 hour. The reaction mixture was concentrated, and the precipitated solid obtained by adding water was collected by filtration and dried to obtain the dextro-compound in the title (32 mg).

Example 2-13

Production of

(-)-N-(4-trifluoromethylphenyl)-4-{3-chloro-5-(1-hydroxyethyl)pyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

Low polar isomer of (2R)-2-phenylpronionate

1-(5,6-dichloropyridine-3-yl)ethyl ester (324 mg) obtained in the Sixth Step of Example 2-12 was used, and the similar process subsequent to the Seventh Step of Example 2-12 was performed to obtain the levo-compound in the title (115 mg).

Example 2-14

Production of

(+)-N-(4-trifluoromethoxyphenyl)-4-{3-chloro-5-(1-hydroxyethyl)pyridin-2-yl}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of

\(N\)-\(4\)-(trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

In the Fourth Step of Example 2-12, 4-trifluoromethoxyaniline (2.25 g) was used in the place of 4-trifluoromethylaniline, and the reaction similar to that of the processes 4 and 5 of Example 2-12 was performed to obtain the title compound (1.77 g).

Second Step

Production of

\((+)-N\)-(4-trifluoromethoxyphenyl)-4-\{3-chloro-5-(1-hydroxyethyl)pyridin-2-yl\}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

\(N\)-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (334 mg) obtained in the First Step and a more polar isomer of \((2R)\)-2-phenylpropionate

1-(5,6-dichloropyridine-3-yl)ethyl ester (320 mg) obtained in the Sixth Step of Example 2-12 were used, and the reaction subsequent to the Seventh Step of Example 2-12 was performed to obtain dextro-compound in the title (197 mg).

Example 2-15

Production of

\((-)-N\)-(4-trifluoromethoxyphenyl)-4-\{3-chloro-5-(1-hydroxyethyl)pyridin-2-yl\}-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

In the Second Step of Example 2-14, the more polar isomer of \((2R)\)-2-phenylpropionate
1-(5,6-dichloropyridine-3-yl)ethyl ester was replaced by the low polar isomer (320 mg), and others were performed similarly to obtain levo-compound in the title (180 mg).

Example 2-16 to Example 2-48

Compounds of Examples 2-16 to 2-48 shown in the following tables were obtained by similarly performing any methods described in the general processes A to C for producing the compound and/or the methods described in Examples 2-01 to 2-15 hereinbefore.

Example 3

N-(4-trifluoromethoxyphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-3-oxo-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of methyl 2-methoxymethoxy-3-nitrobenzoate:

Methyl 3-nitrosalicylate (5.91 g) obtained in the First Step of Example 1-001 was dissolved in N,N-dimethylformamide (60 ml), potassium carbonate (8.29 g) was added, methoxymethyl chloride (2.73 ml) was added with stirring under ice-cooling, and the mixture was stirred for 4.5 hours. The reaction mixture was concentrated and was partitioned between ethyl acetate and water. The ethyl acetate layer was washed with water, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (6.98 g).

Second Step
Production of methyl 3-amino-2-methoxymethoxybenzoate:
Methyl 2-methoxymethoxyloxy-3-nitrobenzoate (6.98 g) obtained in the First Step was subjected to hydrogenation similar to the Second Step of Example 1-001 to obtain the title compound (6.11 g).

Third Step
Production of methyl 3-(3-chloropyridin-2-yl)amino-2-methoxymethoxybenzoate:

Methyl 3- amino-2-methoxymethoxybenzoate (6.11 g) obtained in the Second Step and 2,3-dichloropyridine (4.29 g) were subjected to the reaction similar to that of the First Step of Example 2-01 to obtain the title compound (7.46 g).

Fourth Step
Production of methyl 3-(3-chloropyridin-2-yl)aminosalicylate:

Methyl 3-(3-chloropyridin-2-yl)amino-2-methoxymethoxybenzoate (7.46 g) obtained in the Third Step was dissolved in methanol (50 ml), 6 M hydrochloric acid was added, and the mixture was stirred at 40°C for 1 hour. The reaction mixture was concentrated. The solid substance precipitated by adding water was collected by filtration and dried to obtain the title compound (5.64 g).

Fifth Step
Production of 3-(3-chloropyridin-2-yl)aminosalicylic acid:

Methyl 3-(3-chloropyridin-2-ylamino)salicylate (3.08 g) obtained in the Fourth Step was dissolved in methanol (20
ml), 4 M sodium hydroxide (8.3 ml) was added, and the mixture was stirred at 70°C for 3 hours. The reaction mixture was concentrated. Aqueous citric acid solution was added to the residue, and the solid was substance precipitated collected by filtration and dried to obtain the title compound (2.73 g).

Sixth Step
Production of
3-(3-chloropyridin-2-yl)amino-N-(4-trifluoromethoxy)benzamide obtained in the previous Fifth Step:
(2.73 g) and 4-trifluoromethoxyaniline (1.46 ml) were subjected to condensation similar to the Eighth Step of Example 1-001 to obtain the title compound (2.90 g).

Seventh Step
Production of
3-[(3-chloropyridin-2-yl)-(chloroacetyl)]amino-N-(4-trifluoromethoxy)benzamide:
3-(3-chloropyridin-2-yl)amino-N-(4-trifluoromethoxy)benzamide (0.84 g) obtained in the Sixth Step was dissolved in tetrahydrofuran (10 ml), triethylamine (0.30 ml) and chloroacetyl chloride (0.175 ml) under ice-cooling were added in this order, and the mixture was stirred for 1 hour. The reaction mixture was partitioned between ethyl acetate and water. The ethyl acetate layer was washed with water, aqueous citric acid solution, aqueous sodium hydrogencarbonate and saturated sodium chloride solution, in this order, dried over
anhydrous sodium sulfate and concentrated to obtain residue containing the title compound.

Eighth Step
Production of
N-(4-trifluoromethoxyphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-3-oxo-2H-benzo[1,4]oxazine-8-carboxamide:

The residue obtained in the previous step was dissolved in N,N-dimethylformamide (10 mL), potassium carbonate (0.55 g) was added, and the mixture was stirred at 80°C for 1 hour. After concentrating the reaction solution, the solution was partitioned between ethyl acetate and water. The ethyl acetate layer was washed with water, aqueous citric acid solution, and saturated saline solution, in this order, dried over anhydrous sodium sulfate and then concentrated. The concentrated residue was purified by the use of silica gel chromatography to obtain the title compound (519 mg).

Example 4
Example 4-01
Production of
(R)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of (S)-methyl 3-(3-chloropyridin-2-yl)amino-2-(oxirane-2-yl)methyloxybenzoate:
Methyl 3-((3-chloropyridin-2-yl)aminosalicylate (6.08 g) obtained in the Fourth Step of Example 3 was dissolved in N,N-dimethylformamide (60 ml), potassium carbonate (3.01 g) and (S)-glycidyl nosylate (6.78) were added, and the mixture was stirred at room temperature for 13 hours. The reaction mixture was partitioned between diethyl ether and water. Ether layer was washed with water, aqueous sodium hydrosolcarbonate solution and saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel chromatography to obtain the title compound (7.22 g).

Second Step

Production of (R)-methyl
4-((3-chloropyridin-2-yl)-3-hydroxymethyl-3,4-dihydro-2H-benzole[1,4]oxazine-8-carboxylate:

(S)-Methyl
3-((3-chloropyridin-2-yl)amino-2-(oxirane-2-yl)methoxybenzoate (7.10 g) obtained in the First Step was dissolved in N,N-dimethylformamide (70 ml), potassium carbonate (3.66 g) was added, and the mixture was stirred at 60°C for 2 hours. The reaction mixture was partitioned between water and ethyl acetate. The ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel chromatography to obtain the colorless oily compound in the title (3.42 g).

Third Step
Production of \((R)\)-methyl 4-\((3\text{-chloropyridin-2-yl})\)-3-(\(\text{tetrahydropyran-2-yl}\)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

\((R)\)-methyl 4-\((3\text{-chloropyridin-2-yl})\)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (3.33 g) obtained in the Second Step was dissolved in chloroform (35 ml), 2,3-dihydropyran (1.0 g) and tin chloride dihydrate (225 mg) were added, and the mixture was stirred at room temperature for 17 hours. The reaction mixture was concentrated and purified by the use of silica gel chromatography to obtain the oily compound in the title (3.49 g).

Fourth Step

Production of \((R)\)-4-\((3\text{-chloropyridin-2-yl})\)-3-(\(\text{tetrahydropyran-2-yl}\)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

\((R)\)-methyl 4-\((3\text{-chloropyridin-2-yl})\)-3-(\(\text{tetrahydropyran-2-yl}\)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (3.39 g) obtained in the Third Step was dissolved in tetrahydrofuran (15 ml) and methanol (15 ml), 4 M sodium hydroxide (10 ml) was added, and stirred at 60°C for 0.5 hour. The reaction mixture was neutralized with hydrochloric acid concentrated, and partitioned between water and ethyl acetate. The ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (3.15 g).
Fifth Step

Production of

\((R)\cdot 4\cdot (3\cdot \text{chloropyridin-2-yl})\cdot 3\cdot (\text{tetrahydropyran-2-yl})\text{oxymethyl-N} \cdot (4\cdot \text{trifluoromethoxyphenyl}) \cdot 3,4\cdot \text{dihydro-2H-benzo[1,4]oxazine-8-carboxamide:}\)

\((R)\cdot 4\cdot (3\cdot \text{chloropyridin-2-yl})\cdot 3\cdot (\text{tetrahydropyran-2-yl})\text{oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (1.20 g)} \text{ obtained in the Fourth Step and 4-trifluoromethoxyaniline (525 mg) were subjected to condensation similar to the Eighth Step of Example 1-001 to obtain the white amorphous compound in the title (1.40 g).}\)

Sixth Step

Production of

\((R)\cdot 4\cdot (3\cdot \text{chloropyridin-2-yl})\cdot 3\cdot \text{hydroxymethyl-N} \cdot (4\cdot \text{trifluoromethoxyphenyl}) \cdot 3,4\cdot \text{dihydro-2H-benzo[1,4]oxazine-8-carboxamide:}\)

\((R)\cdot 4\cdot (3\cdot \text{chloropyridin-2-yl})\cdot 3\cdot (\text{tetrahydropyran-2-yl})\text{oxymethyl-N} \cdot (4\cdot \text{trifluoromethoxyphenyl}) \cdot 3,4\cdot \text{dihydro-2H-benzo[1,4]oxazine-8-carboxamide (1.33 g)} \text{ obtained in the Fifth Step was dissolved in tetrahydrofuran (15 ml), 6 M hydrochloric acid (2 ml) was added, and the mixture was stirred at room temperature for 1 hour. The reaction mixture was partitioned between water and ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride solution, in this order, dried over anhydrous sodium sulfate and then concentrated. The concentrated residue was purified by the}
use of silica gel column chromatography to obtain the title compound (930 mg).

Example 4-02

Production of

(R)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of

(R)-4-(3-chloropyridin-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(R)-4-(3-chloropyridin-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (1.63 g) obtained in the Fourth Step of Example 4-01 was dissolved in pyridine (15 ml), thionyl chloride (0.352 ml) was added, and the mixture was stirred at room temperature for 1 hour. A solution of 4-trifluoromethylaniline (0.973 ml) in pyridine (1 ml) was added and the mixture was further stirred for 1 hour. The reaction mixture was concentrated and partitioned between water and ethyl acetate. The thus obtained ethyl acetate layer was washed with aqueous citric acid solution, aqueous sodium hydrogen carbonate and saturated sodium chloride solution, in this order, dried over anhydrous sodium sulfate and then concentrated. The thus obtained residue was purified by the use of silica gel column
chromatography to obtain the white amorphous compound in the title (2.03 g).

Second Step

Production of

\[(R)\cdot 4\cdot (3\text{-chloropyridin}-2\text{-yl})\cdot 3\text{-hydroxylmethyl}\cdot N\cdot (4\text{-trifluoromethylphenyl})\cdot 3,4\text{-dihydro}\cdot 2H\cdot \text{benzo}[1,4]\text{oxazine}-8\text{-carboxamide}:\]

\[(R)\cdot 4\cdot (3\text{-chloropyridin}-2\text{-yl})\cdot 3\cdot (\text{tetrahydropyran}-2\text{-yl})\text{oxymethyl}\cdot N\cdot (4\text{-trifluoromethylphenyl})\cdot 3,4\text{-dihydro}\cdot 2H\cdot \text{benzo}[1,4]\text{oxazine}-8\text{-carboxamide} \] (1.93 g) obtained in the First Step was subjected to the reaction similar to that of the Sixth Step of Example 4-01 to obtain the title compound (1.34 g).

Example 4-03

Production of

\[(S)\cdot 4\cdot (3\text{-chloropyridin}-2\text{-yl})\cdot 3\text{-hydroxylmethyl}\cdot N\cdot (4\text{-trifluoromethoxyphenyl})\cdot 3,4\text{-dihydro}\cdot 2H\cdot \text{benzo}[1,4]\text{oxazine}-8\text{-carboxamide}:\]

First Step

Production of

\[(S)\cdot 4\cdot (3\text{-chloropyridin}-2\text{-yl})\cdot 3\cdot (\text{tetrahydropyran}-2\text{-yl})\text{oxymethyl}-3,4\text{-dihydro}\cdot 2H\cdot \text{benzo}[1,4]\text{oxazine}-8\text{-carboxylic acid}:\]

In the First Step of Example 4-01, (R)\text{-glycidyl nosylate (6.74 g) was used in the place of (S)\text{-glycidyl nosylate, and the reaction was performed in the similar manner from the First Step to the Fourth Step to obtain the title compound (3.88 g).}

Second Step
Production of
(S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(S)-4-(3-chloropyridin-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (1.88 g) obtained in the First Step was used, and the reactions similar to the Fifth Step and the Sixth Step of Example 4-01 were performed to obtain the title compound (1.126 g).

Example 4-04

Production of
(S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(S)-4-(3-chloropyridin-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (2.0 g) obtained in the First Step of Example 4-03 was used, and the reaction similar to that of the process of Example 4-02 was performed to obtain the title compound (0.872 g).

Example 4-05

Production of
(S)-4-(5-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of methyl 3-(5-chloropyridin-2-yl)amino-2-methoxymethoxybenzoate:
Methyl 3-amino-2-methoxymethoxybenzoate (4.2 g) obtained by the same method in the Second Step of Example 3 and 2,5-dichloropyridine (3.0 g) were subjected to the reaction similar to that of the First Step of Example 2-01 to obtain the title compound (1.0 g).

Second Step

Production of
(S)-4-(5-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

Methyl 3-(5-chloropyridin-2-yl)amino-2-methoxymethoxybenzoate (330 mg) obtained in the First Step was used, and others were treated by the similar way to Example 4-03 to obtain the title compound (83 mg).

Example 4-06

Production of
(S)-4-(5-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl 3-(5-chloropyridin-2-yl)amino-2-hydroxybenzoate:

In the Third Step of Example 3, 2,5-dichloropyridine (3.0 g) was used in the place of 2,3-dichloropyridine, methyl 3-amino-2-methoxymethoxybenzoate (4.20 g) was subjected to
the reaction similar to that of the Fourth Step to obtain the title compound (1.0 g).

Second Step

Production of

(S)-4-(5-chloropyridin-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid.

Methyl 3-(5-chloropyridin-2-yl)amino-2-hydroxybenzoate (1.0 g) obtained in the First Step was used in the First Step of Example 4-01 except that (R)-glycidyl nosylate (1.1 g) was used in the place of (S)-glycidyl nosylate and others were performed in a similar manner as from the First Step to the Fourth Step to obtain the title compound (830 mg).

Third Step

Production of

(S)-4-(5-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

The process similar to Example 4-02 was performed by using (S)-4-(5-chloropyridin-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (500 mg) obtained in the Second Step to obtain the title compound (104 mg).

Example 4-07

Production of

(S)-4-(5-picoline-2-yl)-3-hydroxymethyl-N-(4-trifluorome
thoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

In Example 4 - 05, 6-chloro-3-picoline was used in the place of 2,5-dichloropyridine, and was subjected to the reaction similar to that of the process to obtain the title compound (91 mg).

Example 4 - 08

Production of (S)-4-[(5-picoline-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

In Example 4 - 06, 6-chloro-3-picoline was used in the place of 2,5-dichloropyridine, and subjected to the reaction similar to that of the process to obtain the title compound (167 mg).

Example 4 - 09 to Example 4 - 56

Compounds of Examples 4 - 09 to 4 - 56 were obtained by similarly performing any methods described in the general processes A to C for producing the compound and/or the methods described in Examples 4 - 01 to 4 - 08 hereinbefore.

Example 4 - 57

Production of (R)-9-[(3-chloropyridin-2-yl)-7-hydroxy-N-(4-trifluoromethoxyphenyl)]-6,7,8,9-tetrahydro-5-oxa-9-azabenzocycloheptane-4-carboxamide:

First Step
Production of methyl 
(R)-9-(3-chloropyridin-2-yl)-7-hydro-6,7,8,9-tetrahydro-5-oxa-9-azabenzocycloheptane-4-carboxylate: 
(S)-3-(3-chloropyridin-2-yl)amino-2-(oxirane-2-yl)methylloxy benzoate (0.30 g), which was obtained by using (R)-glycidyl nosylate in the place of (S)-glycidyl nosylate in the process of Example 4-01, was dissolved in N,N-dimethylformamide (5 ml), sodium methoxide (72 mg) was added, and the mixture was stirred at room temperature for 1.5 hour. The reaction mixture was partitioned between ethyl acetate and water. The ethyl acetate layer was washed with water and dried over anhydrous sodium sulfate. Solvent was distilled off to obtain the oily substance containing the title compound.

Second Step

Production of 
(R)-9-(3-chloropyridin-2-yl)-7-hydro-6,7,8,9-tetrahydro-5-oxa-9-azabenzocycloheptane-4-carboxylic acid:

Methyl 
(R)-9-(3-chloropyridin-2-yl)-7-hydro-6,7,8,9-tetrahydro-5-oxa-9-azabenzocycloheptane-4-carboxylate (520 mg) obtained by the method similar to the previous step was dissolved in methanol (5 ml), 4 N sodium hydroxide (1.6 ml) was added, and the mixture was stirred at 65°C for 3 hours. The reaction mixture was concentrated and partitioned between water and ethyl acetate. The aqueous layer was adjusted to pH 3 by adding 6 N hydrochloric acid. The precipitated solid substance was
collected by filtration, washed with water and dried to obtain the title compound (415 mg).

Third Step

Production of

(R)·9·(3·chloropyridin-2·yl)·7·hydroxy·N··(4·trifluoromethoxyphenyl)·6,7,8,9·tetracydro·5·oxa·9·azabenzocycloheptane·4·carboxamide:

(R)·9·(3·chloropyridin-2·yl)·7·hydroxy·6,7,8,9·tetracydro·5·oxa·9·azabenzocycloheptane·4·carboxylic acid (321 mg) obtained in the Second Step was subjected to the condensation reaction similar to that of the Eighth Step of Example 1·001 to obtain the title compound (310 mg).

Example 4·58

Production of

(R)·9·(3·chloropyridin-2·yl)·7·hydroxy·N··(4·trifluoromethylphenyl)·6,7,8,9·tetracydro·5·oxa·9·azabenzocycloheptane·4·carboxamide:

In the production process of Example 4·04, when the reaction similar to that of the Sixth Step of Example 4·01 was performed, the title compound of white solid substance was obtained as the by-product.

Example 4·59

Production of

(S)·9·(3·chloropyridin-2·yl)·7·hydroxy·N··(4·trifluoromethoxyphenyl)·6,7,8,9·tetracydro·5·oxa·9·azabenzocycloheptane·4·carboxamide:
(S)-glycidyl nosylate was used in the method similar to Example 4-57 to obtain the title compound.

Example 5

Example 5-01

Production of N-[(4-tert-butylphenyl)-4-(pyrazine-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl 4-[(pyrazine-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (1.0 g) obtained in the Fifth Step of Example 1-001 and chloropyrazine (0.6 g) were used, and the coupling reaction similar to that of the Fourth Step of Example 1-002 was performed to obtain the oily compound in the title (0.84 g).

Second Step

Production of 4-[(pyrazine-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl 4-[(pyrazine-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (0.84 g) obtained in the First Step was dissolved in tetrahydrofuran (2 ml) and methanol (2 ml), 2 N sodium hydroxide (3 ml) was added, and the mixture was stirred at 60°C for 3 hours. The reaction mixture was concentrated and 1N potassium hydrogen sulfate was added. The precipitated yellowish solid
substance was collected by filtration and dried to obtain the title compound (0.40 g).

Third Step

Production of N-(4-tert-butylphenyl)-4-(pyrazine-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(pyrazine-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (150 mg) obtained in the Second Step was dissolved in tetrahydrofuran (3 ml), oxalyl chloride (0.05 ml) and N,N-dimethylformamide (one drop) were added, and the mixture was stirred at room temperature for 0.5 hour. The reaction mixture was concentrated. The concentrated residue was dissolved in tetrahydrofuran (3 ml), tert-butylaniline (74 mg) and triethylamine (0.5 ml) were added, and the mixture was stirred at room temperature for 0.5 hour. The reaction mixture was partitioned between water and ethyl acetate, and the ethyl acetate layer was dried over anhydrous sodium sulfate and then concentrated. The residue was dissolved in methanol with heating. The solution was cooled to obtain a solid substance, which was collected by filtration and dried to obtain the title compound (12 mg).

Example 5-02

Production of N-(4-chlorophenyl)-4-(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of methyl 4-(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (677 mg) obtained in the Fifth Step of Example 1-001 and 2-chloro-4-ethylpyrimidine (500 mg) were used, and the coupling reaction similar to that of the Fourth Step of Example 1-002 was performed to obtain the yellowish oily compound in the title (944 mg).

Second Step

Production of 4-(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl 4-(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (944 mg) obtained in the First Step was subjected to the reaction similar to that of the Second Step of example 5-02 to obtain white solid compound in the title (820 mg).

Third Step

Production of N-(4-chlorophenyl)-4-(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (100 mg) obtained in the Second Step and 4-chloroaniline (45 mg) were subjected to the condensation
reaction similar to that of the Eighth Step of Example 1-001 to obtain the title compound (99 mg).

Example 5-03

Production of N-(4-ethoxyphenyl)-4-(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (100 mg) obtained in the Second Step of example 5-02 and p-phenetidine (48 mg) were subjected to the condensation reaction similar to that of the Eighth Step of Example 1-001 to obtain the title compound (84 mg).

Example 5-04

Production of

4-(6-chloropyridazine-3-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl

4-(6-chloropyridazine-3-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (0.5 g) obtained in the Fifth Step of Example 1-001 and 3,6-dichloropiridazine (0.575 g) were used, and the coupling reaction similar to that of the Fourth Step of Example 1-002 was performed to obtain the title compound (0.24 g).

Second Step
Production of 4-(6-chloropyridazine-3-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl 4-(6-chloropyridazine-3-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (240 mg) obtained in the First Step was subjected to the reaction similar to that of the Second Step of example 5-02 to obtain the solid compound in the title (145 mg).

Third Step

Production of 4-(6-chloropyridazine-3-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(6-chloropyridazine-3-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (145 mg) obtained in the Second Step and 4-trifluoromethylaniline (89 mg) were subjected to the condensation reaction similar to example 5-01, and purified by the use of silica gel chromatography (n-hexane : ethylacetate = 3 : 1) to obtain the title compound (38 mg).

Example 5-05

Production of 4-(4-methylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl 4-thiocarbamide-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:
Methyl 3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (1.93 g) obtained in the Fifth Step of Example 1-001 was dissolved in tetrahydrofuran (20 ml), 9-fluorenylmethyloxycarbonyl isothiocyanate (2.95 g) was added under ice-cooling, and the mixture was stirred for 0.5 hour. Piperidine (5 ml) was added and the mixture was further stirred for 0.5 hour. The thus obtained reaction mixture was concentrated, and the residue was purified by the use of silica gel chromatography to obtain the title compound (2.27 g).

Second Step

Production of methyl 4-(4-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 4-thiocarbamide-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (1.10 g) obtained in the First Step was dissolved in methanol (15 ml), chloroacetone (0.4 ml) was added, and the mixture was refluxed for 5 hours. The reaction mixture was concentrated and partitioned between ethyl acetate and saturated sodium hydrogen carbonate solution. The ethyl acetate layer was washed with water, aqueous citric acid and saturated sodium chloride solution, in this order, dried over anhydrous sodium sulfate and concentrated to obtain the residue containing the title compound.

Third Step
Production of
4-(4-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

The residue containing methyl
4-(4-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate obtained in the Second Step was dissolved in methanol (10 ml), 4 N sodium hydroxide (3 ml) was added, and the mixture was refluxed for 0.5 hour. The reaction mixture was concentrated. The solid precipitated by adding aqueous citric acid was collected by filtration and dried to obtain the title compound (1.10 g).

Fourth Step

Production of
4-(4-methylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(4-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (0.55 g) obtained in the Third Step and 4-trifluoromethoxyaniline (0.28 ml) were subjected to the reaction similar to that of the Eighth Step of Example 1-001 and purified by the use of silica gel chromatography to obtain the title compound (0.722 g).

Example 5-06

Production of
4-(5-methylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of methyl 4-(5-methylthiazol-2-yl)-
3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

2,4,6-tris(1-chloroethyl)-1,3,5-trioxane (0.64 g) and
montmorillonite K-10 (39 mg) were heated at 110°C for 10
minutes. Toluene (10 ml) and methyl 4-thiocarbamide-3,4-
dihydro-2H-benzo[1,4]oxazine-8-carboxylate (1.17 g) obtained
in the First Step of example 5-05 were added, then
2,4,6-tris(1-chloroethyl)-1,3,5-trioxane was added
appropriately under heated to reflux until the reaction was
terminated. The reaction mixture was concentrated and
partitioned between ethyl acetate and saturated sodium
hydrogencarbonate solution. The ethyl acetate layer was
concentrated to obtain the title compound (0.68 g).

Second Step

Production of 4-(5-methylthiazol-2-yl)-
3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl 4-(5-methylthiazol-2-yl)-
3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (0.68 g)
obtained in the Second Step was dissolved in methanol (7 ml),
4 N sodium hydroxide (1.77 ml) was added, and the mixture was
refluxed for 20 minutes with stirring. The reaction mixture was
concentrated. The solid precipitated by adding aqueous citric
acid was collected by filtration and dried to obtain the title
compound (0.66 g).

Third Step
Production of
4-(5-methylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (0.66 g) obtained in the Second Step and 4-trifluoromethoxyaniline (0.28 ml) were subjected to the reaction similar to that of the Eighth Step of Example 1-001 to obtain the title compound (0.878 g).

Example 6
Example 6-01

Production of
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of methyl 2-methoxymethoxy-3-thioureid benzoate:

Methyl 3-amino-2-methoxymethoxybenzoate (2.11 g) obtained in the Second Step of Example 3 was dissolved in tetrahydrofuran (20 ml), and 9-fluorenylmethoxycarbonyl isothiocyanate (3.09 g) was added under ice-cooling, and the mixture was stirred for 3 hours. The reaction mixture was concentrated, N,N-dimethylformamide and piperidine (1 ml) were added, and the mixture was further stirred for 2 hours. The thus obtained reaction mixture was concentrated and partitioned between ethyl acetate and water. The ethyl acetate layer was washed with aqueous citric acid, saturated
sodium hydrogencarbonate solution and saturated sodium chloride solution, in this order, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (1.91 g).

Second Step

Production of methyl 3-(5-methylthiazol-2-yl)aminosalicylate:

Methyl 2-methoxymethoxy-3-thioureid benzoate (1.64 g) obtained in the First Step was subjected to the reaction similar to that of the First Step of example 5-06 to obtain the title compound (734 mg).

Third Step

Production of methyl (R)-3-(5-methylthiazol-2-yl)amino-2-(oxirane-2-yl)methyloxysterbenzoate:

Methyl 3-(5-methylthiazol-2-yl)aminosalicylate (848 mg) obtained by the similar method as in the Second Step was dissolved in N,N-dimethylformamide (8 ml), potassium carbonate (452 mg) and (R)-glycidyl nosylate (915 mg) were added, and the mixture was stirred at room temperature for 5 hours. The reaction mixture was partitioned between ethyl acetate and water. The ethyl acetate layer was washed with water and saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel chromatography (n-hexane : ethyl acetate = 3 : 1) to obtain the title compound (719 mg).

Fourth Step
Production of methyl
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl
(R)-3-(5-methylthiazol-2-yl)amino-2-(oxirane-2-yl)methyloxybenzoate (700 mg) obtained in the Third Step was dissolved in dimethyl sulfoxide (5 ml), 1,8-diazabicyclo[5.4.0]undec-7-ene (166 mg) was added, and the mixture was stirred at room temperature for 1.5 hours. The reaction mixture was partitioned between ethyl acetate and water. The ethyl acetate layer was washed with water and saturated sodium chloride solution, dried over anhydrous sodium sulfate and concentrated to obtain the colorless oily compound in the title (695 mg).

Fifth Step
Production of methyl
(S)-4-(5-methylthiazol-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (695 mg) obtained in the Fourth Step was dissolved in chloroform (7 ml), 2,3-dihydropyran (700 mg) and p-toluenesulfonic acid hydrate (453 mg) were added, and the mixture was stirred at room temperature for 7 hours. The reaction mixture was concentrated and partitioned between ethyl acetate and saturated sodium hydrogen carbonate solution. The thus
obtained ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The concentrated residue was purified by the use of silica gel chromatography (n-hexane : ethyl acetate = 3 : 1) to obtain the title compound (552 mg).

Sixth Step
Production of (S)-4-(5-methylthiazol-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl (S)-4-(5-methylthiazol-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (552 mg) was dissolved in tetrahydrofuran (2.5 ml) and methanol (2.5 ml), 4 M sodium hydroxide (1.0 ml) was added, and the mixture was stirred at 60°C for 1 hour. The reaction mixture was neutralized with diluted hydrochloric acid and extracted with ethyl acetate. The ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (563 mg).

Seventh Step
Production of (S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3-(tetrahydrofuran-2-yl)oxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(S)-4-(5-methylthiazol-2-yl)-3-(tetrahydrofuran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic
acid (250 mg) obtained in the Sixth Step and 4-trifluoromethoxyaniline (170 mg) were subjected to condensation similar to the Eighth Step of Example 1-001 to obtain the title compound (379 mg).

Eighth Step

Production of

(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-4H-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (379 mg) obtained in the Seventh Step was dissolved in tetrahydrofuran (5 ml), added 6 N hydrochloric acid (1 ml) was added and the mixture was stirred at room temperature for 1 hour. The reaction mixture was neutralized with saturated sodium hydrogen carbonate solution and extracted with ethyl acetate. The ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel column chromatography. To the purified substance was added n-hexane, and the precipitated solid substance was collected by filtration and dried to obtain the title compound (216 mg).

Example 6-02

Production of

(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-N-(4-trifluo
romethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(S)-4-(5-methylthiazol-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (260 mg) obtained in the Sixth Step of Example 6-01 was dissolved in pyridine (5 ml), thionyl chloride (0.097 ml) was added, and the mixture was stirred at room temperature for 1 hour. Subsequently, 4-trifluoromethylaniline was added, and the mixture was stirred overnight. The reaction mixture was concentrated and partitioned between ethyl acetate and 5% aqueous citric acid solution. The ethyl acetate layer was washed with water and saturated sodium chloride solution, in this order, dried over anhydrous sodium sulfate and concentrated to obtain the residue containing the title compound.

Second Step

Production of
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

The residue containing
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3-(tetrahydr
ropyran-2-yl)oxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydrro-2H-benzo[1,4]oxazine-8-carboxamide obtained in the First Step was subjected to the reaction similar to that of the Eighth Step of Example 6-01 to obtain the title compound (227 mg).

Example 6-03

Production of
(S)-N-(3,4-dichlorophenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide hydrochloride:

First Step

Production of methyl 2-methoxymethyloxy-3-(3-prop-2-yny1)thioureidobenzoate:

Methyl 3-amino-2-methoxymethyloxybenzoate (21.12 g) obtained in the Second Step of Example 3 was dissolved in ethyl acetate (100 ml), water (100 ml) and sodium hydrogen carbonate (25.2 g) were added, and thiophosgene (7.62 ml) was added dropwise with stirring under ice-cooling. After passing 0.5 hour, propargylamine (7.2 ml) was added, and the mixture was stirred at room temperature for 2 hours. The reaction mixture was partitioned and ethyl acetate layer was washed with water, dried over anhydrous sodium sulfate and concentrated to obtain the brownish oily substance (36.16 g) containing the title compound.

Second Step
Production of methyl 2-methoxymethoxy-3-(5-methylene-4,5-dihydrothiazol-2-yl)aminobenzoate:

The brownish oily substance (36.16 g) obtained in the First Step was dissolved in methanol (200 ml), p-toluenesulfonic acid (1.90 g) was added, and the mixture was refluxed for 2 hours with stirring. The reaction mixture was concentrated and partitioned between ethyl acetate and saturated aqueous sodium hydrogen carbonate. The ethyl acetate layer was washed with water and saturated sodium chloride solution in this order, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (34.42 g).

Third Step

Production of methyl 3-(5-methylthiazol-2-yl)aminosalicylate:

Methyl 2-methoxymethoxy-3-(5-methylene-4,5-dihydrothiazol-2-yl)aminobenzoate (34.42 g) obtained in the Second Step was stirred at 50°C for 30 minutes in a solution of 25% hydrogen bromide/acetic acid (60 ml). The reaction mixture was cooled, 4 N sodium hydroxide (75 ml) was added with stirring, and the precipitated solid substance was collected by filtration and dried to obtain the title compound.

Fourth Step
Production of methyl
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3-(5-methylthiazol-2-yl)aminosalicylate (3.54 g) obtained in the Third Step was subjected to the reaction similar to that of the Third Step and Fourth Step of Example 6-01 to obtain the title compound (4.29 g).

Fifth Step

Production of
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyl
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (4.29 g) obtained in the Fourth Step was dissolved in tetrahydrofuran (10 ml) and methanol (10 ml), 4 N sodium hydroxide solution (10 ml) was added, and the mixture was stirred at 60°C for 0.5 hour. The reaction mixture was neutralized, concentrated and partitioned between ethyl acetate and water. The ethyl acetate layer was washed with saturated sodium chloride solution and then concentrated. Ethyl acetate was added to the residue, the insoluble substance was filtered off and concentrated to obtain the title compound (3.53 g).

Sixth Step

Production of
(S)-3-acetoxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid hydrochloride:
(S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (2.53 g) obtained in the Fifth Step was dissolved in tetrahydrofuran (25 ml), 4-(dimethylamino)pyridine (1.01 g) was added, and the mixture was stirred at room temperature for 1 hour. Subsequently, acetic anhydride (0.779 ml) was added and the mixture was further stirred for 0.5 hour. The reaction mixture was partitioned between ethyl acetate and 5% aqueous citric acid solution. The ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The concentrated residue was dissolved in ethyl acetate (50 ml) and 4 N hydrogen chloride/ethyl acetate solution (2.5 ml) was added. The precipitated solid substance was collected by filtration and dried to obtain the title compound (2.60 g).

Seventh Step
Production of
(S)-3-acetoxymethyl-N-(3,4-dichlorophenyl)-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(S)-3-acetoxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid hydrochloride (445 mg) obtained in the Sixth Step was dissolved in pyridine (5 ml), thionyl chloride (0.168 ml) was added, and the mixture was stirred at room temperature for 1 hour. 3,4-Dichloroaniline (187 mg) was added and the mixture was further stirred for 0.5 hour. The reaction mixture was
concentrated and partitioned between ethyl acetate and 5% aqueous citric acid solution. The ethyl acetate layer was washed with water and saturated sodium chloride solution, in this order, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel chromatography (n-hexane : ethyl acetate = 2 : 1) to obtain the title compound (398 mg).

Eighth Step

Production of

(S)-N-(3,4-dichlorophenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide hydrochloride:

(S)-3-acetoxymethyl-N-(3,4-dichlorophenyl)-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide (398 mg) obtained in the Seventh Step was dissolved in tetrahydrofuran (5 ml) and methanol (5 ml), 1 N sodium hydroxide (1.25 ml) was added, and the mixture was stirred at room temperature for 0.5 hour. The reaction mixture was concentrated, and partitioned between ethyl acetate and water. The ethyl acetate layer was washed with water and saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The residue was dissolved in diethyl ether (10 ml), and 4 N hydrogen chloride/ethyl acetate solution (1 ml) was added. The precipitated solid substance was collected by filtration and dried to obtain the title compound (352 mg).

Example 6-04 to Example 6-10
Compounds of Examples 6.04 to 6.10 shown in the following tables were obtained by similarly performing any methods described in the general processes A to C for producing the compound and/or the methods described in Examples 6.01 to 6.03 hereinbefore.

Example 6.11

Production of
(S)-3-hydroxymethyl-4-(5-methyloxazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl 3-isothiocyanato-2-methoxymethoxybenzoate:

Methyl 3-amino-2-methoxymethoxybenzoate (2.0 g) obtained in the Second Step of Example 3 was dissolved in tetrahydrofuran (20 ml), and triethylamine (4 ml) was added, and thiophosgene (0.76 ml) was added dropwise under ice-coldling and stirred at room temperature for 1 hour. After adding water, the reaction mixture was concentrated and partitioned by adding ethyl acetate. The ethyl acetate layer was washed with water, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (2.4 g).

Second Step

Production of methyl 3-(5-methyloxazol-2-yl)amino-3-methoxymethoxybenzoate:

1-azidoacetone (0.94 g) was dissolved in methylene chloride (10 ml), methyl
3-isothiocyanato-2-methoxymethoxybenzoate (2.4 g) obtained in the First Step and triphenylphosphine (2.5 g) were added, and the mixture was stirred at room temperature for 3 hours. The reaction mixture was concentrated, diethyl ether was added, and the precipitated solid substance was removed off by filtration. The filtrate was concentrated. The residue was purified by the use of silica gel chromatography (n-hexane : ethyl acetate = 5 : 1) to obtain the title compound (1.82 g).

Third Step
Production of methyl 3-(5-methoxyazol-2-yl)aminosalicylate:
Methyl 3-(5-methoxyazol-2-yl)amino-3-methoxymethoxybenzoate (1.77 g) obtained in the Second Step was dissolved in tetrahydrofuran (20 ml), 6 N hydrochloric acid (3 ml) was added, and the mixture was stirred at room temperature for 1 hour. The reaction mixture was concentrated and partitioned between water and ethyl acetate. The aqueous layer was neutralized with hydrochloric acid, collected the precipitated solid substance by filtration and dried to obtain the title compound (1.4 g).

Fourth Step
Production of (S)-4-(5-methoxyazol-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:
Methyl 3-(5-methoxyazol-2-yl)aminosalicylate (1.4 g) obtained in the Third Step was subjected to the reaction similar
to that of the Third Step to Sixth Step of Example 6.01 to obtain the title compound (760 mg).

Fifth Step

Production of

(S)-3-hydroxymethyl-4-(5-methylloxazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(S)-4-(5-methylloxazol-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (200 mg) obtained in the Fourth Step was subjected to the reaction similar to that of the Seventh Step and the Eighth Step of Example 6.01 to obtain the title compound (78 mg).

Example 6.12

Production of

(S)-3-hydroxymethyl-4-(5-methylloxazol-2-yl)-N-(4-trifluoromethylenphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

(S)-4-(5-methylloxazol-2-yl)-3-(tetrahydropyran-2-yl)oxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (200 mg) obtained in the Fourth Step of Example 6.11 was subjected to the reaction similar to that of the First Step and the Second Step of Example 6.02 to obtain the title compound (80 mg).

Example 6.13

Production of

(S)-4-(4,5-dimethylthiazol-2-yl)-3-hydroxymethyl-N-(4-tri
fluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step

Production of methyl 3-(4,5-dimethylthiazol-2-yl)amino-3-methoxymethoxybenzoate:

Methyl 2-methoxymethoxy-3-thioureid benzoate (2.00 g) obtained in the First Step of Example 6-01, 3-bromo-2-butanone (1.34 g) and sodium hydrogen carbonate (746 mg) in ethanol (20 ml) was refluxed for 1.5 hour with stirring. The reaction mixture was concentrated and partitioned between water and ethyl acetate. Ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate and concentrated to obtain the title compound (2.65 g).

Second Step

Production of methyl 3-(4,5-dimethylthiazol-2-yl)aminosalicylate:

Methyl 3-(4,5-dimethylthiazol-2-yl)amino-3-methoxymethoxybenzoate (2.65 g) was dissolved in tetrahydrofuran (10 ml), 6 N hydrochloric acid (2 ml) was added, and the mixture was stirred at 60°C for 1 hour. The reaction mixture was neutralized by adding saturated sodium hydrogen carbonate solution and then concentrated. The precipitated solid substance was collected by filtration and dried to obtain the title compound (2.00 g).

Third Step
Production of 
(S)-4- (4,5-dimethylthiazol-2-yl)-3-hydroxymethyl-N-(4-tri
fluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-c
arboxamide:

Methyl 3- (4,5-dimethylthiazol-2-yl)aminosalicylate
obtained in the Second Step was subjected to the reaction
similar to those of the process Third Step to Eighth Step
of Example 6-001 to obtain the title compound.

Example 6-14

Production of 
(S)-4- (4,5-dimethylthiazol-2-yl)-3-hydroxymethyl-N-(4-tri
fluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-ca
rboxamide:

Methyl 3- (4,5-dimethylthiazol-2-yl)aminosalicylate
obtained in the Second Step of Example 6-13 was subjected to
the reaction similar to those of the process Third Step to
Sixth Step of Example 6-01 and the First Step and Second Step
of Example 6-02 to obtain the title compound.

Example 6-15

Production of 
(S)-3-hydroxymethyl-4- (5-methyl[1,3,4]thiazol-2-yl) -N-(4-
trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-
8-carboxamide:

First Step
Production of methyl
3-(acetylhydrazinothioxomethyl)amino-2-methoxymethyloxybe
nzoate:
Methyl 3-isothicyano-2-methoxymethoxybenzoate (6.12 g) obtained in the First Step of Example 6-11 was dissolved in tetrahydrofuran (100 ml), acetohydrazide (2.5 g) was added, and the mixture was refluxed overnight with stirring. The reaction mixture was partitioned between water and ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride solution, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel chromatography (n-hexane : ethyl acetate = 1 : 1) to obtain the title compound (4.97 g).

Second Step

Production of methyl 3-(5-methyl[1,3,4]thiadizol-2-yl)aminosalicylate:

Methyl 3-(acetylhydrazinothioxomethyl)amino-2-methoxymethoxybenzoate (4.97 g) obtained in the First Step was dissolved in ethanol (100 ml), and concentrated sulfuric acid (50 ml) was added under ice-coding. The mixture was further stirred at room temperature for 1 hour. The reaction mixture was neutralized and partitioned between water and ethyl acetate. The ethyl acetate layer was washed with water and saturated sodium chloride solution in this order, dried over anhydrous sodium sulfate and then concentrated. The residue was purified by the use of silica gel chromatography (n-hexane : ethyl acetate = 2 : 1) to obtain the title compound (1.11 g).

Third Step
Production of 
(S)-3-hydroxymethyl-4-((5-methyl[1,3,4]thiadiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide: 

Methyl 
3-((5-methyl[1,3,4]thiadiazol-2-yl)aminosalicylate obtained in the Second Step was subjected to the reaction similar to those of the process Third Step to Eighth Step of Example 6-001 to obtain the title compound.

Example 6-16

Production of 
(S)-3-hydroxymethyl-4-((5-methyl[1,3,4]thiadiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide: 

Methyl 
3-((5-methyl[1,3,4]thiadiazol-2-yl)aminosalicylate obtained in the Second Step of Example 6-15 was subjected to the reaction similar to those of the process Third Step to Sixth Step of Example 6-001 and the First Step and Second Step of Example 6-02 to obtain the title compound.

Example 6-17

Production of 
4-((4,5-dimethylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxamide:

First Step 
Production of methyl 
2-carbomethoxymethylsulfanyl-3-nitrobenzoate:
Methyl 2-chloro-3-nitrobenzoate (11.6 g) obtained in the First Step of Example 1-004 was dissolved in methanol (100 mL), sodium hydrogen carbonate (6.83 g) and mercaptoacetic acid (2.64 mL) were added, and the mixture was refluxed with stirring for 16 hours. The reaction solution was left to cool, poured into 2 M hydrochloric acid (100 mL)-ethyl acetate (100 mL) with stirring under ice-cooling, and partitioned. The ethyl acetate layer was washed with a saturated sodium chloride solution, dried over anhydrous sodium sulfate, and concentrated. The residue was purified by silica gel chromatography (chloroform : methanol : acetic acid = 19 : 0.9 : 0.1) to obtain the title compound (4.84 g) as a pale orange solid.

Second Step

Production of methyl 3-oxo-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylate:

Ammonium chloride (379 mg) was dissolved in water (10 mL), and reduced iron (3.66 g) was added while heating to 85°C and stirring. A solution of methyl 2-carbomethoxymethylsulfanyl-3-nitrobenzoate obtained in the above step (4.81 g) in N,N-dimethylformamide (20 mL) was added dropwise thereto over 15 minutes, and the mixture was heated and stirred at 85°C for a further one hour. The reaction solution was left to cool, water, ethyl acetate and tetrahydrofuran were added. After removing the insoluble substance, the mixture was partitioned. The organic layer was dried over anhydrous sodium sulfate and concentrated. The
pale gray solid precipitated with diisopropyl ether was collected by filtration and dried to obtain the title compound (2.99 g).

Third Step
Production of methyl
3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylate:
Methyl
3-oxo-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylate obtained in the above step (2.99 g) was subjected to the same reaction as in the Fifth Step of Example 1-001 to obtain the title compound (2.53 g) as an yellow solid.

Fourth Step
Production of methyl
4-thiocarbamide-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylate:
The same reaction as in the First Step of Example 5-05 was performed for methyl
3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylate obtained in the above step to obtain the title compound (1.44 g) as a white solid.

Fifth Step
Production of methyl
4-((4,5-dimethylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylate:
Methyl
4-thiocarbamide-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylate obtained in the above step (1.44 g), 3-bromo-2-butanone
(972 mg) and sodium hydrogencarbonate (541 mg) were refluxed with stirring in methanol (15 mL) and tetrahydrofuran (10 mL) for 17 hours. The reaction solution was concentrated and partitioned between water and ethyl acetate. The resulting ethyl acetate layer was washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate, and concentrated to obtain the title compound (1.91 g).

Sixth Step

Production of

4-((4,5-dimethyl-thiazol-2-yl)3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylic acid:

Methyl

4-((4,5-dimethyl-thiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxylate obtained in the above step (1.91 g) was dissolved in methanol (5 mL) and tetrahydrofuran (5 mL), 4 N sodium hydroxide (2.0 mL) was added, and the mixture was refluxed with stirring for one hour. The reaction solution was concentrated, and neutralized by the addition of 2 N hydrochloric acid under ice-cooling. The precipitated solid was filtered off and dried to obtain the title compound (1.41 g).

Seventh Step

Production of

4-((4,5-dimethylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxamide:

The same reaction as in the Eighth Step of Example 1-001 was performed for
4-\{(4,5\text{-dimethyl\text{-}thiazol\text{-}2\text{-}yl})\text{-}3,4\text{-dihydro\text{-}2H\text{-}benzo[1,4]thiazine\text{-}8\text{-}carboxylic acid obtained in the above step (1.40 g) and 4\text{-}trifluoromethoxyaniline (809 mg) to obtain the title compound (1.03 g) as a white solid.

Example 6-18

Production of

4\text{-}(4,5\text{-dimethylthiazol\text{-}2\text{-}yl})\text{-}N\text{-}(4\text{-}trifluoromethoxyphenyl)\text{-}1\text{-}oxo\text{-}3,4\text{-tetrahydro\text{-}benzo[1,4]thiazine\text{-}8\text{-}carboxamide:}

4\text{-}(4,5\text{-dimethylthiazol\text{-}2\text{-}yl})\text{-}N\text{-}(4\text{-}trifluoromethoxyphenyl)\text{-}3,4\text{-dihydro\text{-}2H\text{-}benzo[1,4]thiazine\text{-}8\text{-}carboxamide obtained in Example 6-17 (500 mg) was suspended in chloroform (15 mL), metachloroperbenzoic acid (247 mg) was added, and the mixture was stirred at room temperature for two hours. A saturated sodium hydrogen carbonate solution was added to the reaction solution and partitioned. The chloroform layer was washed with a saturated sodium chloride solution, dried over anhydrous sodium sulfate and concentrated. The residue was purified by silica gel chromatography (chloroform : methanol = 9 : 1), and the white solid precipitated with diisopropyl ether was filtered off and dried to obtain the title compound (290 mg).

Example 7

Example 7-01

Production of

N\text{-}(4\text{-tert\text{-}butylphenyl})\text{-}4\text{-}(4\text{-methoxyphenyl})\text{-}3,4\text{-dihydro\text{-}2H\text{-}benzo[1,4]oxazine\text{-}8\text{-carboxamide:}

First Step
Production of methyl 4-((4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate:

Methyl 3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (1.0 g) obtained in the Fifth Step of Example 1-001 and 4-bromoanisole (1.0 g) were used, and the coupling reaction similar to that of the Fourth Step of Example 1-002 was performed to obtain the oily compound in the title (0.58 g).

Second Step
Production of 4-((4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid:

Methyldimethyl 4-((4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylate (0.58 g) obtained in the First Step was dissolved in tetrahydrofuran (10 ml) and methanol (10 ml), 2 N sodium hydroxide solution (5 ml) was added, and the mixture was stirred at 60°C for 1.5 hour. After concentrating the reaction mixture, yellow solid substance precipitated by adding 1 N potassium hydrogen sulfate was collected by filtration and dried to obtain the title compound (0.552 g).

Third Step
Production of N-(4-tert-butylphenyl)-4-((4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

4-((4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxylic acid (57 mg) obtained in the Second Step was
dissolved in tetrahydrofuran (5 ml), oxalyl chloride (0.03 ml) and N,N-dimethylformamide (one drop) were added, and the mixture was stirred at room temperature for 0.5 hour, then the reaction mixture was concentrated. The concentrated residue was dissolved in tetrahydrofuran (5 ml), tert-butylaniline (30 mg) and triethylamine (0.5 ml) were added, and stirred at room temperature for 1 hour. The reaction mixture was partitioned between water and ethyl acetate. The ethyl acetate layer was dried over anhydrous sodium sulfate and then concentrated. Precipitated solid obtained by adding hexane to the residue was collected by filtration, and dried to obtain the title compound (48 mg).

Example 7-02

Production of

\[ \text{N-}(4\text{-isobutyloxyphenyl})\text{-}4\text{-}(4\text{-methoxyphenyl})\text{-}3,4\text{-dihydro}\text{-}2\text{H\text{-}benzo[1,4]oxazine\text{-}8\text{-carboxamide:}} \]

\[ 4\text{-}(4\text{-methoxyphenyl})\text{-}3,4\text{-dihydro}\text{-}2\text{H\text{-}benzo[1,4]oxazine\text{-}8\text{-carboxylic acid (57 mg) obtained in the Second Step of}} \]

Example 7-01 and 4-isobutyloxyaniline (40 mg) were subjected to the reaction similar to that of the Third Step of example 7-01 to obtain the title compound (31 mg).

Example 7-03

Production of

\[ \text{N-}(4\text{-chlorophenyl})\text{-}4\text{-}(4\text{-methoxyphenyl})\text{-}3,4\text{-dihydro}\text{-}2\text{H\text{-}benzo[1,4]oxazine\text{-}8\text{-carboxamide:}} \]

\[ 4\text{-}(4\text{-methoxyphenyl})\text{-}3,4\text{-dihydro}\text{-}2\text{H\text{-}benzo[1,4]oxazine\text{-}8\text{-carboxylic acid (57 mg) obtained in the Second Step of}} \]
Example 7-01 and 4-chloroaniline (40 mg) were subjected to the reaction similar to that of the Third Step of example 7-01 to obtain the title compound (29 mg).

Example 7-04

Production of
(S)-4-(2-chlorophenyl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide:

First Step
Production of methyl 3-(2-chlorophenyl)amino-2-methoxymethoxybenzoate:

Methyl 3-amino-2-methoxymethoxybenzoate (3.5 g) obtained in the Second Step of Example 3 and 2-iodochlorobenzene (2.62 ml) were subjected to the reaction similar to that of the First Step of Example 2-01 to obtain the title compound (2.33 g).

Second Step
Production of methyl 3-(2-chlorophenyl)aminosalicylate methyl:

Methyl 3-(2-chlorophenyl)amino-2-methoxymethoxybenzoate (2.33 g) obtained in the First Step was subjected to the reaction similar to that of the Fourth Step of Example 3 to obtain the title compound (1.75 g).

Third Step
Production of
(S)-4-(2-chlorophenyl)-3-hydroxymethyl-N-(4-trifluorometh
oxypyhenyl)-3,4-dihydro-2H-benza[1,4]oxazine-8-carboxamide

Methyl 3-(2-chlorophenyl)aminosalicylate methyl
obtained in the Second Step was subjected to the reaction of
the Third Step below in Example 6-01 to obtain the title
compound.
Example 7-05 to Example 7-11

Compounds of Examples 7-05 to 7-11 hereinbelow were
obtained by similarly performing any methods described in the
general processes A to C for producing the compound and/or
the methods described in Examples 4-01 to 4-59 hereinbefore
and a method described in example 7-04.

Chemical structures, molecular weights thereof and NMR
data of compounds obtained in Examples 1-001 to 7-11 are shown
in Table 1 to Table 50.
[Table 1]
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<th>Ex. No.</th>
<th>Chemical Compounds</th>
<th>NMR</th>
</tr>
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<td>1-001</td>
<td>(400 MHz, CHLOROFORM-D) 1.33 (s, 9 H) 3.88 - 3.97 (m, 2 H) 4.57 - 4.65 (m, 2 H) 6.66 (dd, J=8.00, 1.28 Hz, 1 H) 6.90 (t, J=7.88 Hz, 1 H) 7.11 (dd, J=7.98, 4.97 Hz, 1 H) 7.38 (d, J=8.58 Hz, 2 H) 7.60 (d, J=8.58 Hz, 2 H) 7.77 - 7.82 (m, 2 H) 8.37 (dd, J=4.75, 1.28 Hz, 1 H) 9.57 (s, 1 H)</td>
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<td>1-002</td>
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<td>1-003</td>
<td>(400 MHz, DMSO-D6) 1.23 (s, J=8.49 Hz, 3 H) 1.29 (s, 9 H) 4.67 - 4.75 (m, 1 H) 4.62 - 4.75 (m, 2 H) 6.38 (dd, J=8.12, 1.62 Hz, 1 H) 7.29 (t, J=7.76 Hz, 1 H) 7.34 - 7.44 (m, 3 H) 7.64 - 7.71 (m, 2 H) 8.07 (dd, J=7.58, 1.62 Hz, 1 H) 8.45 (dd, J=4.64, 1.62 Hz, 1 H) 10.10 (s, 1 H)</td>
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<td>1-006</td>
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<td>1-008</td>
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### Table 2

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<td><img src="image1" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-D6) 2.28 (s, 3 H) 3.77 - 3.84 (m, 2 H) 4.40 - 4.48 (m, 2 H) 5.60 (dd, J=8.07, 1.47 Hz, 1 H) 6.79 (t, J=7.70 Hz, 1 H) 7.07 - 7.10 (m, 1 H) 7.14 (d, J=8.44 Hz, 2 H) 7.31 (dd, J=8.07, 4.77 Hz, 1 H) 7.63 (dd, J=8.44 Hz, 2 H) 8.05 (dd, J=8.07, 1.47 Hz, 1 H) 8.43 (dd, J=4.77, 1.47 Hz, 1 H) 10.06 (s, 1 H)</td>
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<tr>
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<td>(300 MHz, DMSO-D6) 0.89 (d, J=6.60 Hz, 3 H) 0.93 - 1.11 (m, 2 H) 1.19 - 1.40 (m, 3 H) 1.54 - 1.76 (m, 2 H) 1.81 - 1.93 (m, 2 H) 3.60 - 3.81 (m, 3 H) 3.85 - 3.95 (m, 1 H) 6.45 (dd, J=7.89, 1.65 Hz, 1 H) 6.72 - 6.77 (m, 1 H) 7.10 (dd, J=7.70, 1.47 Hz, 1 H) 7.28 (dd, J=7.89, 1.65 Hz, 1 H) 8.79 (d, J=7.70 Hz, 1 H) 8.92 (dd, J=7.89, 1.65 Hz, 1 H) 8.43 (dd, J=4.59, 1.65 Hz, 1 H)</td>
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<td>Ex. No.</td>
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<td>(300 MHz, DMSO-D6) 3.81 (m, 2 H) 4.42 (m, 2 H) 6.52 (dd, J=8.10, 1.50 Hz, 1 H) 6.83 (l, J=7.90 Hz, 1 H) 7.08 (dd, J=7.30, 1.40 Hz, 1 H) 7.33 (dd, J=4.80, 8.10 Hz, 1 H) 7.35 (dd, J=8.80 Hz, 2 H) 7.86 (dd, J=8.80 Hz, 2 H) 8.06 (dd, J=1.50, 8.30 Hz, 1 H) 8.45 (dd, J=1.50, 4.80 Hz, 1 H) 10.36 (s, 1 H)</td>
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<td>(300 MHz, DMSO-D6) 3.74 (m, 3 H) 3.80 (m, 2 H) 4.43 (m, 2 H) 6.51 (dd, J=8.12, 1.39 Hz, 1 H) 6.80 (l, J=7.89 Hz, 1 H) 6.91 (dd, J=9.17 Hz, 2 H) 7.08 (dd, J=7.70, 1.47 Hz, 1 H) 7.31 (dd, J=7.89, 4.58 Hz, 1 H) 7.86 (dd, J=8.17 Hz, 2 H) 8.05 (dd, J=7.89, 1.65 Hz, 1 H) 8.43 (dd, J=4.89, 1.65 Hz, 1 H) 10.01 (s, 1 H)</td>
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### Table 4

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<td>(400 MHz, DMSO-D6): 1.42 (s, 6H) 3.78 - 3.85 (m, 2H) 4.37 - 4.48 (m, 2H) 4.98 (s, 1H) 6.49 - 6.52 (m, 1H) 6.78 - 8.82 (m, 1H) 7.07 - 7.09 (m, 1H) 7.30 - 7.33 (m, 1H) 7.37 - 7.46 (m, 2H) 7.60 - 7.71 (m, 2H) 8.04 - 8.06 (m, 1H) 8.42 - 8.44 (m, 1H) 10.06 (s, 1H)</td>
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<td>(300 MHz, DMSO-d6) 3.81 (m, 2 H), 4.44 (m, 2 H), 6.53 (dd, J=8.10, 1.50 Hz, 1 H), 6.82 (t, J=7.90 Hz, 1 H), 7.08 (dd, J=7.30, 1.40 Hz, 1 H), 7.33 (dd, J=7.30, 1.30 Hz, 1 H), 8.01 - 8.07 (m, 2 H), 8.26 (m, 1 H), 8.40 (m, 1 H) 10.50 (s, 1 H)</td>
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<td><img src="image" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) 1.26 (d, J=8.94 Hz, 6 H), 3.81 (m, 2 H), 4.43 (m, 2 H), 4.78 (dt, J=12.00, 5.86 Hz, 1 H), 6.53 (dd, J=8.10, 1.50 Hz, 1 H), 6.80 (t, J=7.90 Hz, 1 H), 7.08 (dd, J=7.30, 1.40 Hz, 1 H), 7.33 (dd, J=7.30, 1.30 Hz, 1 H), 7.94 - 8.14 (m, 3 H), 8.40 (m, 1 H), 10.34 (s, 1 H)</td>
</tr>
<tr>
<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR (300 MHz, DMSO-D6)</td>
</tr>
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<td>56</td>
<td><img src="image1" alt="Chemical Structure" /></td>
<td>3.81 (m, 3H), 4.43 (m, 2H) 4.78 (dt, J=12.00, 5.86 Hz, 1H), 5.60 (dd, J=10.1, 15.0 Hz, 1H), 6.50 (dt, J=7.50 Hz, 1H), 7.08 (dd, J=7.30, 1.40 Hz, 1H), 7.30 (dd, J=4.80, 8.10 Hz, 1H), 7.88 (dd, J=8.80 Hz, 2H), 8.03 - 8.06 (m, 2H), 8.26 (m, 1H), 8.44 (m, 1H), 10.25 (s, 1H)</td>
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<tr>
<td>57</td>
<td><img src="image2" alt="Chemical Structure" /></td>
<td>0.93 - 1.00 (d, J=6.62 Hz, 8H), 2.01 (dd, J=13.33, 6.61 Hz, 1H), 3.72 - 3.81 (m, 2H), 3.84 (d, J=8.26 Hz, 2H), 4.34 - 4.45 (m, 2H), 6.48 (dd, J=8.12, 1.39 Hz, 1H), 6.77 (t, J=7.88 Hz, 1H), 7.05 (dd, J=7.54, 1.51 Hz, 1H), 7.20 (d, J=9.04 Hz, 1H), 7.26 (dd, J=8.00, 4.75 Hz, 1H), 7.30 (dd, J=8.04, 2.55 Hz, 1H), 8.02 (dd, J=7.88, 1.62 Hz, 1H), 8.08 (d, J=2.78 Hz, 1H), 8.40 (dd, J=4.84, 1.62 Hz, 1H), 10.22 (s, 1H)</td>
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<td>58</td>
<td><img src="image3" alt="Chemical Structure" /></td>
<td>3.85 - 4.08 (m, 2H), 4.29 - 4.38 (m, 2H), 5.93 (t, J=7.88 Hz, 2H), 7.13 (s, 1H), 7.21 - 7.28 (m, 1H), 7.29 - 7.35 (m, 2H), 7.46 (dd, J=8.12, 1.39 Hz, 1H), 7.56 - 7.66 (m, 2H), 8.16 (d, J=5.33 Hz, 1H), 10.03 (s, 1H)</td>
</tr>
<tr>
<td>59</td>
<td><img src="image4" alt="Chemical Structure" /></td>
<td>2.37 (s, 3H), 3.97 - 4.07 (m, 2H), 4.24 - 4.34 (m, 2H), 6.76 (d, J=7.42 Hz, 1H), 6.87 - 6.93 (m, 1H), 6.97 (d, J=8.35 Hz, 1H), 7.17 (dd, J=7.42, 1.62 Hz, 1H), 7.28 - 7.35 (m, 2H), 7.43 (dd, J=8.12, 1.62 Hz, 1H), 7.53 (dd, J=8.35, 7.42 Hz, 1H), 7.58 - 7.65 (m, 2H), 10.03 (s, 1H)</td>
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<td>60</td>
<td><img src="image5" alt="Chemical Structure" /></td>
<td>2.22 (s, 3H), 3.97 - 4.01 (m, 2H), 4.30 - 4.34 (m, 2H), 6.87 - 6.93 (m, 1H), 7.11 - 7.20 (m, 2H), 7.34 (dd, J=7.89 Hz, 2H), 7.41 (dd, J=8.23, 1.47 Hz, 1H), 7.52 (dd, J=8.76, 2.20 Hz, 1H), 7.64 (d, J=8.79 Hz, 2H), 8.15 (d, J=2.55 Hz, 1H), 10.04 (s, 1H)</td>
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<td><img src="image6" alt="Chemical Structure" /></td>
<td>5.128 (s, 9H), 3.80 (t, J=4.4 Hz, 2H), 4.38 (t, J=4.4 Hz, 2H), 5.63 (t, J=7.9 Hz, 1H), 6.93 (dd, J=6.1, 1.5 Hz, 1H), 7.07 (dd, J=7.5, 1.7 Hz, 1H), 7.35 (d, J=4.2 Hz, 2H), 7.42 (dt, J=8.8, 2.5 Hz, 1H), 7.53 - 7.71 (m, 3H), 8.32 (dd, J=4.8, 1.5 Hz, 1H), 8.55 (d, J=2.6 Hz, 1H), 10.04 (s, 1H)</td>
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<tr>
<td>62</td>
<td><img src="image7" alt="Chemical Structure" /></td>
<td>3.970 (d, J=4.4 Hz, 2H), 4.31 (t, J=4.4 Hz, 2H), 6.60 (d, J=5.15 Hz, 1H), 7.15 (d, J=2.55 Hz, 1H), 7.44 (dd, J=8.8, 1.4 Hz, 1H), 7.53 (dd, J=8.8, 2.5 Hz, 1H), 7.80 (d, J=8.8, 2H), 7.92 (d, J=8.8 Hz, 2H), 8.15 (d, J=2.55 Hz, 1H), 10.58 (s, 1H)</td>
</tr>
<tr>
<td>Ex. No.</td>
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</tr>
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<td>-----</td>
</tr>
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<td>63 1-063</td>
<td><img src="image1" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) δ: 2.25 (s, 3H), 3.99 (t, J = 4.6 Hz, 2H), 4.31 (t, J = 4.4 Hz, 2H), 5.34 (br s, 2H), 6.81-6.92 (m, 2H), 7.11-7.14 (m, 3H), 7.39 (dd, J = 7.7, 1.7 Hz, 2H), 7.52 (dt, J = 9.5, 1.3 Hz, 1H), 8.15 (t, J = 1.1 Hz, 1H), 9.97 (s, 1H).</td>
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<td>64 1-064</td>
<td><img src="image2" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) δ: 2.08 (s, 3H), 2.23 (s, 3H), 3.98 (t, J = 4.4 Hz, 2H), 4.31 (t, J = 4.4 Hz, 2H), 6.90 (t, J = 7.7 Hz, 1H), 7.15-7.18 (m, 2H), 7.42 (dd, J = 8.3, 1.7 Hz, 2H), 7.53-7.59 (m, 2H), 8.03-8.15 (m, 2H), 9.50 (s, 1H), 10.27 (s, 1H).</td>
</tr>
<tr>
<td>65 1-065</td>
<td><img src="image3" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.70-1.73 (m, 2H), 1.93-1.97 (m, 2H), 2.24 (s, 3H), 2.71-2.74 (m, 1H), 3.24-3.28 (m, 2H), 3.99 (t, J = 4.2 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 6.97 (t, J = 7.9 Hz, 1H), 7.04 (t, J = 10.0 Hz, 1H), 7.17-7.19 (m, 2H), 7.38-7.44 (m, 2H), 7.57-7.69 (m, 2H), 8.16 (s, 1H), 10.14 (s, 1H).</td>
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<tr>
<td>66 1-066</td>
<td><img src="image4" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.75-1.78 (m, 4H), 2.22-2.25 (m, 4H), 2.58 (d, J = 4.4 Hz, 3H), 2.70-2.72 (m, 2H), 3.31-3.35 (m, 2H), 4.00 (t, J = 4.2 Hz, 2H), 4.54 (t, J = 4.2 Hz, 2H), 6.52 (t, J = 7.9 Hz, 1H), 7.07 (t, J = 9.2 Hz, 1H), 7.50 (d, J = 7.7 Hz, 2H), 7.39-7.44 (m, 2H), 7.63-7.72 (m, 3H), 8.14-8.18 (m, 1H), 10.15 (s, 1H).</td>
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<tr>
<td>67 1-067</td>
<td><img src="image5" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.73 (s, 6H), 2.22-2.25 (m, 3H), 2.60-2.63 (m, 4H), 3.04-3.05 (m, 3H), 3.35-3.45 (m, 2H), 4.01 (t, J = 3.7 Hz, 2H), 4.34 (t, J = 4.2 Hz, 2H), 6.03 (t, J = 7.9 Hz, 2H), 7.09-7.12 (m, 1H), 7.19-7.21 (m, 2H), 7.42-7.45 (m, 3H), 7.66 (t, J = 13.9 Hz, 1H), 8.15-8.17 (m, 1H), 10.18 (s, 1H).</td>
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<td>68 1-068</td>
<td><img src="image6" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.13 (t, J = 7.0 Hz, 3H), 1.56-1.60 (m, 2H), 1.93-1.96 (m, 2H), 2.23 (s, 3H), 2.71-2.79 (m, 2H), 3.17-3.20 (m, 2H), 3.40-3.51 (m, 3H), 3.96 (t, J = 4.4 Hz, 2H), 4.32 (t, J = 4.2 Hz, 2H), 6.90 (t, J = 7.7 Hz, 1H), 7.00 (q, J = 8.4 Hz, 1H), 7.15 (dt, J = 10.1, 4.5 Hz, 2H), 7.40 (dt, J = 13.1, 4.8 Hz, 2H), 7.52 (dd, J = 8.6, 2.0 Hz, 1H), 7.65 (dd, J = 15.0, 2.2 Hz, 1H), 8.15 (d, J = 2.6 Hz, 1H), 10.12 (s, 1H).</td>
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<tr>
<td>69 1-069</td>
<td><img src="image7" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.09 (q, J = 6.2 Hz, 6H), 1.52-1.56 (m, 2H), 1.86-1.91 (m, 2H), 2.23 (s, 3H), 2.74-2.77 (m, 2H), 3.17-3.19 (m, 2H), 3.47-3.53 (m, 3H), 3.59-3.61 (m, 1H), 3.70-3.72 (m, 1H), 3.98 (t, J = 4.4 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 6.60 (t, J = 7.9 Hz, 1H), 7.01 (t, J = 9.5 Hz, 1H), 7.14 (dt, J = 10.1, 4.5 Hz, 2H), 7.30 (dd, J = 8.4, 1.8 Hz, 1H), 7.64 (dd, J = 14.9, 2.4 Hz, 8H), 8.15 (d, J = 2.2 Hz, 1H), 10.11 (s, 1H).</td>
</tr>
<tr>
<td>Ex. No.</td>
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</tr>
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</tr>
<tr>
<td>70</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(500 MHz, DMSO-d6) δ: 1.21-1.30 (m, 1H), 1.51-1.63 (m, 1H), 1.78-1.79 (m, 1H), 1.96-2.02 (m, 1H), 2.23 (s, 3H), 2.53-2.68 (m, 1H), 3.08-3.12 (m, 1H), 3.32-3.38 (m, 6H), 3.99 (t, J = 4.4 Hz, 2H), 4.32 (t, J = 4.2 Hz, 2H), 6.30 (t, J = 7.9 Hz, 1H), 7.02 (t, J = 9.4 Hz, 1H), 7.15 (dt, J = 10.1, 4.5 Hz, 2H), 7.40 (dd, J = 12.3, 4.5 Hz, 2H), 7.52 (dd, J = 8.4, 2.0 Hz, 1H), 7.65 (dd, J = 15.0, 2.2 Hz, 1H), 8.16 (d, J = 2.6 Hz, 1H), 10.12 (s, 1H).</td>
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<tr>
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<td><img src="image" alt="Chemical Structure" /></td>
<td>(500 MHz, DMSO-d6) δ: 1.31-1.36 (m, 2H), 1.70-1.73 (m, 2H), 2.23 (s, 3H), 2.60-2.63 (m, 2H), 3.23-3.24 (m, 5H), 3.38 (t, J = 4.4 Hz, 2H), 4.32 (t, J = 4.2 Hz, 2H), 6.30 (t, J = 7.9 Hz, 1H), 7.01 (t, J = 9.2 Hz, 1H), 7.14 (t, J = 8.2 Hz, 2H), 7.40 (dd, J = 13.4, 5.0 Hz, 2H), 7.52 (dd, J = 8.6, 2.4 Hz, 1H), 7.64 (dd, J = 15.0, 2.2 Hz, 1H), 8.14 (s, 1H), 10.11 (s, 1H).</td>
</tr>
<tr>
<td>72</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 1.93-2.04 (m, 2H), 2.24 (s, 3H), 3.24-3.31 (m, 5H), 3.51 (dq, J = 10.8, 2.6 Hz, 1H), 4.02 (dd, J = 7.7, 4.9 Hz, 2H), 4.36 (t, J = 4.4 Hz, 2H), 6.73 (t, J = 9.7 Hz, 1H), 6.93 (t, J = 7.9 Hz, 1H), 7.26-7.31 (m, 3H), 7.42 (dd, J = 8.1, 1.6 Hz, 1H), 7.50-7.62 (m, 1H), 7.74-7.75 (m, 1H), 8.17 (s, 1H), 10.03 (s, 1H).</td>
</tr>
<tr>
<td>73</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 1.60 (t, J = 7.0 Hz, 6H), 1.95-2.08 (m, 1H), 2.23 (s, 3H), 3.77-3.82 (m, 5H), 3.97-4.02 (m, 2H), 4.30-4.35 (m, 2H), 6.91 (t, J = 7.9 Hz, 1H), 7.09-7.20 (m, 3H), 7.42 (dd, J = 7.9 Hz, 1H), 7.53 (d, J = 8.8 Hz, 1H), 7.80-7.84 (m, 1H), 8.07-8.10 (m, 1H), 8.15 (m, 1H), 10.10 (s, 1H).</td>
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<td>74</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 0.98 (t, J = 8.1 Hz, 6H), 1.98-2.05 (m, 1H), 2.23 (s, 3H), 3.79 (d, J = 6.5 Hz, 2H), 3.99 (t, J = 4.4 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 6.91 (t, J = 7.9 Hz, 1H), 7.08 (d, J = 9.3 Hz, 1H), 7.14 (d, J = 8.3 Hz, 1H), 7.19 (dd, J = 7.9, 1.4 Hz, 1H), 7.42 (dd, J = 7.9, 1.4 Hz, 1H), 7.53 (dd, J = 8.3, 2.3 Hz, 1H), 7.81 (dd, J = 8.8, 2.8 Hz, 1H), 8.02 (d, J = 2.8 Hz, 1H), 8.15 (m, 1H), 10.09 (s, 1H), 12.59 (s, 1H).</td>
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<td>75</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 1.01 (d, J = 6.6 Hz, 6H), 2.06-2.15 (m, 1H), 2.23 (s, 3H), 3.90 (d, J = 6.6 Hz, 2H), 3.99 (t, J = 4.4 Hz, 2H), 4.33 (t, J = 4.2 Hz, 2H), 6.91 (t, J = 7.9 Hz, 1H), 7.12-7.13 (m, 2H), 7.19 (dd, J = 7.7, 3.9 Hz, 1H), 7.41 (dd, J = 8.3, 1.7 Hz, 1H), 7.52 (dd, J = 8.8, 2.2 Hz, 1H), 7.58 (br s, 1H), 7.60 (br s, 1H), 7.85 (dd, J = 9.0, 2.8 Hz, 1H), 8.13 (d, J = 2.9 Hz, 1H), 8.15 (br s, 1H), 10.08 (s, 1H).</td>
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<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.01 (d, J = 6.6 Hz, 6H), 2.06-2.14 (m, 1H), 2.23 (s, 3H), 2.82 (d, J = 4.8 Hz, 3H), 3.87 (d, J = 8.8 Hz, 2H), 3.99 (t, J = 4.2 Hz, 2H), 4.32 (t, J = 4.2 Hz, 2H), 6.90 (d, J = 7.9 Hz, 1H), 7.09-7.20 (m, 3H), 7.41 (d, J = 6.0 Hz, 1H), 7.52 (dd, J = 8.4, 2.8 Hz, 1H), 8.03 (dd, J = 9.0, 2.8 Hz, 1H), 8.00-8.07 (m, 2H), 8.15 (br s, 1H), 10.09 (s, 1H).</td>
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<tr>
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<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 0.85 (d, J = 6.6 Hz, 6H), 1.90-2.04 (m, 1H), 2.23 (s, 3H), 2.79 (s, 3H), 2.98 (s, 3H), 3.76 (d, J = 6.2 Hz, 2H), 3.99 (t, J = 4.2 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 6.90 (t, J = 7.7 Hz, 1H), 7.03 (d, J = 8.8 Hz, 1H), 7.13 (d, J = 8.8 Hz, 1H), 7.18 (dd, J = 7.3, 1.5 Hz, 1H), 7.41 (dd, J = 8.1, 1.5 Hz, 1H), 7.52 (dd, J = 8.6, 2.0 Hz, 1H), 7.57 (d, J = 2.6 Hz, 1H), 7.69 (dd, J = 8.0, 2.8 Hz, 1H), 8.15 (d, J = 2.6 Hz, 1H), 10.05 (s, 1H).</td>
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</table>
### Table 11

<table>
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<th>Chemical Compounds</th>
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</tr>
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<tr>
<td>78 1-078</td>
<td><img src="image1" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 2.23 (s, 3H), 2.55 (s, 3H), 3.99 (t, J = 4.4 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 6.92 (t, J = 7.9 Hz, 1H), 7.14 (d, J = 8.8 Hz, 1H), 7.18 (dd, J = 7.9, 1.4 Hz, 1H), 7.44 (dd, J = 8.1, 1.6 Hz, 1H), 7.53 (dd, J = 8.6, 2.6 Hz, 1H), 7.87 (d, J = 8.6 Hz, 2H), 7.96 (d, J = 8.8 Hz, 2H), 8.15 (s, 1H), 10.48 (s, 1H).</td>
</tr>
<tr>
<td>79 1-079</td>
<td><img src="image2" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 1.42 (s, 6H), 2.23 (s, 3H), 4.00 (t, J = 4.4 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 6.91 (t, J = 7.9 Hz, 1H). 7.13 (d, J = 8.4 Hz, 1H), 7.19 (dd, J = 7.7, 1.5 Hz, 1H), 7.39-7.43 (m, 3H), 7.52 (dd, J = 9.0, 2.4 Hz, 1H), 7.64 (d, J = 8.4 Hz, 2H), 8.15 (s, 1H), 10.04 (s, 1H).</td>
</tr>
<tr>
<td>80 1-080</td>
<td><img src="image3" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 2.23 (s, 3H), 2.59 (s, 3H), 3.99 (t, J = 4.4 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 6.92 (t, J = 7.9 Hz, 1H), 7.14 (d, J = 8.3 Hz, 1H), 7.18 (dd, J = 7.7, 1.5 Hz, 1H), 7.44 (dd, J = 8.1, 1.6 Hz, 1H), 7.50-7.55 (m, 2H), 7.87 (dd, J = 8.6, 2.3 Hz, 1H), 8.07 (d, J = 2.3 Hz, 1H), 8.15 (s, 1H), 10.42 (s, 1H).</td>
</tr>
<tr>
<td>81 1-081</td>
<td><img src="image4" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.59 (s, 3H), 2.23 (s, 3H), 3.99 (t, J = 4.4 Hz, 2H), 4.31 (t, J = 4.4 Hz, 2H), 5.28 (s, 1H), 6.91 (t, J = 7.9 Hz, 1H), 7.11-7.17 (m, 2H), 7.32 (d, J = 8.3 Hz, 1H), 7.42 (dd, J = 6.3, 1.4 Hz, 1H), 7.53 (dd, J = 6.6, 2.0 Hz, 1H), 7.70 (dd, J = 6.8, 2.6 Hz, 1H), 8.13-8.18 (m, 2H), 10.24 (s, 1H).</td>
</tr>
<tr>
<td>82 1-082</td>
<td><img src="image5" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 1.44 (s, 3H), 2.23 (s, 3H), 2.96 (s, 3H), 3.98 (t, J = 4.4 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 6.91 (t, J = 7.9 Hz, 1H), 7.14 (d, J = 8.4 Hz, 1H), 7.18 (dd, J = 7.7, 1.5 Hz, 1H), 7.33 (dd, J = 8.8 Hz, 2H), 7.42 (dd, J = 8.1, 1.5 Hz, 1H), 7.52 (dd, J = 8.6, 2.2 Hz, 1H), 7.70 (dd, J = 8.8 Hz, 2H), 8.15 (d, J = 2.6 Hz, 1H), 10.11 (s, 1H).</td>
</tr>
<tr>
<td>83 1-083</td>
<td><img src="image6" alt="Chemical Structure" /></td>
<td>(300 MHz, CHLOROFORM-d) 2.28 (s, 3H), 3.97 (s, 3H), 4.17 (t, J = 4.4 Hz, 2H), 4.52 (t, J = 4.6 Hz, 2H), 8.04 (dd, J = 8.6, 2.4 Hz, 1H), 6.99 (t, J = 8.1 Hz, 1H), 7.10 (d, J = 6.4 Hz, 1H), 7.29 (d, J = 8.8 Hz, 1H), 7.38 (dd, J = 8.4, 2.2 Hz, 1H), 7.44 (dd, J = 8.1, 1.5 Hz, 1H), 7.88 (dd, J = 9.0, 2.2 Hz, 2H), 8.17 (d, J = 1.8 Hz, 1H), 9.71 (s, 1H).</td>
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<tr>
<td>84 1-084</td>
<td><img src="image7" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d) 1.04-1.06 (m, 3H), 1.82-1.86 (m, 2H), 2.28 (s, 3H), 3.99-4.00 (m, 2H), 4.16-4.17 (m, 2H), 4.50-4.51 (m, 2H), 6.92-7.00 (m, 2H), 7.10 (dd, J = 8.3, 2.3 Hz, 1H), 7.27-7.30 (m, 1H), 7.36-7.45 (m, 2H), 7.57-7.60 (m, 1H), 7.87-7.91 (m, 1H), 8.17 (s, 1H), 9.57 (s, 1H).</td>
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<td>(400 MHz, CHLOROFORM-d) 1.04-1.06 (m, 3H), 1.82-1.86 (m, 2H), 2.28 (s, 3H), 3.99-4.00 (m, 2H), 4.16-4.17 (m, 2H), 4.50-4.51 (m, 2H), 6.92-7.00 (m, 2H), 7.10 (dd, J = 8.3, 2.3 Hz, 1H), 7.27-7.30 (m, 1H), 7.36-7.45 (m, 2H), 7.57-7.60 (m, 1H), 7.87-7.91 (m, 1H), 8.17 (s, 1H), 9.57 (s, 1H).</td>
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<tr>
<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<tr>
<td>86</td>
<td><img src="image1" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d) 1.44-1.45 (m, 3H), 2.28 (d, J = 3.2 Hz, 3H), 3.8-4.17 (m, 4H), 4.50-4.51 (m, 2H), 6.93-6.99 (m, 2H), 7.09-7.11 (m, 1H), 7.27-7.28 (m, 1H), 7.36-7.45 (m, 2H), 7.57-7.59 (m, 1H), 7.85-7.86 (m, 1H), 8.17 (s, 1H), 8.37 (s, 1H).</td>
</tr>
<tr>
<td>87</td>
<td><img src="image2" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d) 1.44 (t, J = 7.0 Hz, 3H), 2.29 (s, 3H), 2.29 (s, 3H), 4.12 (q, J = 7.0 Hz, 2H), 4.17 (t, J = 4.4 Hz, 2H), 4.50 (t, J = 4.4 Hz, 2H), 6.82 (dd, J = 7.5, 1.9 Hz, 1H), 6.97 (t, J = 8.1 Hz, 1H), 7.09 (dd, J = 12.8, 8.1 Hz, 2H), 7.37 (dd, J = 8.3, 2.3 Hz, 2H), 7.42 (dd, J = 8.1, 1.9 Hz, 2H), 7.62 (d, J = 2.3 Hz, 1H), 7.89 (dd, J = 7.0, 1.4 Hz, 1H), 8.17 (s, 1H), 9.60 (s, 1H).</td>
</tr>
<tr>
<td>88</td>
<td><img src="image3" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 2.19 (s, 3H), 2.23 (s, 3H), 3.79 (t, J = 3.9 Hz, 2H), 4.32 (t, J = 4.6 Hz, 2H), 4.40 (s, 2H), 6.50 (t, J = 7.9 Hz, 1H), 7.09-7.17 (m, 3H), 7.24 (d, J = 7.9 Hz, 1H), 7.38-7.41 (m, 4H), 7.50 (dd, J = 8.8, 2.3 Hz, 1H), 8.16 (s, 1H), 10.07 (s, 1H).</td>
</tr>
<tr>
<td>89</td>
<td><img src="image4" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 2.12 (s, 3H), 2.28 (s, 3H), 3.70 (t, J = 4.6 Hz, 2H), 4.03-4.06 (m, 4H), 4.38 (t, J = 4.2 Hz, 2H), 6.95 (t, J = 7.9 Hz, 1H), 7.06 (d, J = 7.9 Hz, 1H), 7.20 (dd, J = 8.1, 1.6 Hz, 1H), 7.29 (dd, J = 14.8, 8.3 Hz, 2H), 7.44 (d, J = 11.1 Hz, 2H), 7.76 (d, J = 8.8 Hz, 1H), 8.19 (s, 1H), 10.06 (s, 1H).</td>
</tr>
<tr>
<td>90</td>
<td><img src="image5" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.64 (br s, 1H), 1.79 (t, J = 10.0 Hz, 2H), 1.91 (s, 3H), 2.30 (s, 3H), 3.27 (d, J = 7.4 Hz, 1H), 3.45 (d, J = 10.0 Hz, 3H), 3.81-3.84 (m, 1H), 4.02-4.06 (m, 3H), 4.39 (br s, 2H), 6.88 (t, J = 7.7 Hz, 1H), 7.26-7.35 (m, 2H), 7.48 (d, J = 7.9 Hz, 1H), 7.58 (d, J = 7.4 Hz, 1H), 7.86-7.92 (m, 3H), 8.21 (s, 1H), 10.63 (s, 1H).</td>
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<td>91</td>
<td><img src="image6" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 2.22 (s, 3H), 3.96 (q, J = 6.8 Hz, 2H), 4.29 (q, J = 6.8 Hz, 2H), 6.50 (t, J = 7.9 Hz, 1H), 7.12-7.15 (m, 2H), 7.35-7.43 (m, 3H), 7.52 (dd, J = 9.3, 2.3 Hz, 1H), 7.77 (d, J = 12.0 Hz, 2H), 8.14 (d, J = 2.3 Hz, 1H), 10.27 (s, 1H).</td>
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<td>92</td>
<td><img src="image7" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 2.20 (d, J = 1.1 Hz, 3H), 2.23 (s, 3H), 3.99 (t, J = 4.2 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 6.91 (t, J = 7.7 Hz, 1H), 7.19-7.20 (m, 3H), 7.38-7.42 (m, 2H), 7.53 (dd, J = 8.4, 2.2 Hz, 1H), 7.87 (dd, J = 12.1, 1.8 Hz, 1H), 8.16 (d, J = 2.2 Hz, 1H), 10.23 (s, 1H).</td>
</tr>
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<td>93</td>
<td><img src="image8" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 2.23 (s, 3H), 3.99 (t, J = 4.4 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 6.92 (t, J = 7.9 Hz, 1H), 7.14-7.19 (m, 2H), 7.44-7.55 (m, 3H), 8.15 (q, J = 0.7 Hz, 1H), 8.23 (dd, J = 8.8, 2.6 Hz, 1H), 8.75 (d, J = 2.6 Hz, 1H), 10.48 (s, 1H).</td>
</tr>
<tr>
<td>94</td>
<td><img src="image9" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.31 (t, J = 7.2 Hz, 3H), 3.79 (t, J = 4.4 Hz, 2H), 3.96 (q, J = 7.0 Hz, 2H), 4.42 (t, J = 4.4 Hz, 2H), 6.46 (dd, J = 8.1, 1.6 Hz, 1H), 6.76 (t, J = 7.9 Hz, 1H), 6.88 (d, J = 9.3 Hz, 2H), 7.07 (dd, J = 7.9, 1.4 Hz, 1H), 7.30 (dd, J = 7.5, 4.4 Hz, 1H), 7.64 (d, J = 8.8 Hz, 2H), 8.03 (dd, J = 7.9, 1.9 Hz, 1H), 8.41 (cd, J = 4.4, 1.6 Hz, 1H), 9.99 (s, 1H).</td>
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<tr>
<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<tr>
<td>95</td>
<td><img src="image1.png" alt="Image" /></td>
<td>(400 MHz, DMSO-d6) 1.49-1.54 (m, 2H), 1.61-1.67 (m, 4H), 2.91 (t, J = 5.1 Hz, 4H), 3.79 (t, J = 4.4 Hz, 2H), 4.42 (t, J = 4.4 Hz, 2H), 6.49 (dd, J = 6.1, 1.6 Hz, 1H), 6.79 (t, J = 7.9 Hz, 1H), 6.99 (t, J = 9.3 Hz, 1H), 7.05 (dd, J = 7.4, 1.4 Hz, 1H), 7.30 (dd, J = 8.0, 4.4 Hz, 1H), 7.39 (dd, J = 8.6, 2.1 Hz, 1H), 7.54 (dd, J = 14.0, 2.3 Hz, 1H), 8.03 (dd, J = 7.0, 1.9 Hz, 1H), 8.42 (dd, J = 4.6, 1.4 Hz, 1H), 10.13 (s, 1H).</td>
</tr>
<tr>
<td>96</td>
<td><img src="image2.png" alt="Image" /></td>
<td>(400 MHz, DMSO-d6) 1.08 (t, J = 7.0 Hz, 3H), 1.18-1.37 (m, 4H), 1.85-1.98 (m, 4H), 3.17-3.24 (m, 1H), 3.43 (q, J = 7.0 Hz, 2H), 3.67-3.73 (m, 1H), 3.76 (t, J = 4.4 Hz, 2H), 3.86 (t, J = 4.4 Hz, 2H), 6.44 (dd, J = 7.5, 1.4 Hz, 1H), 6.73 (t, J = 7.9 Hz, 1H), 7.07 (dd, J = 7.7, 1.6 Hz, 1H), 7.27 (dd, J = 8.0, 4.8 Hz, 1H), 7.92 (dd, J = 7.4 Hz, 1H), 8.01 (dd, J = 7.9, 1.9 Hz, 1H), 8.39 (dd, J = 4.8, 2.0 Hz, 1H).</td>
</tr>
<tr>
<td>97</td>
<td><img src="image3.png" alt="Image" /></td>
<td>(400 MHz, DMSO-d6) 1.05 (q, J = 6.0 Hz, 6H), 1.15-1.38 (m, 4H), 1.85-1.90 (m, 4H), 3.27-3.31 (m, 1H), 3.63-3.73 (m, 2H), 3.76 (t, J = 4.6 Hz, 2H), 4.38 (t, J = 4.6 Hz, 2H), 6.44 (dd, J = 7.9, 1.4 Hz, 1H), 6.73 (t, J = 7.6 Hz, 1H), 7.07 (dd, J = 7.7, 1.5 Hz, 1H), 7.27 (dd, J = 8.0, 4.6 Hz, 1H), 7.91 (dd, J = 7.9 Hz, 1H), 8.01 (dd, J = 7.9, 1.4 Hz, 1H), 8.39 (dd, J = 4.6, 1.4 Hz, 1H).</td>
</tr>
<tr>
<td>98</td>
<td><img src="image4.png" alt="Image" /></td>
<td>(400 MHz, DMSO-d6) 1.17-1.70 (m, 12H), 1.84-1.94 (m, 4H), 3.20-3.25 (m, 1H), 3.68-3.72 (m, 1H), 3.76 (t, J = 4.4 Hz, 2H), 3.88-4.01 (m, 1H), 4.38 (t, J = 4.4 Hz, 2H), 6.44 (dd, J = 8.1, 1.6 Hz, 1H), 6.72 (t, J = 7.9 Hz, 1H), 7.07 (dd, J = 7.9, 1.4 Hz, 1H), 7.27 (dd, J = 8.0, 4.8 Hz, 1H), 7.91 (dd, J = 7.9 Hz, 1H), 8.01 (dd, J = 7.9, 1.4 Hz, 1H), 8.39 (dd, J = 4.9, 1.5 Hz, 1H).</td>
</tr>
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<td>99</td>
<td><img src="image5.png" alt="Image" /></td>
<td>(400 MHz, DMSO-d6) 1.12-1.47 (m, 10H), 1.62-1.66 (m, 2H), 1.74-1.78 (m, 2H), 1.85-1.91 (m, 4H), 3.52-3.56 (m, 2H), 3.66-3.72 (m, 1H), 3.76 (t, J = 4.6 Hz, 2H), 4.38 (t, J = 4.4 Hz, 2H), 6.43 (dd, J = 8.1, 1.6 Hz, 1H), 6.72 (t, J = 7.9 Hz, 1H), 7.07 (dd, J = 7.9, 1.4 Hz, 1H), 7.27 (dd, J = 8.0, 4.8 Hz, 1H), 7.91 (dd, J = 7.9 Hz, 1H), 8.01 (dd, J = 7.9, 1.9 Hz, 1H), 8.39 (dd, J = 4.5, 1.5 Hz, 1H).</td>
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<tr>
<td>100</td>
<td><img src="image6.png" alt="Image" /></td>
<td>(400 MHz, DMSO-d6) 0.38 (t, J = 4.4 Hz, 2H), 4.43 (t, J = 4.4 Hz, 2H), 6.53 (dd, J = 8.1, 1.5 Hz, 1H), 6.82 (t, J = 7.9 Hz, 1H), 7.07 (dd, J = 7.5, 1.7 Hz, 1H), 7.32 (dd, J = 7.9, 4.6 Hz, 1H), 7.71 (dd, J = 8.8 Hz, 2H), 7.97 (dd, J = 8.8 Hz, 2H), 8.05 (dd, J = 8.1, 1.5 Hz, 1H), 8.43 (dd, J = 2.1, 1.0 Hz, 1H), 10.52 (s, 1H).</td>
</tr>
<tr>
<td>101</td>
<td><img src="image7.png" alt="Image" /></td>
<td>(300 MHz, DMSO-d6) 0.46 (s, J = 7.9 Hz, 1H), 0.70 (s, J = 7.1 Hz, 1H), 0.77 (s, J = 8.5 Hz, 1H), 0.80 (dd, J = 8.1, 1.5 Hz, 1H), 0.80 (t, J = 7.9 Hz, 1H), 1.70 (s, J = 8.1 Hz, 1H), 2.70 (s, J = 8.1 Hz, 1H), 4.42 (s, J = 7.9 Hz, 1H), 7.06 (dd, J = 14.9, 2.4 Hz, 1H), 8.42 (dd, J = 7.9, 1.7 Hz, 1H), 8.43 (dd, J = 2.2, 1.1 Hz, 1H), 10.13 (s, 1H).</td>
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<tr>
<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<tr>
<td>103 1-103</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.10 (d, J = 5.9 Hz, 6H), 1.84-1.67 (m, 2H), 1.95-2.04 (m, 2H), 2.92-2.95 (m, 2H), 3.26-3.30 (m, 2H), 3.55-3.59 (m, 1H), 3.72-3.76 (m, 3H), 4.43 (t, J = 4.4 Hz, 2H), 6.51 (dd, J = 8.1, 1.8 Hz, 1H), 8.30 (t, J = 7.7 Hz, 1H), 7.06 (dd, J = 7.7, 1.5 Hz, 1H), 7.30-7.33 (m, 2H), 7.45 (d, J = 8.4 Hz, 1H), 7.73 (d, J = 14.7 Hz, 1H), 8.04 (dd, J = 8.1, 1.8 Hz, 1H), 8.43 (dd, J = 4.8, 1.5 Hz, 1H), 10.24 (s, 1H).</td>
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<td>104 1-104</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d) δ: 1.03 (s, 3H), 2.09-2.15 (m, 1H), 3.79 (d, J = 7.0 Hz, 2H), 3.95 (t, J = 4.4 Hz, 2H), 4.63 (t, J = 4.6 Hz, 2H), 6.47 (dd, J = 8.1, 1.6 Hz, 1H), 6.90-6.94 (m, 2H), 7.12 (q, J = 4.2 Hz, 1H), 7.30 (t, J = 5.3 Hz, 1H), 7.58 (dd, J = 13.0, 2.9 Hz, 1H), 7.76-7.80 (m, 2H), 8.38 (dd, J = 4.6, 1.4 Hz, 1H), 9.58 (s, 1H).</td>
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<td>105 1-105</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 0.98 (t, J = 6.8 Hz, 3H), 1.20-1.33 (m, 4H), 1.85-1.97 (m, 4H), 2.22 (s, 3H), 3.14-3.24 (m, 1H), 3.43 (q, J = 6.8 Hz, 2H), 3.66-3.76 (m, 1H), 3.95 (t, J = 4.4 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (dd, J = 8.0, 7.6 Hz, 1H), 7.17 (d, J = 8.8 Hz, 1H), 7.17 (dd, J = 8.0, 2.0 Hz, 1H), 7.33 (dd, J = 8.0, 1.2 Hz, 1H), 7.48 (dd, J = 8.4, 2.4 Hz, 1H), 7.90 (d, J = 8.0 Hz, 1H), 8.11 (d, J = 2.4 Hz, 1H).</td>
</tr>
<tr>
<td>106 1-106</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 0.94 (t, J = 6.5 Hz, 6H), 1.16-1.37 (m, 4H), 1.81-1.91 (m, 4H), 2.20 (s, 3H), 3.23-3.33 (m, 1H), 3.62-3.73 (m, 1H), 3.99 (t, J = 4.4 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (dd, J = 8.4, 7.2 Hz, 1H), 7.07 (d, J = 8.6 Hz, 1H), 7.18 (dd, J = 8.0, 2.0 Hz, 1H), 7.33 (dd, J = 8.0, 2.0 Hz, 1H), 7.48 (dd, J = 8.0, 2.0 Hz, 1H), 7.89 (d, J = 8.0 Hz, 1H), 8.11 (d, J = 1.6 Hz, 1H).</td>
</tr>
<tr>
<td>107 1-107</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 1.05-1.36 (m, 9H), 1.40-1.52 (m, 9H), 2.20 (s, 3H), 3.26-3.38 (m, 1H), 3.64-3.74 (m, 1H), 3.95 (t, J = 4.4 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (dd, J = 8.0, 8.0 Hz, 1H), 7.07 (d, J = 8.5 Hz, 1H), 7.17 (dd, J = 7.4, 1.4 Hz, 1H), 7.33 (dd, J = 8.0, 1.4 Hz, 1H), 7.49 (dd, J = 8.5, 2.6 Hz, 1H), 7.90 (d, J = 8.0 Hz, 1H), 8.12 (d, J = 1.2 Hz, 1H).</td>
</tr>
<tr>
<td>108 1-108</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 1.04-1.34 (m, 4H), 1.72-1.97 (m, 4H), 2.20 (s, 3H), 2.48-2.58 (m, 1H), 3.30 (brs, 3H), 3.59-3.71 (m, 1H), 3.95 (t, J = 4.4 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (dd, J = 8.0, 8.0 Hz, 1H), 7.07 (d, J = 8.5 Hz, 1H), 7.19 (dd, J = 8.0, 1.5 Hz, 1H), 7.33 (dd, J = 8.0, 1.5 Hz, 1H), 7.48 (dd, J = 8.5, 2.6 Hz, 1H), 7.86 (d, J = 8.0 Hz, 1H), 8.12 (d, J = 2.0 Hz, 1H).</td>
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<td>109 1-109</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 1.48-1.84 (m, 8H), 2.20 (s, 3H), 3.78-3.88 (m, 5H), 3.95 (t, J = 4.4 Hz, 2H), 4.28 (t, J = 4.4 Hz, 2H), 6.84 (dd, J = 8.0, 8.0 Hz, 1H), 7.08 (d, J = 8.5 Hz, 1H), 7.18 (dd, J = 8.0, 1.5 Hz, 1H), 7.34 (dd, J = 8.0, 1.5 Hz, 1H), 7.48 (dd, J = 8.5, 2.5 Hz, 1H), 7.97 (d, J = 8.0 Hz, 1H), 8.12 (d, J = 2.5 Hz, 1H).</td>
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<td>110 1-110</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 1.74-1.84 (m, 2H), 2.05-2.09 (m, 2H), 2.20 (s, 3H), 2.48-2.53 (m, 2H), 2.41-2.46 (m, 2H), 3.95 (t, J = 4.4 Hz, 2H), 4.28-4.28 (m, 3H), 6.85 (t, J = 7.9 Hz, 1H), 7.08 (d, J = 8.8 Hz, 1H), 7.18 (dd, J = 7.7, 1.4 Hz, 1H), 7.35 (dd, J = 8.1, 1.6 Hz, 1H), 7.49 (dd, J = 8.8, 2.3 Hz, 1H), 8.11-8.13 (m, 2H).</td>
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<tr>
<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<tr>
<td>111</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.45-1.78 (m, 8H), 2.16-2.20 (m, 4H), 2.43-2.45 (m, 4H), 3.55 (t, J = 4.6 Hz, 4H), 3.98 (s, J = 4.4 Hz, 3H), 4.31 (t, J = 4.6 Hz, 2H), 6.85 (t, J = 7.9 Hz, 1H), 7.05 (d, J = 8.3 Hz, 1H), 7.25 (dd, J = 7.7, 1.6 Hz, 1H), 7.35 (dd, J = 8.1, 1.6 Hz, 1H), 7.49 (dd, J = 8.3, 2.3 Hz, 1H), 8.02 (d, J = 7.4 Hz, 1H), 8.12 (s, 1H).</td>
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<td>112</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.20-1.32 (m, 4H), 1.80 (m, 2H), 1.90 (m, 3H), 2.46-2.50 (m, 3H), 3.64-3.68 (m, 1H), 3.95 (s, J = 4.4 Hz, 2H), 7.07 (d, J = 7.9 Hz, 2H), 7.18 (d, J = 7.9 Hz, 1H), 7.33 (dd, J = 8.1, 1.6 Hz, 1H), 7.48 (dd, J = 8.3, 2.3 Hz, 1H), 7.88 (d, J = 7.9 Hz, 1H), 8.12 (s, 1H).</td>
</tr>
<tr>
<td>113</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.94 (s, J = 7.0 Hz, 6H), 1.25-1.32 (m, 4H), 1.70 (m, 2H), 1.91 (m, 2H), 2.20 (t, 3H), 2.43-2.45 (m, 3H), 3.64-3.68 (m, 1H), 3.95 (s, J = 4.4 Hz, 2H), 7.07 (d, J = 7.9 Hz, 1H), 7.07 (d, J = 7.9 Hz, 1H), 7.19 (dd, J = 7.7, 1.6 Hz, 1H), 7.33 (dd, J = 8.1, 1.6 Hz, 1H), 7.48 (dd, J = 8.3, 2.3 Hz, 1H), 7.87 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 2.3 Hz, 1H).</td>
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<tr>
<td>114</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.55-1.59 (m, 12H), 2.17-2.24 (m, 4H), 2.45-2.53 (m, 4H), 3.88-3.90 (m, 1H), 3.97 (s, J = 4.4 Hz, 2H), 4.31 (t, J = 4.4 Hz, 2H), 6.85 (t, J = 7.9 Hz, 1H), 7.08 (d, J = 8.8 Hz, 1H), 7.24 (dd, J = 7.9, 1.4 Hz, 1H), 7.34 (d, J = 7.9 Hz, 1H), 7.48 (dd, J = 8.6, 2.1 Hz, 1H), 7.97 (d, J = 7.9 Hz, 1H), 8.12 (s, 1H).</td>
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<td>115</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.18-1.34 (m, 4H), 1.84 (m, 4H), 1.95-1.96 (m, 5H), 2.20 (t, 3H), 3.65-3.70 (m, 1H), 3.95 (s, J = 4.4 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (t, J = 7.9 Hz, 1H), 7.07 (d, J = 8.8 Hz, 1H), 7.16 (dd, J = 7.9, 1.4 Hz, 1H), 7.33 (dd, J = 7.9, 1.4 Hz, 1H), 7.48 (dd, J = 8.6, 2.3 Hz, 1H), 7.89 (d, J = 7.9 Hz, 1H), 8.11 (s, 1H).</td>
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<td>116</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.33-2.16 (m, 14H), 2.27 (t, 3H), 2.96-2.98 (m, 2H), 3.33-3.37 (m, 1H), 3.39-3.39 (m, 2H), 3.68-3.76 (m, 3H), 3.42-3.45 (m, 1H), 6.67-6.94 (m, 1H), 7.25-7.41 (m, 1H), 7.79 (d, J = 8.3 Hz, 1H), 7.99-8.12 (m, 1H), 8.17 (t, 1H), 10.43-10.53 (m, 1H).</td>
</tr>
<tr>
<td>117</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.29-1.40 (m, 2H), 1.56-1.65 (m, 2H), 1.94-2.01 (m, 2H), 2.18-2.19 (m, 2H), 2.24 (t, 3H), 3.15 (m, 3H), 3.33-3.38 (m, 1H), 3.44-3.45 (m, 1H), 4.31 (t, J = 4.4 Hz, 2H), 6.88 (t, J = 7.9 Hz, 1H), 7.21 (d, J = 7.9 Hz, 1H), 7.36 (dd, J = 8.3, 1.4 Hz, 1H), 7.67 (d, J = 8.3 Hz, 1H), 8.00 (d, J = 7.9 Hz, 1H), 8.15 (s, 1H), 11.13 (s, 1H).</td>
</tr>
<tr>
<td>118</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.21-1.38 (m, 4H), 1.76-1.88 (m, 7H), 2.20 (t, 3H), 3.44-3.49 (m, 1H), 3.65-3.72 (m, 1H), 3.95 (t, J = 4.2 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (t, J = 7.9 Hz, 1H), 7.08 (d, J = 8.3 Hz, 1H), 7.18 (dd, J = 7.9, 1.4 Hz, 1H), 7.33 (dd, J = 8.3, 1.4 Hz, 1H), 7.48 (dd, J = 8.8, 2.3 Hz, 1H), 7.72 (d, J = 7.9 Hz, 1H), 7.92 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 1.9 Hz, 1H).</td>
</tr>
<tr>
<td>Table 16</td>
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<tr>
<th>Ex. No.</th>
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<th>NMR</th>
</tr>
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<tbody>
<tr>
<td>119 1-119</td>
<td><img src="image1.png" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.83-0.92 (m, 2H), 1.09-1.47 (m, 8H), 1.59-1.69 (m, 5H), 1.84-1.85 (m, 4H), 2.20 (s, 3H), 3.13-3.33 (m, 3H), 3.09-3.74 (m, 1H), 3.95 (J = 4.4 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (t, J = 7.9 Hz, 1H), 7.07 (d, J = 8.3 Hz, 1H), 7.18 (dd, J = 7.7, 1.6 Hz, 1H), 7.33 (dd, J = 7.9, 1.4 Hz, 1H), 7.48 (dd, J = 8.3, 2.3 Hz, 1H), 7.90 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 2.3 Hz, 1H).</td>
</tr>
<tr>
<td>120 1-120</td>
<td><img src="image2.png" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.30 (s, J = 7.0 Hz, 3H), 3.07 (q, J = 7.0 Hz, 2H), 4.02 (t, J = 4.6 Hz, 2H), 4.33 (t, J = 4.6 Hz, 2H), 5.68 (d, J = 9.3 Hz, 2H), 6.63 (d, J = 7.9 Hz, 1H), 7.21-7.24 (m, 2H), 7.47 (d, J = 8.3, 1.0 Hz, 1H), 7.61 (d, J = 9.3 Hz, 2H), 7.74 (dd, J = 9.0, 2.6 Hz, 1H), 8.30 (d, J = 2.8 Hz, 1H), 8.98 (s, 1H).</td>
</tr>
<tr>
<td>121 1-121</td>
<td><img src="image3.png" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 4.02 (t, J = 4.5 Hz, 3H), 4.32 (t, J = 4.5 Hz, 2H), 6.94 (t, J = 7.9 Hz, 1H), 7.21-7.23 (m, 2H), 7.38 (d, J = 8.8 Hz, 2H), 7.49 (dd, J = 7.9, 1.4 Hz, 1H), 7.73-7.77 (m, 4H), 8.30 (d, J = 2.8 Hz, 1H), 10.27 (s, 1H).</td>
</tr>
<tr>
<td>122 1-122</td>
<td><img src="image4.png" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d5) 6.5-1.54-1.63 (m, 2H), 2.16-2.01 (m, 2H), 2.74-2.77 (m, 2H), 3.16-3.18 (m, 2H), 3.29-3.33 (m, 4H), 4.03 (t, J = 4.2 Hz, 2H), 4.34 (t, J = 4.4 Hz, 2H), 6.95-7.02 (m, 2H), 7.23 (d, J = 9.2 Hz, 2H), 7.38 (d, J = 8.4 Hz, 1H), 7.50 (dd, J = 8.1, 0.7 Hz, 1H), 7.64 (dd, J = 15.0, 2.2 Hz, 1H), 7.76 (dd, J = 9.0, 2.8 Hz, 1H), 8.32 (d, J = 2.6 Hz, 1H), 10.12 (s, 1H).</td>
</tr>
<tr>
<td>123 1-123</td>
<td><img src="image5.png" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d5) 6.4.04 (t, J = 4.4 Hz, 2H), 4.35 (t, J = 4.4 Hz, 2H), 6.96 (t, J = 7.9 Hz, 1H), 7.23-7.26 (m, 2H), 7.53 (dd, J = 8.3, 1.7 Hz, 1H), 7.71 (d, J = 8.5 Hz, 2H), 7.77 (dd, J = 9.0, 2.8 Hz, 1H), 7.95 (d, J = 8.4 Hz, 2H), 8.32 (d, J = 2.6 Hz, 1H), 10.49 (s, 1H).</td>
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<tr>
<td>124 1-124</td>
<td><img src="image6.png" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 2.25 (s, 3H), 2.84 (t, J = 7.2 Hz, 2H), 3.30 (d, J = 6.6 Hz, 2H), 3.98 (t, J = 4.4 Hz, 2H), 4.28 (t, J = 4.6 Hz, 2H), 6.89 (t, J = 7.9 Hz, 1H), 7.24-7.35 (m, 5H), 7.66 (d, J = 8.8 Hz, 1H), 8.15-8.20 (m, 2H).</td>
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<td>125 1-125</td>
<td><img src="image7.png" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d) 2.28 (s, 3H), 2.91 (t, J = 7.0 Hz, 2H), 3.73 (q, J = 6.5 Hz, 2H), 4.07 (t, J = 4.6 Hz, 2H), 4.21 (t, J = 4.4 Hz, 2H), 6.92 (t, J = 7.9 Hz, 1H), 7.08 (t, J = 8.3 Hz, 1H), 7.20-7.21 (m, 2H), 7.27-7.30 (m, 2H), 7.38 (dd, J = 7.9, 1.9 Hz, 2H), 7.74 (s, 1H), 7.82 (d, J = 7.9, 1.4 Hz, 1H), 8.14 (s, 1H).</td>
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<td>126 1-126</td>
<td><img src="image8.png" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d) 2.28 (s, 3H), 2.51 (t, J = 6.7 Hz, 2H), 3.74 (q, J = 6.5 Hz, 2H), 4.08 (s, 2H), 4.24 (t, J = 4.4 Hz, 2H), 7.76 (s, 1H), 7.83 (d, J = 8.5 Hz, 1H), 8.14 (s, 1H).</td>
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<td>Ex. No.</td>
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<td>127</td>
<td><img src="image1.png" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 2.26 (s, 3H), 2.98 (t, J = 7.2 Hz, 2H), 3.53 (s, J = 6.5 Hz, 2H), 3.99 (t, J = 4.4 Hz, 2H), 6.90 (t, J = 7.9 Hz, 1H), 7.23-7.33 (m, 4H), 7.38-7.40 (m, 2H), 7.44 (td, J = 7.7, 1.6 Hz, 1H), 7.69 (d, J = 8.8 Hz, 1H), 8.16 (s, 1H), 8.23 (t, J = 5.3 Hz, 1H).</td>
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<tr>
<td>128</td>
<td><img src="image2.png" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 4.16 (t, J = 4.4 Hz, 2H), 4.37 (t, J = 4.2 Hz, 2H), 6.69 (t, J = 7.9 Hz, 1H), 7.24-7.28 (m, 2H), 7.60 (dd, J = 8.3, 1.7 Hz, 1H), 7.71 (d, J = 8.8 Hz, 2H), 7.95 (d, J = 8.4 Hz, 2H), 8.08 (dd, J = 8.8, 2.2 Hz, 1H), 8.77 (d, J = 2.6 Hz, 1H), 10.51 (s, 1H).</td>
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<td>129</td>
<td><img src="image3.png" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.51 (s, 6H), 3.78 (t, J = 4.2 Hz, 2H), 4.43 (t, J = 4.2 Hz, 2H), 5.33 (s, 1H), 6.46-6.52 (m, 1H), 6.83 (t, J = 10.7 Hz, 1H), 7.05 (dd, J = 7.7, 1.5 Hz, 1H), 7.66-7.71 (m, 2H), 7.94-7.97 (m, 2H), 8.04 (d, J = 2.2 Hz, 1H), 8.51 (d, J = 2.2 Hz, 1H), 10.50 (s, 1H).</td>
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<td>130</td>
<td><img src="image4.png" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 1.40 (d, J = 8.6 Hz, 3H), 3.76-3.80 (m, 2H), 4.12-4.14 (m, 2H), 4.62-4.64 (m, 1H), 5.43 (d, J = 4.8 Hz, 1H), 6.47 (dd, J = 8.1, 1.5 Hz, 1H), 6.81 (d, J = 7.9 Hz, 1H), 7.05 (d, J = 3.9 Hz, 1H), 7.70 (d, J = 8.8 Hz, 2H), 7.95-7.96 (m, 4H), 8.40-8.40 (m, 1H), 10.50 (s, 1H).</td>
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<tr>
<td>131</td>
<td><img src="image5.png" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 3.90-3.92 (m, 2H), 4.42-4.43 (m, 2H), 6.77 (d, J = 4.0 Hz, 1H), 6.87 (d, J = 7.7 Hz, 1H), 7.16-7.17 (m, 1H), 7.73 (d, J = 9.2 Hz, 2H), 7.98 (d, J = 14.3 Hz, 1H), 8.30 (d, J = 1.8 Hz, 1H), 8.83 (d, J = 1.8 Hz, 1H), 10.72 (s, 1H).</td>
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<td>132</td>
<td><img src="image6.png" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d) δ: 1.33 (s, 6H), 3.92 (s, 3H), 4.36 (t, J = 4.6 Hz, 2H), 4.55 (t, J = 4.6 Hz, 2H), 7.06 (t, J = 7.9 Hz, 1H), 7.21 (d, J = 8.6 Hz, 1H), 7.38 (t, J = 4.4 Hz, 2H), 7.52 (dd, J = 6.1, 1.8 Hz, 3H), 7.58 (dd, J = 6.7, 4.9 Hz, 3H), 8.02 (dd, J = 7.9, 1.4 Hz, 2H), 8.08 (dd, J = 8.8, 2.3 Hz, 2H), 8.93 (d, J = 2.3 Hz, 1H), 9.48 (s, 1H).</td>
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<tr>
<td>133</td>
<td><img src="image7.png" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d) δ: 1.28 (s, 3H), 3.96 (t, J = 4.2 Hz, 2H), 4.37 (t, J = 4.2 Hz, 2H), 6.98 (t, J = 7.9 Hz, 1H), 7.26 (dd, J = 20.4, 8.3 Hz, 2H), 7.35 (d, J = 8.8 Hz, 2H), 7.57 (d, J = 8.3 Hz, 1H), 7.64 (d, J = 8.8 Hz, 2H), 8.07 (dd, J = 8.8, 1.8 Hz, 1H), 8.77 (d, J = 2.3 Hz, 1H), 10.07 (s, 1H).</td>
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<tr>
<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<tr>
<td>134</td>
<td><img src="image" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) 3.92 (s, J = 4.2 Hz, 2H), 4.43 (t, J = 4.4 Hz, 2H), 6.76 (dd, J = 8.1, 1.5 Hz, 2H), 6.86 (t, J = 7.9 Hz, 1H), 7.17 (dd, J = 7.5, 1.7 Hz, 1H), 7.71 (d, J = 8.0 Hz, 2H), 7.97 (d, J = 7.1 Hz, 2H), 8.31 (d, J = 7.3 Hz, 1H), 8.83 (d, J = 2.9 Hz, 1H), 10.53 (s, 1H).</td>
</tr>
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<td>135</td>
<td><img src="image" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) 1.26 (s, 3H), 3.92 (t, J = 4.2 Hz, 2H), 4.44 (t, J = 4.2 Hz, 2H), 6.73 (dd, J = 8.1, 1.5 Hz, 1H), 8.84 (J, J = 7.9 Hz, 1H), 7.18 (dd, J = 7.5, 1.7 Hz, 1H), 7.30 (d, J = 8.6 Hz, 2H), 7.65 (d, J = 8.6 Hz, 2H), 8.30 (d, J = 2.2 Hz, 1H), 8.83 (d, J = 7.2 Hz, 1H), 10.07 (s, 1H).</td>
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<td>136</td>
<td><img src="image" alt="Chemical Compound" /></td>
<td>(300 MHz, DMSO-d6) 5.2 (s, 3H), 3.94 (t, J = 4.6 Hz, 2H), 4.44 (t, J = 4.4 Hz, 2H), 6.76 (dd, J = 8.1, 1.5 Hz, 1H), 6.87 (d, J = 7.9 Hz, 1H), 7.17 (dd, J = 7.3, 1.5 Hz, 1H), 7.71 (d, J = 8.4 Hz, 2H), 7.97 (d, J = 8.4 Hz, 3H), 8.30 (d, J = 2.2 Hz, 1H), 5.91 (d, J = 1.8 Hz, 1H), 10.52 (s, 1H).</td>
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<td>137</td>
<td><img src="image" alt="Chemical Compound" /></td>
<td>(400 MHz, DMSO-d6) 1.27 (s, 3H), 2.78 (d, J = 4.6 Hz, 2H), 4.14 (t, J = 4.2 Hz, 2H), 4.36 (t, J = 4.6 Hz, 2H), 5.08 (t, J = 7.9 Hz, 1H), 7.25 (d, J = 7.4 Hz, 2H), 7.35 (d, J = 8.8 Hz, 2H), 7.56 (d, J = 8.3 Hz, 1H), 7.64 (d, J = 8.6 Hz, 2H), 8.05 (dd, J = 8.8, 2.3 Hz, 1H), 8.39 (d, J = 5.6 Hz, 1H), 8.73 (d, J = 2.3 Hz, 1H), 10.05 (s, 1H).</td>
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<td>138</td>
<td><img src="image" alt="Chemical Compound" /></td>
<td>(400 MHz, DMSO-d6) 1.28 (s, 3H), 4.15 (t, J = 4.4 Hz, 2H), 4.36 (t, J = 4.4 Hz, 2H), 5.06 (t, J = 7.9 Hz, 1H), 7.25-7.33 (m, 9H), 7.55 (d, J = 8.3 Hz, 1H), 7.64 (d, J = 8.6 Hz, 2H), 7.93 (s, 1H), 8.00 (dd, J = 8.8, 2.6 Hz, 1H), 8.76 (d, J = 2.3 Hz, 1H), 10.07 (s, 1H).</td>
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<td>139</td>
<td><img src="image" alt="Chemical Compound" /></td>
<td>(400 MHz, DMSO-d6) 1.08 (t, J = 7.0 Hz, 6H), 1.27 (s, 9H), 3.31-3.40 (m, 4H), 3.82 (t, J = 4.4 Hz, 2H), 4.34 (t, J = 4.4 Hz, 2H), 6.81 (t, J = 7.9 Hz, 1H), 7.02-7.16 (m, 4H), 7.34 (d, J = 8.8 Hz, 2H), 7.64 (d, J = 8.3 Hz, 2H), 7.86 (d, J = 3.2 Hz, 1H), 10.02 (s, 1H).</td>
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<td>140</td>
<td><img src="image" alt="Chemical Compound" /></td>
<td>(400 MHz, CHLOROFORM-d) 1.33 (s, 9H), 4.44 (dd, J = 6.0, 3.2 Hz, 2H), 4.59 (t, J = 4.9 Hz, 2H), 7.10-7.12 (m, 1H), 7.25-7.26 (m, 1H), 7.30 (dd, J = 8.8, 1.6 Hz, 2H), 7.52-7.54 (m, 1H), 7.56 (dd, J = 8.6, 1.6 Hz, 2H), 8.00-8.11 (m, 1H), 8.26-8.28 (m, 1H), 8.17 (t, J = 1.6 Hz, 1H), 8.40 (s, 1H).</td>
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<td>Ex. No.</td>
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<td>141 1-141</td>
<td><img src="image1" alt="Chemical Structure" /></td>
<td>(400 MHz, CD$_3$OD) δ 1.32 (s, 9H), 3.54 (s, 2H), 4.02 (t, J = 4.4 Hz, 2H), 4.49 (t, J = 4.4 Hz, 2H), 6.92 (t, J = 8.1 Hz, 1H), 7.00-7.03 (m, 2H), 7.22-7.26 (m, 1H), 7.37 (t, J = 4.4 Hz, 2H), 7.59 (dd, J = 8.3 Hz, 2H), 7.80 (dd, J = 7.8, 1.5 Hz, 1H), 7.91 (d, J = 8.3 Hz, 1H), 8.52 (s, 1H), 8.69 (s, 1H), 9.53 (s, 1H).</td>
</tr>
<tr>
<td>142 1-142</td>
<td><img src="image2" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) δ 1.28 (s, 9H), 3.05 (s, 3H), 3.98 (t, J = 4.2 Hz, 2H), 4.33 (t, J = 4.2 Hz, 2H), 6.91 (t, J = 7.8 Hz, 1H), 7.19 (dd, J = 13.0, 8.3 Hz, 2H), 7.33 (dd, J = 12.3, 6.7 Hz, 2H), 7.40 (d, J = 8.3 Hz, 1H), 7.64 (d, J = 8.3 Hz, 2H), 7.92 (t, J = 5.6 Hz, 1H), 8.40 (s, 1H), 10.05 (s, 2H).</td>
</tr>
<tr>
<td>143 1-143</td>
<td><img src="image3" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) δ 3.29 (s, 3H), 4.06 (t, J = 4.4 Hz, 2H), 4.34 (t, J = 4.4 Hz, 2H), 6.36 (s, 2H), 6.95 (t, J = 7.8 Hz, 1H), 7.22 (d, J = 8.3 Hz, 2H), 7.52 (dd, J = 8.1, 1.6 Hz, 1H), 7.65 (dd, J = 8.6, 2.6 Hz, 1H), 7.71 (d, J = 8.8 Hz, 2H), 7.95 (d, J = 8.8 Hz, 2H), 8.26 (d, J = 1.9 Hz, 1H), 10.50 (s, 1H).</td>
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<td>144 1-144</td>
<td><img src="image4" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d$_6$) δ 1.34 (t, J = 7.0 Hz, 3H), 3.91 (t, J = 4.4 Hz, 2H), 4.08 (q, J = 7.1 Hz, 2H), 4.33 (t, J = 4.2 Hz, 2H), 6.88 (t, J = 7.9 Hz, 1H), 7.12-7.17 (m, 2H), 7.28-7.41 (m, 4H), 7.65 (d, J = 9.2 Hz, 2H), 8.06 (d, J = 3.3 Hz, 1H), 10.31 (s, 1H).</td>
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<tr>
<td>145 1-145</td>
<td><img src="image5" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) δ 3.71 (t, J = 4.4 Hz, 2H), 3.90 (s, 3H), 4.44 (t, J = 4.4 Hz, 2H), 6.36 (dd, J = 7.9, 1.4 Hz, 1H), 6.78 (t, J = 7.8 Hz, 1H), 6.99 (dd, J = 7.0, 1.4 Hz, 1H), 7.71 (d, J = 8.3 Hz, 2H), 7.77 (d, J = 2.8 Hz, 1H), 7.97 (d, J = 8.3 Hz, 2H), 8.22 (d, J = 2.8 Hz, 1H), 10.50 (s, 1H).</td>
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<tr>
<td>146 1-146</td>
<td><img src="image6" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) δ 3.88 (t, J = 4.4 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 6.66 (t, J = 7.9 Hz, 1H), 7.07-7.10 (m, 2H), 7.16-7.23 (m, 2H), 7.70 (d, J = 8.3 Hz, 2H), 7.94-7.95 (m, 3H), 9.72 (brs, 1H), 10.48 (s, 1H).</td>
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<td>146 1-146</td>
<td><img src="image7" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) δ 3.81 (s, 3H), 3.92 (t, J = 4.4 Hz, 2H), 4.34 (t, J = 4.4 Hz, 2H), 6.89 (t, J = 7.8 Hz, 1H), 7.12 (dd, J = 7.5, 1.8 Hz, 1H), 7.20 (d, J = 9.3 Hz, 1H), 7.31 (dd, J = 7.9, 1.6 Hz, 1H), 7.41 (dd, J = 9.3, 3.2 Hz, 1H), 7.71 (d, J = 8.8 Hz, 2H), 7.96 (d, J = 8.8 Hz, 2H), 8.08 (d, J = 3.2 Hz, 1H), 10.50 (s, 1H).</td>
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<tr>
<td>147 1-147</td>
<td><img src="image8" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) δ 1.28 (s, 9H), 3.09 (s, 3H), 3.98 (t, J = 4.4 Hz, 2H), 4.34 (t, J = 4.4 Hz, 2H), 6.80 (t, J = 7.8 Hz, 1H), 7.02 (dd, J = 5.3, 1.7 Hz, 1H), 7.20 (d, J = 1.5 Hz, 1H), 7.28 (dd, J = 7.7, 1.5 Hz, 1H), 7.35 (d, J = 8.8 Hz, 2H), 7.56 (dd, J = 8.4, 1.5 Hz, 1H), 7.64 (d, J = 8.8 Hz, 2H), 8.27 (d, J = 5.1 Hz, 1H), 10.04 (s, 1H).</td>
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<td>149 1-149</td>
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<td>(300 MHz, DMSO-d6) 3.99 (t, J = 4.4 Hz, 2H), 4.35 (t, J = 4.2 Hz, 2H), 6.84 (q, J = 7.7 Hz, 1H), 7.10-7.28 (m, 2H), 7.22 (s, 2H), 7.45 (d, J = 4.0 Hz, 1H), 7.65 (dd, J = 8.6, 3.1 Hz, 1H), 7.71 (d, J = 8.8 Hz, 2H), 7.96 (d, J = 8.8 Hz, 2H), 8.31 (d, J = 2.9 Hz, 1H), 10.49 (s, 1H).</td>
</tr>
<tr>
<td>150 1-150</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 3.83 (t, J = 4.2 Hz, 2H), 4.43 (t, J = 4.4 Hz, 2H), 6.74-6.77 (m, 1H), 6.85 (d, J = 7.9 Hz, 1H), 7.09 (dd, J = 7.3, 1.5 Hz, 1H), 7.71 (d, J = 8.8 Hz, 2H), 8.00-8.07 (m, 1H), 8.35 (d, J = 2.2 Hz, 1H), 10.51 (s, 1H).</td>
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<tr>
<td>151 1-151</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, CHLOROFORM-d) 1.32 (s, 3H), 3.82 (s, 3H), 3.93 (t, J = 4.4 Hz, 2H), 4.58 (t, J = 4.4 Hz, 2H), 6.68 (dd, J = 8.1, 1.5 Hz, 1H), 6.85 (t, J = 7.9 Hz, 1H), 7.10 (dd, J = 8.1, 1.1 Hz, 1H), 7.37 (d, J = 8.8 Hz, 2H), 7.60 (d, J = 8.4 Hz, 2H), 7.75 (dd, J = 7.9, 1.7 Hz, 1H), 8.03 (dd, J = 4.6, 1.3 Hz, 1H), 8.63 (s, 1H).</td>
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<tr>
<td>152 1-152</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 5.06 (s, 5H), 9.86-1.00 (m, 1H), 1.27-1.39 (m, 2H), 1.64 (d, J = 13.3 Hz, 2H), 2.22 (s, 3H), 2.58 (d, J = 10.6 Hz, 2H), 3.07 (d, J = 10.3 Hz, 2H), 3.96 (t, J = 4.2 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (t, J = 7.7 Hz, 1H), 7.05-7.10 (m, 2H), 7.34 (dd, J = 8.1, 1.5 Hz, 1H), 7.50 (dd, J = 8.6, 2.0 Hz, 1H), 8.13 (d, J = 2.2 Hz, 1H), 8.88 (s, 1H).</td>
</tr>
<tr>
<td>153 1-153</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 3.72 (q, J = 5.1 Hz, 2H), 3.92 (t, J = 4.2 Hz, 2H), 4.04 (t, J = 5.1 Hz, 2H), 4.34 (t, J = 4.2 Hz, 2H), 4.89 (t, J = 5.1 Hz, 1H), 6.89 (d, J = 7.9 Hz, 1H), 7.01-7.13 (m, 1H), 7.19 (d, J = 8.8 Hz, 1H), 7.31-7.32 (m, 1H), 7.42 (dd, J = 8.8, 3.2 Hz, 1H), 7.71 (dd, J = 8.3 Hz, 2H), 7.96 (d, J = 8.3 Hz, 2H), 8.08 (d, J = 2.8 Hz, 1H), 10.49 (s, 1H).</td>
</tr>
<tr>
<td>154 1-154</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 2.77 (t, J = 6.4 Hz, 2H), 3.67 (q, J = 5.9 Hz, 2H), 3.77 (q, J = 4.4 Hz, 2H), 4.43 (t, J = 4.2 Hz, 2H), 4.72 (t, J = 5.1 Hz, 1H), 6.47 (dd, J = 8.1, 1.8 Hz, 1H), 6.80 (t, J = 7.9 Hz, 1H), 7.04 (dd, J = 7.7, 1.5 Hz, 1H), 7.09-7.12 (m, 3H), 7.96 (dd, J = 9.9, 5.1 Hz, 3H), 8.30 (d, J = 1.8 Hz, 1H), 10.50 (s, 1H).</td>
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<tr>
<td>155 1-155</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 5.1205(s, 5H), 2.318s, 3H), 3.65(s, 2H), 4.04 (d, J=4.4Hz, 2H), 4.37(t, J=4.4Hz, 2H), 6.72 (m, 3H), 7.28(m, 2H), 7.30 (dd, J = 8.1, 1.8 Hz, 1H), 7.63 (dd, J = 9.0, 2.4 Hz, 1H), 7.72 (dd, J = 9.0, 2.4 Hz, 2H), 8.15 (d, J = 2.6 Hz, 1H), 10.00 (s, 1H).</td>
</tr>
<tr>
<td>156 1-156</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 5.1.50-1.51 (m, 2H), 1.60-1.66 (m, 4H), 2.21 (s, 3H), 2.89-2.90 (m, 4H), 3.97 (t, J = 4.4 Hz, 2H), 4.30 (t, J = 4.4 Hz, 2H), 6.89 (t, J = 7.9 Hz, 1H), 6.99 (d, J = 9.5 Hz, 1H), 7.12-7.14 (m, 2H), 7.38 (dd, J = 13.6, 8.2, 1.7 Hz, 2H), 7.51 (dd, J = 8.8, 2.3 Hz, 1H), 7.63 (dd, J = 14.8, 2.3 Hz, 1H), 8.13-8.14 (m, 1H), 10.11 (s, 1H).</td>
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<td>Ex. No.</td>
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<td>157 1-157</td>
<td><img src="image1.png" alt="Image" /></td>
<td><strong>(400 MHz, DMSO-d6)</strong> δ: 2.21 (s, 3H), 2.90-2.96 (m, 4H), 3.71-3.72 (m, 4H), 3.96 (t, J = 20.4 Hz, 2H), 4.30 (t, J = 4.2 Hz, 2H), 8.89 (dd, J = 5.3, 2.6 Hz, 1H), 7.00 (d, J = 9.5 Hz, 1H), 7.11-7.14 (m, 2H), 7.36-7.41 (m, 2H), 7.51 (d, J = 4.2, 2.0 Hz, 1H), 7.65 (dd, J = 15.1, 2.1 Hz, 1H), 8.13-8.14 (m, 1H), 10.15 (s, 1H).</td>
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<td>158 1-158</td>
<td><img src="image2.png" alt="Image" /></td>
<td><strong>(400 MHz, DMSO-d6)</strong> δ: 2.21 (s, 3H), 3.91 (t, J = 4.4 Hz, 2H), 4.30 (t, J = 4.4 Hz, 2H), 6.50 (dd, J = 11.1, 4.6 Hz, 1H), 7.13 (dd, J = 6.0, 4.3 Hz, 2H), 7.36-7.53 (m, 4H), 7.69-7.92 (m, 1H), 8.13-8.14 (m, 1H), 10.35 (s, 1H).</td>
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<td>159 1-159</td>
<td><img src="image3.png" alt="Image" /></td>
<td><strong>(400 MHz, DMSO-d6)</strong> δ: 2.21 (s, 3H), 3.80 (s, 3H), 3.97 (t, J = 4.4 Hz, 2H), 6.50 (dd, J = 11.1, 4.6 Hz, 1H), 7.15-7.35 (m, 3H), 7.41 (d, J = 4.6 Hz, 2H), 7.51 (dd, J = 8.3, 2.3 Hz, 1H), 7.70 (dd, J = 13.7, 2.6 Hz, 1H), 8.13 (s, 1H), 10.14 (s, 1H).</td>
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<td>160 1-160</td>
<td><img src="image4.png" alt="Image" /></td>
<td><strong>(400 MHz, DMSO-d6)</strong> δ: 2.12 (t, J = 7.0 Hz, 3H), 2.25 (d, J = 10.6 Hz, 3H), 3.95-4.03 (m, 4H), 4.32 (t, J = 4.4 Hz, 2H), 6.86-6.93 (m, 3H), 7.16 (d, J = 9.4, 3.8 Hz, 2H), 7.41 (dd, J = 8.1, 1.8 Hz, 1H), 7.52 (d, J = 9.0, 2.4 Hz, 1H), 7.63 (dd, J = 9.9, 2.8 Hz, 2H), 8.15 (d, J = 2.6 Hz, 1H), 9.98 (s, 1H).</td>
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<td>161 1-161</td>
<td><img src="image5.png" alt="Image" /></td>
<td><strong>(300 MHz, DMSO-d6)</strong> δ: 1.83-1.86 (m, 4H), 2.23 (s, 3H), 3.56-3.60 (m, 2H), 3.96-4.01 (m, 2H), 4.31-4.34 (m, 2H), 5.81-5.94 (m, 1H), 7.12-7.23 (m, 4H), 7.47 (dd, J = 31.2, 7.7 Hz, 2H), 7.72 (d, J = 8.6 Hz, 2H), 8.15 (s, 1H), 10.16 (s, 1H).</td>
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<tr>
<td>162 1-162</td>
<td><img src="image6.png" alt="Image" /></td>
<td><strong>(300 MHz, DMSO-d6)</strong> δ: 1.83-1.91 (m, 2H), 2.23 (s, 3H), 2.63-2.85 (m, 2H), 2.80 (s, 3H), 3.16-3.18 (m, 2H), 3.98 (t, J = 4.4 Hz, 2H), 4.32 (t, J = 4.2 Hz, 2H), 6.81 (d, J = 8.1 Hz, 1H), 6.90 (dd, J = 9.2, 6.6 Hz, 2H), 7.02-7.06 (m, 1H), 7.14-7.17 (m, 2H), 7.40 (d, J = 8.4 Hz, 1H), 7.52 (d, J = 8.4 Hz, 1H), 8.14-8.15 (m, 1H), 8.63 (s, 1H).</td>
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<tr>
<td>163 1-163</td>
<td><img src="image7.png" alt="Image" /></td>
<td><strong>(300 MHz, DMSO-d6)</strong> δ: 2.23 (s, 3H), 2.73 (s, 6H), 3.99 (t, J = 4.2 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 6.90-6.95 (m, 2H), 7.13-7.17 (m, 2H), 7.27-7.43 (m, 2H), 7.62 (dd, J = 8.4, 2.2 Hz, 1H), 7.63 (dd, J = 15.4, 2.2 Hz, 1H), 8.15 (d, J = 1.8 Hz, 1H), 10.08 (s, 1H).</td>
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<td>164 1-164</td>
<td><img src="image8.png" alt="Image" /></td>
<td><strong>(300 MHz, DMSO-d6)</strong> δ: 1.09-1.15 (m, 7H), 2.28 (s, 3H), 3.11 (s, 3H), 3.38-3.41 (m, 2H), 3.56-3.59 (m, 1H), 4.01-4.05 (m, 2H), 4.37-4.39 (m, 2H), 6.98 (t, J = 7.9 Hz, 1H), 7.27 (dd, J = 18.5, 7.9 Hz, 2H), 7.46 (d, J = 8.1 Hz, 1H), 7.54-7.56 (m, 1H), 7.75 (d, J = 8.8 Hz, 1H), 7.88 (d, J = 13.2 Hz, 1H), 8.17-8.21 (m, 1H), 10.52 (s, 1H).</td>
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<td>165 1-165</td>
<td><img src="image9.png" alt="Image" /></td>
<td><strong>(300 MHz, DMSO-d6)</strong> δ: 1.05-1.07 (m, 6H), 2.26 (s, 3H), 2.26 (s, 3H), 3.31 (s, 1H), 3.95-4.00 (m, 3H), 4.27-4.42 (m, 2H), 6.93 (t, J = 7.9 Hz, 1H), 7.19-7.23 (m, 2H), 7.39-7.44 (m, 2H), 7.63-7.69 (m, 2H), 8.17-8.19 (m, 1H), 10.21 (s, 1H).</td>
</tr>
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</table>
### [Table 22]

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<tr>
<th>Ex. No.</th>
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<th>NMR</th>
</tr>
</thead>
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<tr>
<td>166</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ 1.17-1.20 (m, 6H), 2.26 (s, 3H), 3.45-3.50 (m, 4H), 3.77-3.80 (m, 1H), 3.99-4.04 (m, 2H), 4.35 (t, J = 4.4 Hz, 2H), 6.95 (t, J = 7.9 Hz, 1H), 7.22 (dt, J = 10.5, 4.7 Hz, 2H), 7.46 (dd, J = 8.1, 1.5 Hz, 1H), 7.57-7.66 (m, 3H), 7.69 (d, J = 13.8 Hz, 1H), 8.18 (d, J = 1.1 Hz, 1H), 10.52 (s, 1H).</td>
</tr>
<tr>
<td>167</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ 1.23 (t, J = 5.1 Hz, 3H), 1.66-1.79 (m, 2H), 1.91-1.96 (m, 2H), 2.25 (s, 3H), 2.66-2.72 (m, 2H), 3.24-3.33 (m, 1H), 3.99 (t, J = 4.8 Hz, 2H), 4.09 (a, J = 26.4 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 4.60 (t, J = 7.9 Hz, 1H), 7.01 (t, J = 9.4 Hz, 1H), 7.13-7.18 (m, 2H), 7.40 (dd, J = 8.1, 4.0 Hz, 2H), 7.52 (dt, J = 8.4, 1.3 Hz, 1H), 7.65 (dd, J = 14.9, 2.4 Hz, 1H), 8.15 (t, J = 1.3 Hz, 1H), 10.12 (s, 1H).</td>
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<tr>
<td>168</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ 1.61-1.75 (m, 2H), 1.91-1.94 (m, 2H), 2.23 (s, 3H), 2.33-2.38 (m, 1H), 2.67-2.71 (m, 2H), 3.13-3.25 (m, 2H), 3.99 (t, J = 4.2 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 6.90 (t, J = 7.9 Hz, 1H), 7.01 (t, J = 9.5 Hz, 1H), 7.17-7.17 (m, 2H), 7.40 (dd, J = 8.4, 1.7 Hz, 2H), 7.52 (dt, J = 9.5, 1.3 Hz, 1H), 7.65 (dd, J = 14.9, 2.4 Hz, 1H), 8.15 (t, J = 1.3 Hz, 1H), 10.12 (s, 1H).</td>
</tr>
<tr>
<td>169</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ 1.55-1.58 (m, 2H), 1.91-1.95 (m, 2H), 2.20 (s, 3H), 2.74 (dt, J = 15.0, 5.4 Hz, 2H), 3.13-3.18 (m, 2H), 3.31 (s, 3H), 3.97 (t, J = 4.4 Hz, 2H), 4.30 (t, J = 4.4 Hz, 2H), 6.97 (t, J = 7.9 Hz, 1H), 7.00 (t, J = 9.5 Hz, 1H), 7.13 (dt, J = 11.3, 5.0 Hz, 2H), 7.30 (dd, J = 13.9, 8.3, 1.9 Hz, 2H), 7.51 (dd, J = 8.3, 2.3 Hz, 1H), 7.73 (dd, J = 14.8, 2.3 Hz, 1H), 8.14 (d, J = 2.3 Hz, 1H), 10.12 (s, 1H).</td>
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<td>170</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ 1.60-1.73 (m, 2H), 1.93-1.98 (m, 2H), 2.02 (s, 3H), 2.22 (s, 3H), 2.82-2.88 (m, 2H), 3.13-3.19 (m, 2H), 3.97 (t, J = 4.4 Hz, 2H), 4.30 (t, J = 4.4 Hz, 2H), 4.77-4.81 (m, 1H), 6.89 (t, J = 7.9 Hz, 1H), 7.03 (t, J = 9.3 Hz, 1H), 7.13 (dt, J = 11.1, 4.9 Hz, 2H), 7.39 (dd, J = 13.0, 4.9 Hz, 2H), 7.51 (dd, J = 8.3, 2.3 Hz, 1H), 7.65 (dd, J = 14.8, 2.3 Hz, 1H), 8.14 (d, J = 1.2 Hz, 1H), 10.13 (s, 1H).</td>
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<td>171</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ 1.50-1.57 (m, 2H), 1.84-1.85 (m, 2H), 2.25 (s, 3H), 2.70-2.74 (m, 2H), 3.18-3.20 (m, 2H), 3.59-3.62 (m, 1H), 3.99 (t, J = 4.2 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 4.66 (d, J = 4.4 Hz, 1H), 6.89 (t, J = 7.9 Hz, 1H), 7.01 (t, J = 5.4 Hz, 1H), 7.13-7.16 (m, 2H), 7.35-7.43 (m, 2H), 7.52 (dd, J = 8.4, 2.6 Hz, 1H), 7.64 (dd, J = 15.0, 2.2 Hz, 1H), 8.15 (d, J = 2.6 Hz, 1H), 10.11 (s, 1H).</td>
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<td>172</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ 2.11 (s, 3H), 2.23 (s, 3H), 3.69 (t, J = 4.2 Hz, 2H), 4.32 (t, J = 4.2 Hz, 2H), 6.91 (t, J = 7.9 Hz, 1H), 7.06 (d, J = 7.9 Hz, 1H), 7.14 (d, J = 8.3 Hz, 1H), 7.17-7.20 (m, 3H), 7.41 (d, J = 7.9 Hz, 1H), 7.48 (s, 1H), 7.53 (dd, J = 8.3, 2.3 Hz, 1H), 8.15 (s, 1H), 10.06 (s, 1H).</td>
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<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ 2.13 (s, 3H), 2.23 (s, 3H), 3.75 (t, J = 5.3 Hz, 2H), 3.97 (dt, J = 15.3, 4.8 Hz, 4H), 4.33 (t, J = 4.4 Hz, 2H), 4.83 (t, J = 5.6 Hz, 1H), 6.91 (t, J = 7.9 Hz, 1H), 7.06 (d, J = 8.3 Hz, 1H), 7.16 (dd, J = 18.6, 7.9 Hz, 3H), 7.43 (t, J = 9.0 Hz, 2H), 7.53 (dd, J = 8.8, 1.9 Hz, 1H), 8.15 (s, 1H), 10.03 (s, 1H).</td>
</tr>
</tbody>
</table>
## Table 23

<table>
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<tr>
<th>Ex. No.</th>
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</tr>
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<tr>
<td>174</td>
<td><img src="image" alt="Chemical 174" /></td>
<td>(400 MHz, DMSO-d6) 1.55 (br s, 2H), 1.67-1.90 (m, 2H), 2.23 (s, 3H), 2.54 (s, 6H), 3.17-3.19 (m, 2H), 3.63 (br s, 2H), 3.99 (t, J = 4.4 Hz, 2H), 4.31 (t, J = 4.4 Hz, 2H), 4.47 (s, 1H), 6.91 (t, J = 7.9 Hz, 1H), 7.15-7.21 (m, 3H), 7.40-7.43 (m, 2H), 7.51-7.54 (m, 1H), 7.73 (dd, J = 13.2, 2.1 Hz, 1H), 8.15 (d, J = 1.9 Hz, 1H), 10.18 (s, 1H).</td>
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<tr>
<td>175</td>
<td><img src="image" alt="Chemical 175" /></td>
<td>(400 MHz, DMSO-d6) 1.66-1.91 (m, 3H), 2.08-2.13 (m, 2H), 2.29 (s, 3H), 3.06 (brs, 2H), 3.20 (br s, 2H), 4.08 (t, J = 4.2 Hz, 2H), 4.38 (t, J = 4.2 Hz, 2H), 4.55-4.59 (m, 1H), 6.06 (l, J = 7.9 Hz, 1H), 7.25 (dd, J = 12.3, 5.0 Hz, 2H), 7.73 (d, J = 8.8 Hz, 1H), 7.45 (dd, J = 10.7, 3.7 Hz, 2H), 7.75-7.80 (m, 2H), 8.20 (s, 1H), 8.12 (d, J = 32.5 Hz, 2H), 10.25 (s, 1H).</td>
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<td>(400 MHz, DMSO-d6) 1.19 (t, J = 7.2 Hz, 3H), 1.25-1.31 (m, 4H), 1.85-1.89 (m, 4H), 2.25 (s, 3H), 2.36-3.34 (m, 1H), 3.70-3.73 (m, 1H), 3.95 (t, J = 4.4 Hz, 2H), 4.07-4.13 (m, 4H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (l, J = 7.9 Hz, 1H), 7.07 (d, J = 8.3 Hz, 1H), 7.17 (dd, J = 7.6, 1.4 Hz, 1H), 7.33 (dd, J = 8.1, 1.6 Hz, 1H), 7.48 (d, J = 8.3, 1.9 Hz, 1H), 7.90 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 2.3 Hz, 1H).</td>
</tr>
<tr>
<td>177</td>
<td><img src="image" alt="Chemical 177" /></td>
<td>(400 MHz, DMSO-d6) 1.21-1.35 (m, 4H), 1.85-1.99 (m, 4H), 2.20 (s, 3H), 3.22-3.36 (m, 1H), 3.69-3.74 (m, 1H), 3.95 (t, J = 4.2 Hz, 2H), 4.00 (t, J = 4.2 Hz), 4.27 (t, J = 4.2 Hz, 2H), 6.84 (l, J = 7.9 Hz, 1H), 7.07 (d, J = 8.3 Hz, 1H), 7.18 (d, J = 7.4 Hz, 1H), 7.33 (dd, J = 8.1, 1.6 Hz, 1H), 7.48 (d, J = 8.8 Hz, 1H), 7.90 (d, J = 7.4 Hz, 1H), 8.12 (s, 1H), 12.46 (brs, 1H).</td>
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<td><img src="image" alt="Chemical 178" /></td>
<td>(400 MHz, DMSO-d6) 1.05-1.18 (m, 4H), 1.20-1.24 (m, 4H), 1.99 (brs, 2H), 2.25 (s, 3H), 2.36-3.30 (m, 1H), 3.70-3.74 (m, 1H), 3.75 (s, 2H), 3.85 (brs, 2H), 3.85 (t, J = 7.9 Hz, 1H), 7.04 (brs, 1H), 7.07 (d, J = 8.3 Hz, 1H), 7.17 (d, J = 7.4 Hz, 1H), 7.22 (brs, 1H), 7.33 (d, J = 8.3 Hz, 1H), 7.48 (d, J = 8.3, 1.9 Hz, 1H), 7.91 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 2.3 Hz, 1H).</td>
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<tr>
<td>179</td>
<td><img src="image" alt="Chemical 179" /></td>
<td>(400 MHz, DMSO-d6) 1.83-1.99 (m, 4H), 1.88 (brs, 2H), 1.99 (brs, 2H), 2.20 (s, 3H), 2.61 (d, J = 4.6 Hz, 3H), 3.28-3.39 (m, 1H), 3.71-3.74 (m, 1H), 3.84 (s, 2H), 3.85 (t, J = 4.4 Hz, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (l, J = 7.9 Hz, 1H), 7.07 (d, J = 8.3 Hz, 1H), 7.17 (dd, J = 7.6, 1.6 Hz, 1H), 7.33 (dd, J = 8.1, 1.6 Hz, 1H), 7.48 (dd, J = 8.3, 2.3 Hz, 1H), 7.53 (brs, 1H), 7.92 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 2.3 Hz, 1H).</td>
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<td>180</td>
<td><img src="image" alt="Chemical 180" /></td>
<td>(400 MHz, DMSO-d6) 1.25-1.31 (m, 4H), 3.05-4.00 (m, 1H), 2.20 (s, 3H), 2.79 (s, 3H), 2.92 (s, 3H), 3.26-3.29 (m, 1H), 3.70-3.74 (m, 1H), 3.95 (t, J = 4.4 Hz, 2H), 4.10 (s, 2H), 4.27 (t, J = 4.4 Hz, 2H), 6.84 (l, J = 7.9 Hz, 1H), 7.07 (d, J = 8.3 Hz, 1H), 7.17 (dd, J = 7.4, 1.4 Hz, 1H), 7.33 (dd, J = 8.3, 1.4 Hz, 1H), 7.48 (dd, J = 8.3, 2.3 Hz, 1H), 7.91 (d, J = 7.4 Hz, 1H), 8.11 (s, 1H).</td>
</tr>
<tr>
<td>181</td>
<td><img src="image" alt="Chemical 181" /></td>
<td>(400 MHz, DMSO-d6) 1.14-1.31 (m, 4H), 1.68-1.93 (m, 4H), 2.16 (s, 3H), 3.18-3.21 (m, 1H), 3.35-3.42 (m, 4H), 3.65-3.86 (m, 1H), 3.91 (t, J = 4.0 Hz, 2H), 4.23 (t, J = 4.0 Hz, 2H), 4.47 (t, J = 5.3 Hz, 1H), 6.78 (l, J = 7.9 Hz, 1H), 7.03 (d, J = 8.8 Hz, 1H), 7.13 (d, J = 7.9 Hz, 1H), 7.29 (d, J = 7.9 Hz, 1H), 7.44 (d, J = 8.3 Hz, 1H), 7.96 (d, J = 7.9 Hz, 1H), 8.07 (s, 1H).</td>
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<td>Ex. No.</td>
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<td>NMR</td>
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<tr>
<td><strong>182</strong></td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.96-1.06 (m, 2H), 1.20-1.30 (m, 2H), 1.45-1.52 (m, 1H), 1.72-1.75 (m, 2H), 1.86-1.91 (m, 2H), 2.21 (s, 3H), 3.14 (d, J = 6.5 Hz, 2H), 3.21 (s, 3H), 3.63-3.71 (m, 1H), 3.96 (d, J = 4.4 Hz, 2H), 4.28 (t, J = 4.4 Hz, 2H), 6.84 (d, J = 7.9 Hz, 1H), 7.08 (d, J = 8.8 Hz, 1H), 7.19 (dd, J = 7.7, 1.6 Hz, 1H), 7.34 (dd, J = 7.9, 1.4 Hz, 1H), 7.49 (dd, J = 8.8, 2.3 Hz, 1H), 7.90 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 1.9 Hz, 1H).</td>
</tr>
<tr>
<td><strong>183</strong></td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.94-1.03 (m, 2H), 1.08 (d, J = 6.5 Hz, 6H), 1.19-1.30 (m, 2H), 1.38-1.43 (m, 1H), 1.75-1.78 (m, 2H), 1.86-1.91 (m, 2H), 2.21 (s, 3H), 3.16 (d, J = 6.5 Hz, 2H), 3.44-3.50 (m, 1H), 3.63-3.72 (m, 1H), 3.96 (d, J = 4.4 Hz, 2H), 4.28 (t, J = 4.4 Hz, 2H), 6.84 (d, J = 7.9 Hz, 1H), 7.08 (d, J = 8.3 Hz, 1H), 7.20 (dd, J = 7.7, 1.6 Hz, 1H), 7.34 (dd, J = 8.1, 1.6 Hz, 1H), 7.49 (dd, J = 8.3, 2.3 Hz, 1H), 7.88 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 2.3 Hz, 1H).</td>
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<tr>
<td><strong>184</strong></td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.27-1.36 (m, 2H), 1.50-1.70 (m, 7H), 2.23 (s, 3H), 3.20 (d, J = 6.5 Hz, 2H), 3.22 (s, 3H), 3.69-4.04 (m, 3H), 4.35 (t, J = 4.4 Hz, 2H), 6.86 (d, J = 7.9 Hz, 1H), 7.16 (d, J = 8.6 Hz, 1H), 7.22 (dd, J = 7.7, 1.6 Hz, 1H), 7.37 (dd, J = 8.1, 1.6 Hz, 1H), 7.59 (d, J = 7.4 Hz, 1H), 8.01 (d, J = 7.9 Hz, 1H), 8.14 (s, 1H).</td>
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<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.90-1.37 (m, 14H), 1.75-1.91 (m, 4H), 2.21 (s, 3H), 3.11 (d, J = 6.5 Hz, 2H), 3.63-3.72 (m, 1H), 3.96 (d, J = 4.4 Hz, 2H), 4.28 (t, J = 4.4 Hz, 2H), 6.84 (d, J = 7.9 Hz, 1H), 7.08 (d, J = 8.3 Hz, 1H), 7.20 (dd, J = 7.7, 1.6 Hz, 1H), 7.34 (dd, J = 8.1, 1.6 Hz, 1H), 7.49 (dd, J = 8.3, 2.3 Hz, 1H), 7.88 (d, J = 7.9 Hz, 1H), 8.12 (d, J = 2.3 Hz, 1H).</td>
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<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.84-1.37 (m, 14H), 1.75-1.91 (m, 4H), 1.85-1.97 (m, 4H), 2.22 (s, 3H), 3.15 (m, 3H), 3.96 (d, J = 4.4 Hz, 2H), 4.28 (t, J = 4.4 Hz, 2H), 6.85 (d, J = 8Hz, 1H), 7.09 (d, J = 8.4 Hz, 1H), 7.19 (d, J = 8 Hz, 1H), 7.49 (d, J = 8.4 Hz, 1H), 7.91 (d, J = 8 Hz, 1H), 8.12 (s, 1H).</td>
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<tr>
<td><strong>187</strong></td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.86 (d, J = 8.4 Hz, 6H), 1.21-1.34 (m, 4H), 1.85-1.97 (m, 4H), 2.22 (s, 3H), 3.15 (m, 3H), 3.96 (d, J = 4.4 Hz, 2H), 4.28 (t, J = 4.4 Hz, 2H), 6.85 (d, J = 8Hz, 1H), 7.09 (d, J = 8 Hz, 1H), 7.28 (m, 2H), 7.38 (d, J = 8 Hz, 1H), 7.76 (d, J = 8 Hz, 1H), 7.94 (d, J = 8 Hz, 1H), 8.17 (s, 1H).</td>
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<td><strong>188</strong></td>
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<td>(400 MHz, DMSO-d6) 0.92 (s, 6H), 1.17-1.47 (m, 6H), 1.82-1.72 (m, 2H), 2.27 (s, 3H), 3.73 (m, 3H), 4.02 (d, J = 4.4 Hz, 2H), 4.37 (d, J = 4.4 Hz, 2H), 5.91 (d, J = 8 Hz, 1H), 7.28 (m, 2H), 7.38 (d, J = 8 Hz, 1H), 7.76 (d, J = 8 Hz, 1H), 7.94 (d, J = 8 Hz, 1H), 8.17 (s, 1H).</td>
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<td><strong>189</strong></td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.83 (d, J = 8.4 Hz, 6H), 1.24-1.4 (m, 6H), 1.8-2.0 (m, 4H), 2.20 (s, 3H), 3.20 (m, 1H), 3.41 (d, J = 8 Hz, 2H), 3.72 (m, 1H), 3.87 (s, J = 4.4 Hz, 2H), 4.29 (q, J = 4.4 Hz, 2H), 6.55 (d, J = 8 Hz, 1H), 7.09 (d, J = 8 Hz, 1H), 7.19 (d, J = 8 Hz, 1H), 7.34 (d, J = 8 Hz, 1H), 7.49 (d, J = 8 Hz, 1H), 7.91 (d, J = 8 Hz, 1H), 8.12 (s, 1H).</td>
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<td>Ex. No.</td>
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<tr>
<td>190</td>
<td><img src="Image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.3-1.4(m, 4H), 1.85-1.86(m, 2H), 2.0-2.1(m, 2H), 2.22(s, 3H), 3.34(m, 1H), 3.76(m, 1H), 3.96(t, J=4.6Hz, 2H), 4.29(t, J=4.6Hz, 2H), 4.65(s, 2H), 6.85(t, J=8Hz, 1H), 7.10(d, J=8Hz, 1H), 7.19(d, J=8Hz, 1H), 7.22-7.44(m, 6H), 7.50(d, J=8Hz, 1H), 7.94(d, J=8Hz, 1H), 8.13(s, 1H)</td>
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<td>191</td>
<td><img src="Image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.85(d, J=6.4Hz, 6H), 1.0-1.15(m, 3H), 1.2-1.35(m, 2H), 1.35-1.5(m, 1H), 1.6-1.75(m, 2H), 1.85-1.95(m, 2H), 2.22(s, 3H), 3.66(m, 1H), 3.97(t, J=4Hz, 2H), 4.29(t, J=4Hz, 2H), 6.85(t, J=8Hz, 1H), 7.09(d, J=8Hz, 1H), 7.20(d, J=8Hz, 1H), 7.34(d, J=8Hz, 1H), 7.49(d, J=8Hz, 1H), 7.86(d, J=8Hz, 1H), 8.13(s, 1H)</td>
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<td>192</td>
<td><img src="Image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 0.86(t, J=7.2Hz, 3H), 0.8-1.0(m, 2H), 1.1-1.35(m, 6H), 1.7-1.9(m, 2H), 1.8-1.9(m, 2H), 2.22(s, 3H), 3.7(m, 1H), 3.86(t, J=4.4Hz, 2H), 4.29(t, J=4.4Hz, 2H), 6.85(t, J=8Hz, 1H), 7.06(d, J=8Hz, 1H), 7.34(d, J=8Hz, 1H), 7.48(d, J=8Hz, 1H), 7.88(d, J=8Hz, 1H), 8.13(s, 1H)</td>
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<td>193</td>
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<td>(400 MHz, DMSO-d6) 0.86(s, 9H), 1.2-1.4(m, 4H), 1.8-2.0(m, 4H), 2.22(s, 3H), 3.07(s, 2H), 3.17(m, 1H), 3.73(m, 1H), 3.86(t, J=4Hz, 2H), 4.29(t, J=4Hz, 2H), 6.85(t, J=8Hz, 1H), 7.06(d, J=8Hz, 1H), 7.34(dd, J=8Hz, 1H), 7.48(dd, J=8Hz, 1H), 7.50(ddd, J=8, 2.4Hz, 1H), 7.92(d, J=8Hz, 1H), 8.13(s, 1H)</td>
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<tr>
<td>194</td>
<td><img src="Image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 1.28 (s, 9H), 3.29 (s, 3H), 4.06 (t, J = 4.4 Hz, 2H), 4.31-4.36 (m, 4H), 6.93 (t, J = 7.9 Hz, 1H), 7.18-7.25 (m, 2H), 7.34 (d, J = 8.8 Hz, 2H), 7.48 (dd, J = 8.1, 1.5 Hz, 1H), 7.60-7.86 (m, 3H), 8.25 (d, J = 2.2 Hz, 1H), 10.05 (s, 1H)</td>
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<td>195</td>
<td><img src="Image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 3.92 (t, J = 4.2 Hz, 2H), 4.43 (t, J = 4.4 Hz, 2H), 6.76 (dd, J = 8.3, 1.4 Hz, 1H), 6.86 (dd, J = 7.9 Hz, 1H), 7.17 (dd, J = 7.4, 1.4 Hz, 1H), 7.46 (dd, J = 7.9 Hz, 1H), 7.78 (dd, J = 9.0, 2.6 Hz, 1H), 8.14 (d, J = 2.8 Hz, 1H), 8.30 (d, J = 1.9 Hz, 1H), 8.83 (d, J = 1.9 Hz, 1H), 10.51 (s, 1H), 13.47 (br s, 1H)</td>
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<td>(300 MHz, DMSO-d6) 3.92 (t, J = 4.2 Hz, 2H), 4.43 (t, J = 4.0 Hz, 2H), 6.74 (dd, J = 8.1, 1.5 Hz, 1H), 6.85 (t, J = 7.9 Hz, 1H), 7.17 (dd, J = 7.7, 1.5 Hz, 1H), 7.35 (d, J = 8.8 Hz, 2H), 7.66 (d, J = 8.8 Hz, 2H), 8.30 (d, J = 2.2 Hz, 1H), 8.83 (d, J = 2.2 Hz, 1H), 10.35 (s, 1H), 13.46 (br s, 1H)</td>
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<td>197</td>
<td><img src="Image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 2.70 (d, J = 5.1 Hz, 3H), 3.78 (l, J = 4.2 Hz, 2H), 4.33 (l, J = 4.4 Hz, 2H), 6.69 (q, J = 5.1 Hz, 1H), 8.02 (l, J = 7.9 Hz, 1H), 6.97-7.06 (m, 4H), 7.34 (d, J = 8.8 Hz, 2H), 7.75 (d, J = 2.8 Hz, 1H), 7.84 (d, J = 8.8 Hz, 2H), 10.35 (s, 1H)</td>
</tr>
</tbody>
</table>
### Table 26

<table>
<thead>
<tr>
<th>Ex. No.</th>
<th>Chemical Compounds</th>
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</tr>
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<tr>
<td>198 1-198</td>
<td><img src="image1" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 6: 1.34 (t, J = 7.5 Hz, 3H), 3.91 (t, J = 4.4 Hz, 2H), 4.08 (q, J = 7.1 Hz, 2H), 4.33 (t, J = 4.2 Hz, 2H), 6.88 (t, J = 7.9 Hz, 1H), 7.12-7.17 (m, 2H), 7.26-7.41 (m, 4H), 7.85 (d, J = 9.2 Hz, 2H), 8.06 (d, J = 3.3 Hz, 1H), 10.31 (s, 1H).</td>
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<tr>
<td>199 1-199</td>
<td><img src="image2" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 3.71 (t, J = 4.4 Hz, 2H), 3.69 (s, 3H), 4.43 (t, J = 4.4 Hz, 2H), 5.35 (dd, J = 7.9, 1.4 Hz, 1H), 6.77 (t, J = 7.9 Hz, 1H), 6.98 (dd, J = 7.7, 1.6 Hz, 1H), 7.35 (dd, J = 8.8 Hz, 2H), 7.77 (d, J = 3.2 Hz, 1H), 7.86 (d, J = 8.8 Hz, 2H), 8.22 (d, J = 2.8 Hz, 1H), 10.33 (s, 1H).</td>
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<td>200 1-200</td>
<td><img src="image3" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 3.72 (c, J = 5.1 Hz, 2H), 3.91 (t, J = 4.4 Hz, 2H), 4.04 (t, J = 5.1 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 4.89 (t, J = 5.1 Hz, 1H), 6.88 (t, J = 7.9 Hz, 1H), 7.11 (dd, J = 7.9, 1.6 Hz, 1H), 7.18 (dd, J = 8.8 Hz, 1H), 7.31 (dd, J = 7.9, 1.5 Hz, 1H), 7.35 (dd, J = 8.8 Hz, 2H), 7.41 (dd, J = 8.8, 3.0 Hz, 1H), 7.85 (d, J = 8.8 Hz, 2H), 8.08 (d, J = 3.0 Hz, 1H), 10.32 (s, 1H).</td>
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<td>201 1-201</td>
<td><img src="image4" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 3.85 (t, J = 4.2 Hz, 2H), 4.33 (t, J = 4.2 Hz, 2H), 6.85 (t, J = 7.9 Hz, 1H), 7.07-7.10 (m, 2H), 7.20-7.21 (m, 2H), 7.34 (d, J = 8.8 Hz, 2H), 7.85 (d, J = 9.2 Hz, 2H), 7.93 (d, J = 2.9 Hz, 1H), 10.30 (s, 1H).</td>
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<td>202 1-202</td>
<td><img src="image5" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) 3.81 (s, 3H), 3.91 (t, J = 4.6 Hz, 2H), 4.33 (t, J = 4.2 Hz, 2H), 6.88 (t, J = 7.9 Hz, 1H), 7.12 (dd, J = 7.7, 1.5 Hz, 1H), 7.19 (dd, J = 8.8 Hz, 1H), 7.28-7.43 (m, 4H), 7.85 (d, J = 9.2 Hz, 2H), 8.08 (d, J = 3.3 Hz, 1H), 10.31 (s, 1H).</td>
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<tr>
<td>203 1-203</td>
<td><img src="image6" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 3.95 (t, J = 4.2 Hz, 2H), 4.39 (t, J = 4.4 Hz, 2H), 6.86 (t, J = 7.9 Hz, 1H), 7.05 (dd, J = 7.9, 1.4 Hz, 1H), 7.18 (dd, J = 7.7, 1.6 Hz, 1H), 7.27 (dd, J = 7.9, 5.1 Hz, 1H), 7.34 (dd, J = 8.8 Hz, 2H), 7.54 (d, J = 8.9 Hz, 2H), 8.29 (dd, J = 7.9, 1.9 Hz, 1H), 8.59 (dd, J = 4.4, 2.0 Hz, 1H), 10.34 (s, 1H).</td>
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<td><img src="image7" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 3.78 (t, J = 4.2 Hz, 2H), 4.43 (t, J = 4.2 Hz, 2H), 6.74 (t, J = 7.9 Hz, 1H), 8.81 (dd, J = 8.1, 1.6 Hz, 1H), 7.55 (dd, J = 7.4, 1.6 Hz, 1H), 7.24 (dd, J = 7.7, 4.6 Hz, 1H), 7.34 (d, J = 8.9 Hz, 2H), 7.43 (s, 1H), 7.84 (q, J = 8.1 Hz, 3H), 7.92 (dd, J = 7.4, 1.9 Hz, 1H), 8.45 (dd, J = 4.4, 2.0 Hz, 1H), 10.30 (s, 1H).</td>
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<td><img src="image8" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 2.55 (d, J = 4.6 Hz, 3H), 3.78 (t, J = 4.2 Hz, 2H), 4.41 (t, J = 4.2 Hz, 2H), 6.74 (t, J = 7.9 Hz, 1H), 6.83 (dd, J = 8.1, 1.6 Hz, 1H), 7.09 (dd, J = 7.7, 1.6 Hz, 1H), 7.20 (dd, J = 7.4, 4.6 Hz, 1H), 7.34 (d, J = 8.8 Hz, 2H), 7.83-7.87 (m, 3H), 8.27-8.30 (m, 1H), 8.43 (dd, J = 4.8, 2.0 Hz, 1H), 10.27 (s, 1H).</td>
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<td>206</td>
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<td>(400 MHz, DMSO-d6) 2.81 (s, 3H), 2.86 (s, 3H), 3.71 (s, 2H), 4.34 (s, 2H), 6.80 (t, J = 7.9 Hz, 1H), 6.91 (dd, J = 8.1, 1.6 Hz, 1H), 7.11 (dd, J = 7.4, 1.4 Hz, 1H), 7.21 (dd, J = 7.4, 4.6 Hz, 1H), 7.34 (d, J = 8.3 Hz, 2H), 7.76 (dd, J = 7.4, 1.9 Hz, 1H), 7.85 (d, J = 9.3 Hz, 2H), 8.41 (dd, J = 5.2, 2.0 Hz, 1H), 10.28 (s, 1H).</td>
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<td>(400 MHz, DMSO-d6) 3.64 (t, J = 4.2 Hz, 2H), 4.25 (t, J = 4.2 Hz, 2H), 4.93 (s, 2H), 6.77 (t, J = 7.9 Hz, 1H), 8.87 (dd, J = 8.1, 1.6 Hz, 1H), 7.16 (dd, J = 7.4, 1.4 Hz, 1H), 7.21 (dd, J = 8.0, 4.8 Hz, 1H), 7.28-7.37 (m, 7H), 7.86 (d, J = 8.8 Hz, 2H), 8.12 (dd, J = 7.9, 1.9 Hz, 1H), 8.53 (dd, J = 5.2, 2.4 Hz, 1H), 10.26 (s, 1H).</td>
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<td>(400 MHz, DMSO-d6) 3.82 (t, J = 4.2 Hz, 2H), 4.39 (t, J = 4.2 Hz, 2H), 6.77 (t, J = 7.9 Hz, 1H), 8.87 (dd, J = 8.3, 1.4 Hz, 1H), 7.08 (dd, J = 7.7, 1.6 Hz, 1H), 7.20 (dd, J = 7.4, 4.6 Hz, 1H), 7.34 (d, J = 8.3 Hz, 2H), 7.86 (d, J = 9.3 Hz, 2H), 8.10 (dd, J = 7.9, 1.9 Hz, 1H), 8.46 (dd, J = 4.4, 2.0 Hz, 1H), 10.29 (s, 1H), 13.28 (d, J = 155.4 Hz, 1H).</td>
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<td>(400 MHz, DMSO-d6) 3.79 (t, J = 4.2 Hz, 2H), 4.38 (t, J = 4.4 Hz, 2H), 5.15 (s, 2H), 6.61 (dd, J = 8.3, 1.4 Hz, 1H), 6.76 (t, J = 7.9 Hz, 1H), 7.03 (dd, J = 7.4, 1.4 Hz, 1H), 7.21 (dd, J = 8.0, 4.8 Hz, 1H), 7.27-7.35 (m, 7H), 7.61 (dd, J = 7.9, 1.4 Hz, 1H), 7.85 (d, J = 8.8 Hz, 2H), 8.00 (dd, J = 4.9, 1.8 Hz, 1H), 10.29 (s, 1H), 10.29 (s, 1H).</td>
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<td>210</td>
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<td>(400 MHz, DMSO-d6) 3.75 (t, J = 4.4 Hz, 2H), 4.41 (t, J = 4.2 Hz, 2H), 6.50 (t, J = 3.9 Hz, 1H), 6.76 (t, J = 7.9 Hz, 1H), 6.97 (d, J = 6.5 Hz, 1H), 7.10 (dd, J = 8.0, 4.8 Hz, 1H), 7.29 (dd, J = 7.9, 1.4 Hz, 1H), 7.34 (d, J = 8.8 Hz, 2H), 7.85 (d, J = 8.8 Hz, 2H), 7.90 (dd, J = 4.4, 1.2 Hz, 1H), 9.97 (s, 1H), 10.31 (s, 1H).</td>
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<td>211</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, CHLOROFORM-d6) 3.70 (brs, 2H), 4.15 (brs, 2H), 4.33 (brs, 2H), 4.65 (brs, 2H), 6.73 (d, J = 7.4 Hz, 1H), 6.94 (t, J = 7.4 Hz, 1H), 7.22 (d, J = 8.3 Hz, 2H), 7.34 (brs, 1H), 7.64 (d, J = 8.0 Hz, 1H), 7.73 (d, J = 8.8 Hz, 2H), 7.87 (d, J = 7.4 Hz, 1H), 8.12 (brs, 1H), 9.56 (s, 1H).</td>
</tr>
<tr>
<td>212</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) 3.81 (t, J = 4.4 Hz, 2H), 4.39 (t, J = 4.2 Hz, 2H), 4.79 (s, 2H), 6.70-6.77 (m, 2H), 7.00 (dd, J = 7.2, 2.1 Hz, 1H), 7.18 (dd, J = 8.0, 4.8 Hz, 1H), 7.34 (d, J = 8.3 Hz, 2H), 7.42 (dd, J = 8.3, 1.4 Hz, 1H), 7.85 (d, J = 9.3 Hz, 2H), 7.99 (dd, J = 4.6, 1.4 Hz, 1H), 10.32 (s, 1H), 13.12 (s, 1H).</td>
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<tr>
<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<tr>
<td>213 1-213</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, (DMSO-d6) 3.82 (t, J = 4.2 Hz, 2H), 4.39 (t, J = 4.2 Hz, 2H), 4.54 (s, 2H), 6.68 (dd, J = 8.1, 1.6 Hz, 1H), 6.76 (t, J = 7.9 Hz, 1H), 7.02 (dd, J = 7.4, 1.4 Hz, 1H), 7.09 (s, 1H), 7.18 (dd, J = 8.0, 4.8 Hz, 1H), 7.32-7.42 (m, 4H), 7.85 (d, J = 9.3 Hz, 2H), 7.99 (dd, J = 4.6, 1.4 Hz, 1H), 10.29 (s, 1H).</td>
</tr>
<tr>
<td>214 1-214</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, (DMSO-d6) 2.58 (d, J = 4.6 Hz, 3H), 3.83 (t, J = 4.2 Hz, 2H), 4.39 (t, J = 4.2 Hz, 2H), 4.58 (s, 2H), 6.99 (dd, J = 8.1, 1.6 Hz, 1H), 6.77 (t, J = 7.9 Hz, 1H), 7.03 (dd, J = 7.4, 1.4 Hz, 1H), 7.17 (dd, J = 8.4, 4.8 Hz, 1H), 7.33 (d, J = 8.8 Hz, 2H), 7.40-7.46 (m, 2H), 7.65 (d, J = 9.3 Hz, 2H), 8.00 (dd, J = 4.6, 1.4 Hz, 1H), 10.31 (s, 1H).</td>
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<tr>
<td>215 1-215</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, (DMSO-d6) 2.83 (s, 3H), 2.95 (s, 3H), 3.83 (s, 2H), 4.36 (s, 2H), 4.96 (s, 2H), 6.71-6.79 (m, 2H), 6.97-7.00 (m, 1H), 7.12-7.16 (m, 1H), 7.32-7.37 (m, 3H), 7.84 (d, J = 8.8 Hz, 2H), 7.93-7.95 (m, 1H), 10.30 (s, 1H).</td>
</tr>
<tr>
<td>216 1-216</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, (DMSO-d6) 3.64 (t, J = 4.1 Hz, 2H), 4.40 (t, J = 4.1 Hz, 2H), 5.31 (s, 2H), 6.65 (dd, J = 8.1, 1.5 Hz, 1H), 6.75 (t, J = 7.7 Hz, 1H), 7.02 (dd, J = 7.4, 1.5 Hz, 1H), 7.22 (dd, J = 8.0, 4.8 Hz, 1H), 7.33 (t, J = 6.3 Hz, 2H), 7.42-7.46 (m, 1H), 7.64 (dd, J = 8.0, 1.5 Hz, 1H), 7.84-7.91 (m, 3H), 8.03 (dd, J = 4.6, 1.4 Hz, 1H), 8.80 (d, J = 5.1 Hz, 1H), 10.31 (s, 1H).</td>
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<tr>
<td>217 1-217</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, (DMSO-d6) 3.78 (t, J = 4.3 Hz, 2H), 4.37 (t, J = 4.3 Hz, 2H), 5.23 (s, 2H), 6.60 (dd, J = 8.3, 1.5 Hz, 1H), 6.77 (t, J = 7.9 Hz, 1H), 7.02 (dd, J = 7.7, 1.5 Hz, 1H), 7.23 (dd, J = 8.0, 4.8 Hz, 1H), 7.32-7.36 (m, 3H), 7.64-7.80 (m, 2H), 7.85 (d, J = 9.3 Hz, 2H), 8.02 (dd, J = 4.8, 1.2 Hz, 1H), 8.50 (dd, J = 4.8, 2.0 Hz, 1H), 8.54 (d, J = 2.0 Hz, 1H), 10.26 (s, 1H).</td>
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<td>218 1-218</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, (DMSO-d6) &amp; H1-NMR (DMSO-d6) δ: 4.16 (t, J = 4.5 Hz, 2H), 4.37 (t, J = 4.5 Hz, 2H), 6.98 (t, J = 8.0 Hz, 1H), 7.28-7.35 (m, 4H), 7.51 (dd, J = 8.0, 1.6 Hz, 1H), 7.82 (d, J = 8.3 Hz, 2H), 8.22 (dd, J = 8.8, 2.3 Hz, 1H), 8.68 (d, J = 2.3 Hz, 1H), 10.33 (s, 1H).</td>
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<tr>
<td>219 1-219</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, (DMSO-d6) 5: 1.85 (s, 3H), 4.02 (t, J = 4.4 Hz, 2H), 4.19 (d, J = 5.9 Hz, 2H), 4.32 (t, J = 4.6 Hz, 2H), 6.93 (t, J = 7.9 Hz, 1H), 7.17-7.21 (m, 2H), 7.34 (dd, J = 8.4 Hz, 2H), 7.46 (dd, J = 8.3, 1.7 Hz, 1H), 7.58 (dd, J = 8.6, 2.4 Hz, 1H), 7.84 (d, J = 9.2 Hz, 2H), 8.20 (d, J = 2.2 Hz, 1H), 8.30 (d, J = 6.1 Hz, 1H), 10.32 (s, 1H).</td>
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<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<td>220</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 3.34 (s, 3H), 3.80 (t, J = 4.2 Hz, 2H), 4.42 (t, J = 4.2 Hz, 2H), 4.47 (s, 2H), 6.51 (dd, J = 8.1, 1.5 Hz, 1H), 6.80 (t, J = 7.7 Hz, 1H), 7.00 (dd, J = 7.7, 1.5 Hz, 1H), 7.35 (d, J = 8.4 Hz, 2H), 7.68 (d, J = 8.8 Hz, 2H), 8.38 (d, J = 1.8 Hz, 1H), 10.33 (s, 1H).</td>
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<td>221</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 2.05 (s, 2H), 2.10 (s, 1H), 2.79 (s, 1H), 2.82 (s, 1H), 2.94 (s, 2H), 4.03-4.04 (m, 2H), 4.32-4.35 (m, 2H), 4.43 (s, 1.5H, keto), 4.50 (s, 0.6H, enol), 5.91-5.94 (m, 1H), 7.16-7.25 (m, 2H), 7.34 (d, J = 8.8 Hz, 2H), 7.50-7.55 (m, 2H), 7.85 (d, J = 9.2 Hz, 2H), 8.18-8.22 (m, 1H), 10.32 (s, 1H).</td>
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<tr>
<td>222</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 3.66 (s, 2H), 4.01 (t, J = 4.2 Hz, 2H), 4.32 (t, J = 4.4 Hz, 2H), 6.92 (t, J = 7.9 Hz, 1H), 7.17 (d, J = 8.8 Hz, 2H), 7.34 (d, J = 8.3 Hz, 2H), 7.44 (dd, J = 8.3, 1.4 Hz, 1H), 7.66 (dd, J = 8.3, 2.3 Hz, 1H), 7.84 (d, J = 9.3 Hz, 2H), 8.23 (d, J = 1.9 Hz, 1H), 10.33 (s, 1H).</td>
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<td>223</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(300 MHz, DMSO-d6) δ: 2.13 (s, 6H), 3.33 (s, 2H), 4.03 (t, J = 4.4 Hz, 2H), 4.34 (t, J = 4.4 Hz, 2H), 6.61 (q, J = 8.7 Hz, 1H), 7.19 (d, J = 3.9 Hz, 2H), 7.35 (d, J = 8.4 Hz, 2H), 7.49 (dd, J = 8.1, 1.5 Hz, 1H), 7.60 (dd, J = 8.4, 2.2 Hz, 1H), 7.85 (d, J = 9.2 Hz, 2H), 8.18 (d, J = 1.8 Hz, 1H), 10.32 (s, 1H).</td>
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<td>224</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 2.65 (d, J = 4.6 Hz, 3H), 3.91-4.00 (m, 2H), 5.12 (d, J = 4.6 Hz, 1H), 6.54 (dd, J = 8.1, 1.5 Hz, 1H), 6.89 (t, J = 7.9 Hz, 1H), 7.24 (dd, J = 7.7, 1.5 Hz, 1H), 7.30 (dd, J = 7.7, 4.6 Hz, 1H), 7.38 (d, J = 8.3 Hz, 2H), 7.39 (d, J = 8.5 Hz, 2H), 8.01 (dd, J = 7.9, 1.5 Hz, 1H), 8.15 (q, J = 4.6 Hz, 1H), 8.43 (dd, J = 4.8, 2.0 Hz, 1H), 10.56 (s, 1H).</td>
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<td>225</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 4.05 (t, J = 4.4 Hz, 2H), 4.33 (t, J = 4.4 Hz, 2H), 4.48 (d, J = 5.6 Hz, 2H), 5.33 (t, J = 5.6 Hz, 1H), 6.94 (t, J = 7.9 Hz, 1H), 7.02 (d, J = 7.9 Hz, 1H), 7.07 (d, J = 7.9 Hz, 1H), 7.21 (dd, J = 7.7, 1.6 Hz, 1H), 7.51 (dd, J = 7.7, 1.6 Hz, 1H), 7.57 (d, J = 7.7 Hz, 1H), 7.71 (d, J = 8.3 Hz, 2H), 7.95 (d, J = 8.3 Hz, 2H), 10.51 (s, 1H).</td>
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<tr>
<td>226</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 2.88 (s, 3H), 3.05 (s, 3H), 3.58 (dd, J = 13.0, 8.6 Hz, 1H), 4.20 (dd, J = 13.0, 4.0 Hz, 1H), 5.94 (dd, J = 8.6, 4.0 Hz, 1H), 6.56 (dd, J = 8.0, 1.4 Hz, 1H), 6.89 (t, J = 7.9 Hz, 1H), 7.24 (dd, J = 7.9, 1.4 Hz, 1H), 7.31 (dd, J = 8.0, 4.8 Hz, 1H), 7.37 (d, J = 8.3 Hz, 2H), 7.88 (d, J = 8.3 Hz, 2H), 8.03 (dd, J = 8.0, 1.4 Hz, 1H), 8.44 (dd, J = 4.8, 1.4 Hz, 1H), 10.30 (s, 1H).</td>
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<td>227</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d6) δ: 3.88 (s, 3H), 4.22 (t, J = 4.2 Hz, 2H), 6.68 (dd, J = 7.9, 1.6 Hz, 1H), 6.84 (t, J = 7.9 Hz, 1H), 7.13 (dd, J = 7.9, 1.6 Hz, 1H), 7.36 (d, J = 8.3 Hz, 2H), 7.66 (s, 1H), 7.87 (d, J = 8.3 Hz, 2H), 8.17 (s, 1H), 8.38 (d, J = 2.3 Hz, 1H), 8.52 (d, J = 2.3 Hz, 1H), 10.36 (s, 1H).</td>
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<tr>
<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<td>228: 1-228</td>
<td><img src="image1" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) 1.80 (s, 3H), 3.23-3.29 (m, 1H), 3.45-3.51 (m, 1H), 4.05-4.06 (m, 1H), 4.17 (d, J = 10.5 Hz, 1H), 4.47 (d, J = 10.5 Hz, 1H), 6.48 (dd, J = 8.0, 1.2 Hz, 1H), 6.84 (t, J = 7.7 Hz, 1H), 7.07 (dd, J = 7.7, 1.2 Hz, 1H), 7.36 (dd, J = 8.0, 4.8 Hz, 1H), 7.72 (d, J = 8.3 Hz, 2H), 8.00 (d, J = 8.3 Hz, 2H), 8.08 (dd, J = 8.0, 1.2 Hz, 1H), 8.16 (t, J = 5.6 Hz, 1H), 8.46 (dd, J = 4.8, 1.2 Hz, 1H), 10.45 (s, 1H).</td>
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<td>229: 1-229</td>
<td><img src="image2" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) 4.41 (dd, J = 10.7, 2.8 Hz, 1H), 4.68-4.75 (m, 2H), 6.58 (d, J = 7.9 Hz, 1H), 6.84 (dd, J = 7.9 Hz, 1H), 7.05 (dd, J = 7.9, 1.5 Hz, 1H), 7.26 (dd, J = 8.0, 4.5 Hz, 1H), 7.69 (d, J = 6.0 Hz, 2H), 7.97-8.00 (m, 3H), 8.38 (dd, J = 4.5, 1.5 Hz, 1H), 10.53 (s, 1H), 13.10 (s, 1H).</td>
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<tr>
<td>230: 1-230</td>
<td><img src="image3" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) 3.63 (s, 3H), 5.50 (dd, J = 7.9, 1.4 Hz, 1H), 6.87 (t, J = 7.9 Hz, 1H), 7.23 (dd, J = 7.9, 1.4 Hz, 1H), 7.33 (dd, J = 8.0, 4.8 Hz, 1H), 7.38 (d, J = 8.5 Hz, 2H), 7.88 (d, J = 8.5 Hz, 2H), 8.04 (dd, J = 7.6, 4.8 Hz, 1H), 8.44 (dd, J = 4.8, 4.6 Hz, 1H), 10.33 (s, 1H).</td>
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<tr>
<td>231: 1-231</td>
<td><img src="image4" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) 3.67 (dd, J = 13.0, 7.4 Hz, 1H), 3.74 (t, J = 4.5 Hz, 2H), 3.92 (dd, J = 13.0, 2.8 Hz, 1H), 4.48-4.54 (m, 1H), 5.22 (t, J = 5.5 Hz, 1H), 6.56 (dd, J = 7.9, 1.5 Hz, 1H), 6.85 (d, J = 7.9 Hz, 1H), 7.27 (dd, J = 7.9, 1.5 Hz, 1H), 7.32 (dd, J = 7.9, 4.5 Hz, 1H), 7.38 (d, J = 8.8 Hz, 2H), 7.89 (d, J = 8.8 Hz, 2H), 8.05 (dd, J = 7.9, 1.9 Hz, 1H), 8.44 (dd, J = 4.5, 1.4 Hz, 1H), 10.40 (s, 1H).</td>
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<td>232: 1-232</td>
<td><img src="image5" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) 3.93-4.08 (m, 2H), 5.31 (t, J = 4.2 Hz, 1H), 6.53 (dd, J = 7.9, 1.4 Hz, 1H), 6.87 (t, J = 7.8 Hz, 1H), 7.27-7.34 (m, 2H), 7.38 (d, J = 8.3 Hz, 2H), 7.89 (d, J = 8.3 Hz, 2H), 8.03 (dd, J = 8.3, 1.4 Hz, 1H), 8.44 (dd, J = 4.8, 2.0 Hz, 1H), 10.45 (s, 1H), 13.59 (s, 1H).</td>
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<td>233: 1-233</td>
<td><img src="image6" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) 4.27 (dd, J = 10.7, 2.5 Hz, 1H), 4.56-4.60 (m, 1H), 4.74 (dd, J = 10.7, 2.5 Hz, 1H), 6.58 (dd, J = 7.9, 1.4 Hz, 1H), 6.65 (t, J = 7.9 Hz, 1H), 7.07 (dd, J = 7.4, 1.4 Hz, 1H), 7.33-7.37 (m, 2H), 7.59 (s, 1H), 7.79 (d, J = 8.8 Hz, 2H), 7.96 (d, J = 8.3 Hz, 2H), 8.06 (dd, J = 7.9, 1.4 Hz, 1H), 8.42 (dd, J = 4.6, 1.4 Hz, 1H), 10.53 (s, 1H).</td>
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<td>234: 1-234</td>
<td><img src="image7" alt="Chemical Structure" /></td>
<td>(400 MHz, DMSO-d$_6$) 2.60 (d, J = 4.6 Hz, 3H), 4.26 (d, J = 8.5 Hz, 1H), 4.60 (s, 1H), 4.74 (d, J = 8.5 Hz, 1H), 6.80 (d, J = 7.9 Hz, 1H), 6.86 (t, J = 7.9 Hz, 1H), 7.08 (d, J = 6.5 Hz, 1H), 7.33 (q, J = 4.2 Hz, 1H), 7.70 (d, J = 8.8 Hz, 2H), 7.95-8.06 (m, 4H), 8.42 (dd, J = 4.6, 1.4 Hz, 1H), 10.52 (s, 1H).</td>
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<td>Ex. No.</td>
<td>Chemical Compounds</td>
<td>NMR</td>
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<td>235 1-235</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, DMSO-d6: 3.83 (dd, J=13, 6.5 Hz, 1H), 4.10 (dd, J = 13, 4.0 Hz, 1H), 5.07 (dd, J = 6.5, 4.0 Hz, 1H), 6.50 (dd, J = 8.0, 1.5 Hz, 1H), 6.89 (t, J = 8.0 Hz, 1H), 7.28 (dd, J = 7.7, 1.5 Hz, 1H), 7.31 (dd, J = 7.7, 4.0 Hz, 1H), 7.37 (d, J = 8.3 Hz, 2H), 7.71 (d, J = 6.5 Hz, 2H), 7.90 (d, J = 8.3 Hz, 2H), 8.03 (dd, J = 8.0, 1.5 Hz, 1H), 8.44 (dd, J = 4.0, 1.5 Hz, 1H), 10.77 (s, 1H).</td>
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<tr>
<td>236 1-236</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, DMSO-d6: 2.27 (s, 3H), 3.34 (s, 3H), 4.10 (br s, 4H), 6.78 (s, J = 7.0 Hz, 1H), 6.96 (d, J = 6.5 Hz, 1H), 7.18-7.29 (m, 6H), 7.70 (d, J = 7.9 Hz, 1H), 8.16 (s, 1H).</td>
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<td>237 2-01</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, DMSO-d6: 3.75 (t, J = 4.4 Hz, 2H), 4.40 (t, J = 4.4 Hz, 2H), 4.53 (d, J = 5.6 Hz, 2H), 5.39 (t, J = 5.6 Hz, 1H), 5.97 (s, 2H), 6.42 (dd, J = 8.1, 1.6 Hz, 1H), 6.75 (dd, J = 8.1, 7.6 Hz, 1H), 6.85 (d, J = 5.4 Hz, 1H), 7.02 (dt, J = 7.5, 1.6 Hz, 1H), 7.13 (dd, J = 8.4, 2.1 Hz, 1H), 7.42 (d, J = 2.1 Hz, 1H), 7.92 (d, J = 2.0 Hz, 1H), 8.34 (d, J = 2.0 Hz, 1H), 10.01 (s, 1H).</td>
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<td>238 2-02</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, DMSO-d6: 3.77 (t, J = 4.4 Hz, 2H), 4.21-4.24 (m, 4H), 4.42 (t, J = 4.4 Hz, 2H), 4.59 (d, J = 5.6 Hz, 2H), 5.43 (t, J = 5.6 Hz, 1H), 6.45 (dd, J = 8.1, 1.6 Hz, 1H), 7.37 (d, J = 8.3 Hz, 1H), 8.10 (d, J = 8.8 Hz, 1H), 7.04 (dd, J = 7.9, 1.6 Hz, 1H), 7.15 (dd, J = 8.6, 2.6 Hz, 1H), 7.38 (d, J = 2.6 Hz, 1H), 7.95 (d, J = 2.1 Hz, 1H), 8.37 (d, J = 2.1 Hz, 1H), 9.96 (s, 1H).</td>
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<td>239 2-03</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, DMSO-d6: 3.78 (t, J = 4.2 Hz, 2H), 4.42 (t, J = 4.4 Hz, 2H), 4.56 (d, J = 4.6 Hz, 2H), 5.44 (t, J = 5.1 Hz, 1H), 6.47 (dd, J = 8.3, 1.4 Hz, 1H), 6.68 (t, J = 7.9 Hz, 1H), 7.04 (dd, J = 7.4, 1.4 Hz, 1H), 7.35 (d, J = 8.8 Hz, 2H), 7.86 (d, J = 8.8 Hz, 2H), 7.95 (d, J = 2.3 Hz, 1H), 8.37 (d, J = 1.0 Hz, 1H), 10.35 (s, 1H).</td>
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<td>240 2-04</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, DMSO-d6: 3.78 (t, J = 4.2 Hz, 2H), 4.43 (t, J = 4.2 Hz, 2H), 4.56 (d, J = 5.6 Hz, 2H), 5.44 (t, J = 5.6 Hz, 1H), 6.49 (dd, J = 7.9, 1.4 Hz, 1H), 6.81 (t, J = 7.7 Hz, 1H), 7.05 (dd, J = 7.7, 1.2 Hz, 1H), 7.56 (dd, J = 9.0, 1.2 Hz, 1H), 7.78 (dd, J = 8.8, 2.3 Hz, 1H), 7.95 (d, J = 1.9 Hz, 1H), 8.14 (d, J = 2.3 Hz, 1H), 8.37 (d, J = 1.4 Hz, 1H), 10.49 (s, 1H).</td>
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<tr>
<td>241 2-05</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, DMSO-d6: 3.79 (t, J = 4.4 Hz, 2H), 4.43 (t, J = 4.2 Hz, 2H), 4.56 (d, J = 4.2 Hz, 2H), 5.44 (br s, 1H), 6.49 (dd, J = 7.9, 1.4 Hz, 1H), 6.81 (t, J = 7.9 Hz, 1H), 7.05 (dd, J = 7.7, 1.6 Hz, 1H), 7.71 (d, J = 8.6 Hz, 2H), 7.97 (t, J = 4.8 Hz, 3H), 8.37 (d, J = 2.3 Hz, 1H), 10.52 (s, 1H).</td>
</tr>
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<td>242 2-06</td>
<td><img src="image" alt="Chemical Structure" /></td>
<td>400 MHz, DMSO-d6: 3.79 (t, J = 4.2 Hz, 2H), 4.43 (t, J = 4.2 Hz, 2H), 4.56 (d, J = 4.2 Hz, 2H), 5.44 (br s, 1H), 6.49 (dd, J = 7.9, 1.4 Hz, 1H), 6.81 (t, J = 7.9 Hz, 1H), 7.05 (dd, J = 7.7, 1.6 Hz, 1H), 7.71 (d, J = 8.6 Hz, 2H), 7.97 (t, J = 4.8 Hz, 3H), 8.37 (d, J = 1.9 Hz, 1H), 10.69 (s, 1H).</td>
</tr>
</tbody>
</table>
DEMANDES OU BREVETS VOLUMINEUX

LA PRÉSENTE PARTIE DE CETTE DEMANDE OU CE BREVETS COMPREND PLUS D'UN TOME.

CECI EST LE TOME _1_ DE _2_

NOTE: Pour les tomes additionnels, veillez contacter le Bureau Canadien des Brevets.

________________________________________

JUMBO APPLICATIONS / PATENTS

THIS SECTION OF THE APPLICATION / PATENT CONTAINS MORE THAN ONE VOLUME.

THIS IS VOLUME _1_ OF _2_

NOTE: For additional volumes please contact the Canadian Patent Office.
CLAIMS

1. A condensed benzamide compound represented by the following general formula [1] or a pharmaceutically acceptable salt thereof:

\[
\begin{align*}
\text{[wherein } Z \text{ is} \\
(1) \cdot \text{O} \cdot, \\
(2) \cdot \text{NR}^5 \cdot \text{ (wherein } R^5 \text{ is a hydrogen atom or a C1-6 alkyl group)}, \\
(3) \cdot \text{S} \cdot, \\
(4) \cdot \text{SO} \cdot \text{ or} \\
(5) \cdot \text{SO}_2 \cdot; \\
\text{1 is 0, 1 or 2;} \\
\text{m is 0, 1 or 2;} \\
\text{R}^1 \text{ is} \\
(1) \text{ a hydrogen atom or} \\
(2) \text{ a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the following Group A:} \\
\text{Group A:} \\
(a) \text{ a halogen atom,} \\
(b) \text{ a hydroxyl group,} \\
(c) \text{ a C1-6 alkoxy group,}
\end{align*}
\]
(d) a carboxyl group,
(e) a C1-6 alkoxy carbonyl group,
(f) \(-\text{CONR}^6\text{R}^7\) (wherein \(\text{R}^6\) and \(\text{R}^7\) are the same or different and each represents a hydrogen atom, a C1-6 alkyl group or an acyl group and said alkyl group may be substituted with a hydroxyl group or an acyloxy group),
(g) \(-\text{NR}^6\text{R}^7\) (wherein \(\text{R}^6\) and \(\text{R}^7\) are the same as above),
(h) \(-\text{NR}^6\text{COR}^7\) (wherein \(\text{R}^6\) and \(\text{R}^7\) are the same as above),
(i) \(-\text{NR}^6\text{CONR}^6\text{R}^7\) (wherein \(\text{R}^6\) and \(\text{R}^7\) are the same as above, and \(\text{R}^6\) is a hydrogen atom or a C1-6 alkyl group); and
(j) \(-\text{NR}^6\text{SO}_3\text{R}^9\) (wherein \(\text{R}^6\) is the same as above, and \(\text{R}^9\) is a C1-6 alkyl group);

\(\text{R}^2\) is
(1) a hydrogen atom,
(2) a hydroxyl group,
(3) a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A (wherein the Group A is the same as above),
(4) a carboxyl group,
(5) a C1-6 alkoxy carbonyl group or
(6) \(-\text{CONR}^{10}\text{R}^{11}\) (wherein \(\text{R}^{10}\) and \(\text{R}^{11}\) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group);

or \(\text{R}^2\) together with \(\text{R}^1\) forms \(-\text{O}\);

\(\text{R}^3\) is
(1) a hydrogen atom, or
(2) a C1-6 alkyl group;

\(\text{R}^4\) is
(1) a hydrogen atom,
(2) a halogen atom
(3) a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the following Group B,
(4) a C1-6 alkoxy group which may be substituted with 1 to 5 substituents selected from the following Group B,
(5) a cycloalkyl group which may be substituted with 1 to 5 substituents selected from the following Group B,
(6) an aralkyl group which may be substituted with 1 to 5 substituents selected from the following Group B,
(7) an alkoxy group which may be substituted with 1 to 5 substituents selected from the following Group B, or
(8) a cycloalkylalkoxy group which may be substituted with 1 to 5 substituents selected from the following Group B,

Group B:
(a) a halogen atom,
(b) a halo C1-6 alkyl group,
(c) a hydroxyl group,
(d) a halo C1-6 alkoxy group,
(e) a C1-6 alkoxy carbonyl group,
(f) a C1-6 alkoxy group,
(g) a carboxyl group,
(h) -CONR^{12}R^{13} (wherein R^{12} and R^{13} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group);
(i) -NR^{12}R^{13} (wherein R^{12} and R^{13} are the same as above),
(j) -NR^{12}COR^{13} (wherein R^{12} and R^{13} are the same as above),
(k) \(-\text{NR}^{14}\text{CONR}^{12}\text{R}^{13}\) (wherein \(\text{R}^{12}\) and \(\text{R}^{13}\) are the same as above, and \(\text{R}^{14}\) is a hydrogen atom or a C1-6 alkyl group),

(l) \(-\text{SO}_2\text{R}^{15}\) (wherein \(\text{R}^{15}\) is a C1-6 alkyl group), and

(m) \(-\text{NR}^{12}\text{SO}_2\text{R}^{15}\) (wherein \(\text{R}^{12}\) and \(\text{R}^{15}\) are the same as above);

(n) a hydroxyl group,

(o) \(-\text{NR}^{16}\text{R}^{17}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{17}\) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group)

(p) \(-\text{COR}^{18}\) (wherein \(\text{R}^{18}\) is a C1-6 alkyl group, a C1-6 alkoxy group, a cycloalkyl group, an aralkyl group, an aralkoxy group, a cycloalkylalkoxy group or a hydroxyl group),

(q) \(-\text{CONR}^{16}\text{R}^{17}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{17}\) are the same as above),

(r) \(-\text{NR}^{19}\text{CONR}^{16}\text{R}^{17}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{17}\) are the same as above, and \(\text{R}^{19}\) is a hydrogen atom or a C1-6 alkyl group),

(s) \(-\text{NR}^{16}\text{COOR}^{20}\) (wherein \(\text{R}^{16}\) is the same as above, and \(\text{R}^{20}\) is a C1-6 alkyl group or a cycloalkyl group),

(t) \(-\text{SR}^{20}\) (wherein \(\text{R}^{20}\) is the same as above),

(u) \(-\text{SOR}^{20}\) (wherein \(\text{R}^{20}\) is the same as above),

(v) \(-\text{SO}_2\text{R}^{20}\) (wherein \(\text{R}^{20}\) is the same as above),

(w) \(-\text{SO}_2\text{NR}^{16}\text{R}^{17}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{17}\) are the same as above) or

(x) \(-\text{NR}^{16}\text{COR}^{18}\) (wherein \(\text{R}^{16}\) and \(\text{R}^{18}\) are the same as above);

\(V\) is

(1) a single bond or

(2) \(-\text{(CR}^{21}\text{R}^{22})_n\) - (wherein \(\text{R}^{21}\) and \(\text{R}^{22}\) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, and \(n\) is 1 or 2);

P1 and P2 rings are the same or different and each represents
(1) a carbocyclic group which may be substituted with 1 to 5 substituents selected from the following group C or
(2) a heterocyclic group which may be substituted with 1 to 5 substituents selected from the following group
Group C:
(a) a halogen atom,
(b) a hydroxyl group,
(c) a C1-6 alkoxy group which may be substituted with 1 to 5 substituents selected from the Group A,
(d) an C1-6 alkylthio group,
(e) a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A,
(f) -CONR^{23}R^{24} (wherein R^{23} and R^{24} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
(g) -NR^{123}R^{124} (wherein R^{123} and R^{124} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
(h) -NR^{223}COR^{224} (wherein R^{223} and R^{224} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),
(i) -NR^{25}CONR^{323}R^{324} (wherein R^{323} and R^{324} are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which
may be substituted with 1 to 5 substituents selected from the Group A),

(j) \( \cdot SR^{26} \) (wherein \( R^{26} \) is a C1-6 alkyl group),

(k) \( \cdot SOR^{126} \) (wherein \( R^{126} \) is a C1-6 alkyl group),

(l) \( \cdot SO_2R^{226} \) (wherein \( R^{226} \) is a C1-6 alkyl group),

(m) \( \cdot NR^{223}SO_2R^{326} \) (wherein \( R^{423} \) is a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A, and \( R^{326} \) is a C1-6 alkyl group),

(n) \( \cdot SO_2NR^{523}R^{524} \) (wherein \( R^{523} \) and \( R^{524} \) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A),

(o) \( \cdot COR^{27} \) (wherein \( R^{27} \) is a C1-6 alkyl group, C1-6 alkoxy group, a cycloalkyl group, an aralkyl group, an aralkoxy group, a cycloalkylalkoxy group or a hydroxyl group),

(p) a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (o) of the Group C,

(q) a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (o) of the Group C,

(r) \( \cdot 0-R^{28} \) (wherein \( R^{28} \) is an acyl group, a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (o) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (o) of the Group C),

(s) \( \cdot 0-(CR^{121}R^{122})_n-R^{128} \) (wherein \( R^{121} \) and \( R^{122} \) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, \( n \) is 1 or 2, and \( R^{128} \) is an acyl group, a carbocyclic
group which may be substituted with 1 to 5 substituents selected from (a) to (i) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (l) of the Group C,
(t) a nitro group, and
(u) a cyano group;

wherein in the above formula I, "cycloalkyl" is C3-8 cycloalkyl; "arylalkyl" is a group comprising a C6-14 aryl and a C1-6 alkyl; "aralkoxy" is a group comprising a C14 aryl and a C1-6 alkoxy; "cycloalkylalkoxy group" is a group comprising a C3-8 cycloalkyl and a C1-6 alkoxy; "carbocyclic group" is a saturated or unsaturated cyclic hydrocarbon group having 3 to 14 carbon atoms and which comprises a C6-14 aryl group, a cycloalkyl group, a cycloalkenyl group having 3-8 carbon atoms optionally comprising one or two double bonds, or a condensed carbocyclic ring; "heterocyclic group" is a saturated or partially or fully unsaturated 5-membered or 6-membered monocyclic heterocyclic ring containing 1 to 4 hetero atoms comprising a nitrogen atom, an oxygen atom or a sulfur atom besides the carbon atoms, condensed rings of said heterocyclic ring, or a condensed ring of said heterocyclic ring and a carbocyclic ring selected from benzene, cyclopentane and cyclohexane.

2. The condensed benzamide compound or pharmaceutically acceptable salt thereof according to claim 1 wherein Z is -O-, -NR²-, -S- or -SO-. 
3. The condensed benzamide compound or pharmaceutically acceptable salt thereof according to claim 1 or 2 wherein R³ is a hydrogen atom or a C1-4 alkyl group.

4. The condensed benzamide compound or pharmaceutically acceptable salt thereof according to any one of claims 1 to 3 wherein the P1 ring is a saturated or unsaturated 5-membered or 6-membered heterocyclic ring having 1 to 4 hetero atoms selected from a nitrogen atom, an oxygen atom and a sulfur atom, or a condensed ring of these heterocyclic rings, or a condensed heterocyclic ring of said heterocyclic ring and a carbocyclic ring selected from benzene, cyclopentane and cyclohexane, or a phenyl group (wherein said heterocyclic ring may be substituted with a halogen atom, a hydroxyl group, a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A, and a C1-6 alkoxy group
which may be substituted with 1 to 5 substituents selected from the Group A).

5. The condensed benzamide compound or pharmaceutically acceptable salt thereof according to claim 4 wherein the P1 ring is a saturated or unsaturated 5-membered or 6-membered heterocyclic ring having at least 1 to 3 nitrogen atoms or a phenyl group (wherein said heterocyclic ring may be substituted with a C1-6 alkyl group which may be substituted with a hydroxyl group or a C1-6 alkyl group, a halogen atom, a hydroxyl group, and a C1-6 alkoxy group which may be substituted with 1 to 5 substituents selected from the Group A).

6. The condensed benzamide compound or a pharmaceutically acceptable salt thereof according to claim 5 wherein the P1 ring is a heterocyclic group selected from a pyridyl group, a pyrazinyl group and a thiazolyl group or a phenyl group (wherein these heterocyclic groups may be substituted with a C1-6 alkyl group which may be substituted with a hydroxyl group, a halogen atom, a hydroxyl group, and a C1-6 alkoxy group which may be substituted with 1 to 5 substituents selected from the Group A).

7. The condensed benzamide compound or a pharmaceutically acceptable salt thereof according to claims 1 to 3 wherein
the P2 ring is a carbocyclic group which may be substituted with a substituent group selected from

- a halogen atom,
- a hydroxyl group,
- a C1-6 alkoxy group (wherein said alkoxy group may be substituted with a halogen atom, \( \cdot \text{CONR}^{623} \text{R}^{624} \) (wherein \( \text{R}^{623} \) and \( \text{R}^{624} \) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group A), a C3-8 cycloalkyl group, a C1-6 alkoxy group, a carboxyl group or a C1-6 alkoxy carbonyl group),
- a C1-6 alkyl group (wherein said alkyl group may be substituted with a halogen atom, a hydroxyl group or a C1-6 alkoxy group),

\( \cdot \text{NR}^{123} \text{R}^{124} \) (wherein \( \text{R}^{123} \) and \( \text{R}^{124} \) are the same as above),

\( \cdot \text{NR}^{223} \text{COR}^{224} \) (wherein \( \text{R}^{223} \) and \( \text{R}^{224} \) are the same as above),

\( \cdot \text{COR}^{27} \) (wherein \( \text{R}^{27} \) is a C1-6 alkyl group, C1-6 alkoxy group, a cycloalkyl group, an aralkyl group, an aralkoxy group, a cycloalkylalkoxy group or a hydroxyl group),

\( \cdot \text{CONR}^{23} \text{R}^{24} \) (wherein \( \text{R}^{23} \) and \( \text{R}^{24} \) are the same as above),

- a heterocyclic group as a saturated or unsaturated substituent group which has 1 to 3 nitrogen atoms as heteroatoms (wherein said heterocyclic group may be substituted with a substituent group selected from a hydroxyl group, \( \cdot \text{CONR}^{723} \text{R}^{724} \) (wherein \( \text{R}^{723} \) and \( \text{R}^{724} \) are the same or different and each represents a hydrogen atom or a C1-6 alkyl group which may be substituted with 1 to 5 substituents selected from the Group
A), a C1-6 alkoxy group, a carboxyl group, a C1-6 alkyl group which may be substituted with a C1-6 alkoxy group, a C1-6 alkoxy carbonyl group and an acyloxy group),

- O-R²⁸ (wherein R²⁸ is an acyl group, a carbocyclic group as defined in claim 1, but which is optionally substituted with 1 to 5 substituents selected only from (a) to (i) of the Group C or a heterocyclic group as defined in claim 1, but which is optionally substituted with 1 to 5 substituents selected only from (a) to (l) of the Group C),

- O-(CR¹²¹R¹²²)ₙ-R¹²⁸ (wherein R¹²¹ and R¹²² are the same or different and each represents a hydrogen atom or a C1-6 alkyl group, n is 1 or 2, and R¹²⁸ is an acyl group, a carbocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (i) of the Group C or a heterocyclic group which may be substituted with 1 to 5 substituents selected from (a) to (l) of the Group C),

- a nitro group and
- a cyano group, or
- a heterocyclic group as defined in claim 1.
8. The condensed benzamide compound or a pharmaceutically acceptable salt thereof according to claim 7 wherein the P2 ring in claim 7 is a phenyl group, a cyclohexyl group, a thiazolyl group, a pyridyl group, a piperidyl group, a piperidino group, a quinolyl group, a benzo[1,3]dioxo group, a 2,3-dihydrobenzo[1,4]dioxo group or a 1,2,3,4-tetrahydroquinolyl group.

9. The condensed benzamide compound according to claim 1 which is:
   1) N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
   2) 8-(4-tert-butylphenyl)carbamoyl-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylic acid,
   3) N-(4-tert-butylphenyl)-4-(3-chloropyridin-2-yl)-3-methyl 1-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
   4) N-(4-tert-butylphenyl)-1-(3-chloropyridin-2-yl)-4-methyl 1-1,2,3,4-tetrahydroquinazoline-5-carboxamide,
   5) N-(4-tert-butylphenyl)-9-(3-chloropyridin-2-yl)-6,7,8,9-tetrahydro-5-oxa-9-aza-benzocycloheptene-4-carboxamide,
   6) N-(4-chlorophenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
   7) 4-(3-chloropyridin-2-yl)-N-(4-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
8) N·(1-tert-butylpiperidin-4-yl)-4·(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
9) 4·(3-chloropyridin-2-yl)·N·(4-methylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
10) 4·(3-chloropyridin-2-yl)·N·(trans-4-methylcyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
11) 4·(3-chloropyridin-2-yl)·N·(4-isobutoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
12) N·benzyl·4·(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
13) N·(4-chlorophenyl)methyl·4·(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
14) N·(4-tert-butylyphenyl)methyl·4·(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
15) 4·(3-chloropyridin-2-yl)·N·(4-trifluoromethylphenyl)methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
16) N·(4-tert-butylyphenyl)·4·(pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
17) N·(4-tert-butylyphenyl)·4·(3-trifluoromethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
18) N·(4-tert-butylyphenyl)·4·(3-chloropyridin-2-yl)-2,2-dimethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
19) N·(4-tert-butylyphenyl)·4·(3-chloropyridin-2-yl)-2-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
20) N·(4-tert-butylyphenyl)-6-chloro-4·(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
21) 4-(3-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
22) 4-(3-chloropyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
23) 4-(3-chloropyridin-2-yl)-N-(4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
24) N-(trans-4-tert-butylcyclohexyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
25) 4-(3-chloropyridin-2-yl)-N-(4-chloro-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
26) 4-(3-chloropyridin-2-yl)-N-(4-fluorophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
27) 4-(3-chloropyridin-2-yl)-N-(3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
28) 4-(3-chloropyridin-2-yl)-N-(4-trifluoromethylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
29) 4-(3-chloropyridin-2-yl)-N-(5-trifluoromethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
30) 4-(3-chloropyridin-2-yl)-N-(2-trifluoromethylpyridin-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
31) 4-(3-chloropyridin-2-yl)-N-(4-isopropylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
32) 4-(3-chloropyridin-2-yl)-N-(quinolin-7-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
33) 4-(3-chloropyridin-2-yl)-N-(1-methyl-1,2,3,4-tetrahydrouquinolin-7-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
34) 4-(3-chloropyridin-2-yl)-N-(4-methoxycarbonylphenyl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
35) 4-(4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carbonyl]aminobenzoic acid,
36) N-(4-carbamoylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
37) 4-(3-chloropyridin-2-yl)-N-(4-methylcarbamoylphenyl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
38) 4-(3-chloropyridin-2-yl)-N-(4-dimethylcarbamoylphenyl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
39) N-(4-acetylphenyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
40) 4-(3-chloropyridin-2-yl)-N-[4-(1-hydroxy-1-methyl)ethylphenyl]-3,4-dihydro-2H-benzoxazine-8-carboxamide,
41) 4-(3-chloropyridin-2-yl)-N-[4-(1-hydroxy-1-methyl)propylphenyl]-3,4-dihydro-2H-benzoxazine-8-carboxamide,
42) 4-(3-chloropyridin-2-yl)-N-(1-isobutyrylpiperidin-4-yl)phenyl-3,4-dihydro-2H-benzoxazine-8-carboxamide,
43) N-(trans-4-tert-butoxycyclohexyl)-4-(3-chloropyridin-2-yl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
44) 4-(3-chloropyridin-2-yl)-N-(3-isopropoxyphenyl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
45) 4-(3-chloropyridin-2-yl)-N-(3-isobutyl)oxyphenyl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
46) N-(4-tert-butylphenyl)-4-(pyridin-4-yl)-3,4-dihydro-2H-benzoxazine-8-carboxamide,
47) N-(4-tert-butylphenyl)-4-(3-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
48) 4-(3-chloropyridin-2-yl)-N-(4-piperidinophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
49) 4-(3-chloropyridin-2-yl)-N-(4-dimethylaminophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
50) 4-(3-chloropyridin-2-yl)-N-(4-morpholinophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
51) 4-(3-chloropyridin-2-yl)-N-(3-fluoro-4-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
52) 4-(3-chloropyridin-2-yl)-N-(2-fluoro-4-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
53) 4-(3-chloropyridin-2-yl)-N-(4-fluoro-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
54) 4-(3-chloropyridin-2-yl)-N-(4-dimethylamino-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
55) 4-(3-chloropyridin-2-yl)-N-(4-isopropoxy-3-trifluorothylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
56) 4-(3-chloropyridin-2-yl)-N-(4-methoxy-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
57) 4-(3-chloropyridin-2-yl)-N-(4-isobutoxy-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
58) N-(4-tert-butylphenyl)-4-(4-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
59) N-(4-tert-butylphenyl)-4-(6-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
60) N-(4-tert-butylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
61) N-(4-tert-butylphenyl)-4-(pyridin-3-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
62) N-(4-cyanophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
63) N-(3-amino-4-chlorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
64) N-(3-acetamido-4-chlorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
65) N-[4-(4-carbamoylpiperidin-1-yl)-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
66) N-[3-fluoro-4-(4-methylcarbamoylpiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
67) N-[4-(4-dimethylcarbamoylpiperidin-1-yl)-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
68) N-[4-(4-ethoxypiperidin-1-yl)-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
69) N-[3-fluoro-4-(4-isopropoxypiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
70) N-[3-fluoro-4-(3-methoxypiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
71) N-[3-fluoro-4-(4-methoxymethylpiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
72) N-[3-fluoro-4-(3-methoxypyrrolidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
73) N-(4-isobutyloxy-3-methoxycarbonylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
74) 2-isobutyloxy·{[4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carbonyl]amino}benzoic acid,
75) N-(3-carbamoyl-4-isobutyloxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
76) N-(4-isobutyloxy-3-methylcarbamoylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
77) N-(3-dimethylcarbamoyl-4-isobutyloxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
78) N-(4-acetylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
79) N-[4-(1-hydroxy-1-methyl)ethylphenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
80) N-(3-acetyl-4-chlorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
81) N-[4-chloro-3-(1-hydroxy-1-methyl)ethylphenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
82) N-(4-(1-methoxy-1-methyl)ethylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
83) N-(3-chloro-4-methoxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
84) 4-(5-methylpyridin-2-yl)-N-(4-propoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
85) N-(3-fluoro-4-propoxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
86) N-(4-ethoxy-3-fluorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
87) N-(3-ethoxy-4-methyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
88) N-(3-carbamoylmethoxy-4-methyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
89) N-(3-methoxyethoxy-4-methyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
90) N-[3-fluoro-4-(2-methoxymethylpiperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
91) N-(4-chlorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
92) N-(3-fluoro-4-methylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
93) N-(2-chloropyridin-5-yl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
94) 4-(3-chloropyridin-2-yl)-N-(4-ethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
95) 4-(3-chloropyridin-2-yl)N-(3-fluoro-4-piperidinophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
96) 4-(3-chloropyridin-2-yl)-N-(trans-4-ethoxycyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
97) 4-(3-chloropyridin-2-yl)-N-(trans-4-isopropoxycyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
98) 4-(3-chloropyridin-2-yl)-N-(trans-4-cyclopentyloxy cyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
99) 4-(3-chloropyridin-2-yl)-N-(trans-4-cyclohexyloxy cyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
100) 4-(3-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
101) 4-(3-chloropyridin-2-yl)-N-[3-fluoro-4-(4-methoxypiperidin-1-yl)phenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
102) 4-(3-chloropyridin-2-yl)-N-[4-(4-ethoxypiperidin-1-yl)-3-fluorophenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
103) 4-(3-chloropyridin-2-yl)N-[3-fluoro-4-(4-isopropoxypiperidin-1-yl)phenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
104) N-(3-chloropyridin-2-yl)-N-(3-fluoro-4-isobutyloxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
105) N-(trans-4-ethoxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
106) N-(trans-4-isopropoxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
107) N-(trans-4-cyclohexyl-oxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
108) N-(trans-4-amino-cyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
109) N-(1,4-dioxaspiro[4,5]deca-8-yl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
110) 4-(5-methylpyridin-2-yl)-N-(4-oxocyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
111) 4-(5-methylpyridin-2-yl)-N-(cis-4-morpholino-cyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
112) N-(trans-4-dimethylaminocyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
113) N-(trans-4-diethylaminocyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
114) 4-(5-methylpyridin-2-yl)-N-(cis-4-(pyrrolidin-1-yl)cyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
115) 4-(5-methylpyridin-2-yl)-N-(trans-4-(pyrrolidin-1-yl)cyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
116) 4-(5-methylpyridin-2-yl)-N-(4-piperidino-cyclohexyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
117) 4-[(5-methylpyridin-2-yl)-N-(cis-4-morpholinocyclohexyl-1)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
118) N-(trans-4-acetamidocyclohexyl)-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
119) N-(trans-4-cyclohexyloxyocyclohexyl)-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
120) 4-[(5-chloropyridin-2-yl)-N-(4-ethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
121) N-(4-chlorophenyl)-4-[(5-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
122) 4-[(5-chloropyridin-2-yl)-N-[3-fluoro-4-(4-methoxypiperidin-1-yl)phenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
123) 4-[(5-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
124) 4-[(5-methylpyridin-2-yl)-N-(2-phenylethyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
125) N-[(2-(4-chlorophenyl)ethyl)]-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
126) N-[(2-(3-chlorophenyl)ethyl)]-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
127) N-[(2-(2-chlorophenyl)ethyl)]-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,  
128) 6-[(8-(4-trifluoromethylphenyl)carbamoyl)-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
129)4·[(3-chloro·5·(1-hydroxy·1·methyl)ethylpyridin·2·yl)·N·(4·trifluoromethylphenyl)·3·4·dihydro·2H·benzo[1·4]oxazine·8·carboxamide,
130)4·[(3-chloro·5·(1-hydroxyethyl)pyridin·2·yl)·N·(4·trifluoromethylphenyl)·3·4·dihydro·2H·benzo[1·4]oxazine·8·carboxamide,
131)5·chloro·6·[8·(3·fluoro·4·trifluoromethylphenyl)carbamoyl·2·3·dihydro·benzo[1·4]oxazin·4·yl]nicotinic acid,
132)N·(4·tert·butylphenyl)·4·(5·methoxycarbonylpyridin·2·yl)·3·4·dihydro·2H·benzo[1·4]oxazine·8·carboxamide,
133)6·[8·(4·tert·butylphenyl)carbamoyl·2·3·dihydro·benzo[1·4]oxazin·4·yl]nicotinic acid,
134)5·chloro·6·[8·(4·trifluoromethylphenyl)carbamoyl·2·3·dihydro·benzo[1·4]oxazin·4·yl]nicotinic acid,
135)5·chloro·6·[8·(4·tert·butylphenyl)carbamoyl·2·3·dihydro·benzo[1·4]oxazin·4·yl]nicotinic acid,
136)4·(5·acetyl·3·chloropyridin·2·yl)·N·(4·trifluoromethylphenyl)·3·4·dihydro·2H·benzo[1·4]oxazine·8·carboxamide,
137)N·(4·tert·butylphenyl)·4·(5·methylcarbamoylpyridin·2·yl)·3·4·dihydro·2H·benzo[1·4]oxazine·8·carboxamide,
138)N·(4·tert·butylphenyl)·4·(5·carbamoylpyridin·2·yl)·3·4·dihydro·2H·benzo[1·4]oxazine·8·carboxamide,
139)N·(4·tert·butylphenyl)·4·(5·diethylaminopyridin·2·yl)·3·4·dihydro·2H·benzo[1·4]oxazine·8·carboxamide,
140)N·(4·tert·butylphenyl)·4·(5·nitropyridin·2·yl)·3·4·dihydro·2H·benzo[1·4]oxazine·8·carboxamide,
141) 4-(5-aminopyridin-2-yl)-N-(4-tert-butylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
142) 4-(5-acetamidopyridin-2-yl)-N-(4-tert-butylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
143) 4-(5-methoxymethylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
144) 4-(5-ethoxypyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
145) 4-(3-chloro-5-methoxypyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
146) 4-(5-hydroxypyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
147) 4-(5-methoxypyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
148) N-(4-tert-butylphenyl)-4-(4-chloropyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
149) 4-(5-fluoropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
150) 4-(3,5-difluoropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
151) N-(4-tert-butylphenyl)-4-(3-methoxypyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
152) N-(4-tert-butylpiperidin-1-yl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
153) 4-[5-(2-hydroxyethoxy)pyridin-2-yl]-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
154) 4-[(3-chloro-5-(2-hydroxyethyl)pyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
155) 4-(5-methylpyridin-2-yl)-N-(4-neopentyloxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
156) N-(3-fluoro-4-piperidinophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
157) N-(3-fluoro-4-morpholinophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
158) N-(3,4-difluorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
159) N-(3-fluoro-4-methoxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
160) N-(4-ethoxyphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
161) N-[4-(2-oxo-piperidin-1-yl)phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
162) 4-(5-methylpyridin-2-yl)-N-(1-methyl-1,2,3,4-tetrahydroquinolin-7-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
163) N-(4-dimethylamino-3-fluorophenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
164) N-[3-fluoro-4-[N-(2-methoxyethyl)-isopropylamino]phenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
165) N·[4·[N·(2-acetoxyethyl)·isopropylamino]·3·fluorophenyl]·4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carboxamide,
166) N·[3·fluoro·4·[N·(2-hydroxyethyl)·isopropylamino]phenyl]·4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carboxamide,
167) N·[4·(4-ethoxycarbonylpiperidin-1-yl)·3·fluorophenyl]·4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carboxamide,
168) 1·(4·{[4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carbonylamino]phenyl)piperidine·4·carboxylic acid,
169) N·[3·fluoro·4·(4-methoxypiperidin-1-yl)phenyl]·4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carboxamide,
170) N·[4·(4-acetoxy Piperidin-1-yl)·3·fluorophenyl]·4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carboxamide,
171) N·[3·fluoro·4·(4-hydroxypiperidin-1-yl)phenyl]·4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carboxamide,
172) N·[3·methoxy·4·methylphenyl]·4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carboxamide,
173) N·[3·(2-hydroxyethoxy)·4·methylphenyl]·4·(5-methylpyridin-2-yl)·3,4·dihydro-2H·benzo[1,4]oxazine·8·carboxamide,
174) N-[4-(1-tert-butoxycarbonylpiperidin-4-yl)oxy-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
175) N-[4-(piperidin-4-yl)oxy-3-fluorophenyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
176) N-(trans-4-ethoxycarbonylmethoxyoxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
177) (trans-4-[(4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carbonyl]amino)cyclohexyloxy)acetic acid,
178) N-(trans-4-carbamoylmethoxyoxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
179) N-(trans-4-methylcarbamoylmethoxyoxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
180) N-(trans-4-dimethylcarbamoylmethoxyoxycyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
181) N-[trans-4-(2-hydroxyethoxy)oxycyclohexyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
182) N-(trans-4-methoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
183) N-(trans-4-isopropoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
184) N-(cis-4-methoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
185) N-(cis-4-isopropoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
186) N-(trans-4-tert-butoxymethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
187) N-(trans-4-isobutoxy cyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
188) N-(4,4-dimethylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
189) N-[trans-4-(3-methylbutyloxy)cyclohexyl]-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
190) N-(trans-4-benzyloxy cyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
191) N-(trans-4-isopropylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
192) N-(trans-4-propylcyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
193) N-(trans-4-neopentyloxy cyclohexyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
194) N-(4-tert-butylphenyl)-4-(5-methoxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
5-chloro-6-[8-(3-chloro-4-trifluoromethoxyphenyl)carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
5-chloro-6-[8-(4-trifluoromethoxyphenyl)carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
4-(5-methylaminopyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4-(5-ethoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4-(3-chloro-5-methoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
5-(2-hydroxyethoxy)pyridin-2-yl]-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4-(5-hydroxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4-(5-methoxypyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4-(3-cyanopyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4-(3-carbamoylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4-(3-methylcarbamoylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
4-(3-dimethylcarbamoylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
207) 4·(3-benzylloxycarbonylpyridin-2-yl)-N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
208) 2·[8·(4-trifluoromethoxyphenyl)carbamoyl-2,3-dihydro-benzo[1,4]oxazin-4-yl]nicotinic acid,
209) 4·(3-benzylloxypyridin-2-yl)-N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
210) 4·(3-hydroxypyridin-2-yl)-N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
211) 4·[3·(2-hydroxyethoxy)pyridin-2-yl]-N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
212) 2·[8·(4-trifluoromethoxyphenyl)carbamoyl]·2,3-dihydro-benzo[1,4]oxazin-4-yl]pyridin-3-yl]oxyacetic acid,
213) 4·(3-carbamoylmethoxypyridin-2-yl)-N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
214) 4·(3-methylcarbamoylmethoxypyridin-2-yl)-N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
215) 4·(3-dimethylcarbamoylmethoxypyridin-2-yl)-N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
216) 4·[3·(pyridin-2-yl)methoxypyridin-2-yl]-N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
217) 4·{(3-(pyridin-3-yl)methyloxy)pyridin-2-yl}·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
218) 4·{(5-cyanopyridin-2-yl)·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
219) 4·{(5-acetimidomethyl)pyridin-2-yl}·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
220) 4·{(3-chloro-5-methoxymethyl)pyridin-2-yl}·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
221) 4·{(5-(N-methylacetamido)methyl)pyridin-2-yl}·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
222) 4·{(5-aminomethyl)pyridin-2-yl}·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
223) 4·{(5-dimethylaminomethyl)pyridin-2-yl}·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
224) 4·{(3-chloropyridin-2-yl)·2-methylcarbamoyl}·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
225) 4·{(6-hydroxymethyl)pyridin-2-yl}·N·{(4-trifluoromethylphenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
226) 4·{(3-chloropyridin-2-yl)·2-dimethylcarbamoyl}·N·{(4-trifluoromethoxy)phenyl}·3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
227) 4-(5-carbamoyl-3-chloropyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
228) (S)-3-acetamidomethyl-4-(3-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
229) (R)-4-(3-chloropyridin-2-yl)-8-(4-trifluoromethylphenylcarbamoyl)-3,4-dihydro-2H-benzo[1,4]oxazine-3-carboxylic acid,
230) 4-(3-chloropyridin-2-yl)-2-methoxycarbonyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
231) 4-(3-chloropyridin-2-yl)-2-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
232) 4-(3-chloropyridin-2-yl)-8-(4-trifluoromethoxyphenylcarbamoyl)-3,4-dihydro-2H-benzo[1,4]oxazine-2-carboxylic acid,
233) (S)-3-carbamoyl-4-(3-chloropyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
234) (S)-4-(3-chloropyridin-2-yl)-3-methylcarbamoyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
235) 2-carbamoyl-4-(3-chloropyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
236) N-(4-chlorophenyl)-N-methyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
237) N-(benzo[1,3]dioxol-5-yl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
238) N-(2,3-dihydrobenzo[1,4]dioxin-6-yl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
239) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
240) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3-chloro-4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
241) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
242) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3-fluoro-4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
243) 4-(5-hydroxymethylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
244) 4-(5-hydroxymethylpyridin-2-yl)-N-(4-isobutyloxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
245) 4-(3-chloro-5-methoxymethylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
246) 4-(4-hydroxymethylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
247) 4-(3-hydroxymethylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
248) (+)-4-{3-chloro-5-(1-hydroxyethyl)pyridin-2-yl}-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
249) (-)-4-{3-chloro-5-(1-hydroxyethyl)pyridin-2-yl}-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
250) (+)-4-{3-chloro-5-(1-hydroxyethyl)pyridin-2-yl}-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
251) (-)-4-{3-chloro-5-(1-hydroxyethyl)pyridin-2-yl}-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
252) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(2-trifluoromethylpyridin-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
253) N-(4-bromo-3-chlorophenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
254) 4-[5-(1-hydroxy-1-methyl)ethylpyridin-2-yl]-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
255) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-isopropoxy-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
256) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(2,3-dichloropyridin-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
257) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3,4,5-trichlorophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
258) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(6-fluorobenzothiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
259) N-(4-bromo-3-trifluoromethylphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
260) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(2-trifluoromethylbenzimidazol-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
261) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3,4,5-trifluorophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
262) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-fluoro-3-nitrophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
263) N-(3-amino-4-fluorophenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
264) N-(tert-butylphenyl)-4-(5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
265) N-(3,5-bistrifluoromethylphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
266) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-[2,2,2-trifluoroethoxy)phenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
267) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-[3-chloro-(2,2,2-trifluoroethoxy)phenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
268) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(4-trifluoromethylmercaptophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
269) N-(trans-4-tert-butylcyclohexyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
270) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3-fluoro-4-isopropoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
271) 4-(3-chloro-5-hydroxymethylpyridin-2-yl)-N-(3-fluoro-4-piperidinophenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
272) N-(4-tert-butylphenyl)-4-(3-chloro-5-hydroxymethylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
273) 4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-N-[4-(1-hydroxy-2,2,2-trifluoro-1-trifluoromethyl)ethylphenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
274) 4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-N-[2,2,3,3-tetrafluoro-2,3-dihydrobenzo[1,4]dioxin-6-yl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
275) 4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-N-[4-isobutyloxophenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
276) 4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-N-[2,3-dichlorophenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
277) N-[4-bromo-3-fluorophenyl]-4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
278) N-[4-bromophenyl]-4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
279) 4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-N-[3,5-dichlorophenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
280) 4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-N-[4-difluoromethoxyphenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
281) 4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-N-[4-chlorophenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
282) 4-[(3-chloro-5-hydroxymethyl pyridin-2-yl)-N-[4-isopropylphenyl]-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
283) 4·(3-chloro-5-hydroxymethylpyridin-2-yl)-N·(3-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

284) 4·[3-chloro-5-(1-hydroxyethyl)pyridin-2-yl]-N·(3-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

285) 4·(3-chloropyridin-2-yl)-N·(4-trifluoromethoxyphenyl)-3,4-dihydro-3-oxo-2H-benzo[1,4]oxazine-8-carboxamide,

286) (R)·4·(3-chloropyridin-2-yl)-3-hydroxymethyl-N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

287) (R)·4·(3-chloropyridin-2-yl)-3-hydroxymethyl-N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

288) (S)·4·(3-chloropyridin-2-yl)-3-hydroxymethyl-N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

289) (S)·4·(3-chloropyridin-2-yl)-3-hydroxymethyl-N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

290) (S)·4·(5-chloropyridin-2-yl)-3-hydroxymethyl-N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,

291) (S)·4·(5-chloropyridin-2-yl)-3-hydroxymethyl-N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
292) (S) - 3-hydroxymethyl - 4 - (5-methylpyridin - 2-yl) - N - (4 - trifluoromethoxyphenyl) - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
293) (S) - 3-hydroxymethyl - 4 - (5-methylpyridin - 2-yl) - N - (4 - trifluoromethylphenyl) - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
294) (S) - 4 - (3-chloropyridin - 2-yl) - N - (3-fluoro - 4-isopropoxyphenyl) - 3-hydroxymethyl - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
295) (S) - 4 - (3-chloropyridin - 2-yl) - N - (3-fluoro - 4-trifluoromethoxyphenyl) - 3-hydroxymethyl - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
296) (S) - 4 - (3-chloropyridin - 2-yl) - N - (3-chloro - 4-piperidinophenyl) - 3-hydroxymethyl - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
297) (S) - 4 - (3-chloropyridin - 2-yl) - N - (4-dimethylamino - 3-fluorophenyl) - 3-hydroxymethyl - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
298) (S) - 4 - (3-chloropyridin - 2-yl) - N - (3-fluoro - 4-methylphenyl) - 3-hydroxymethyl - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
299) (S) - N - (trans - 4-tert-butylcyclohexyl) - 4 - (3-chloropyridin - 2-yl) - 3-hydroxymethyl - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
300) (S) - 4 - (3-chloropyridin - 2-yl) - 3-hydroxymethyl - N - (trans-4-neopentyloxy cyclohexyl) - 3,4-dihydro - 2H-benzo[1,4]oxazine - 8-carboxamide,
301) (S) - N - (4 - chlorophenyl) - 4 - (3 - chloropyridin - 2 - yl) - 3 - hydroxymethyl - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
302) (S) - 4 - (3 - chloropyridin - 2 - yl) - 3 - hydroxymethyl - N - (4 - isopropylphenyl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
303) (S) - 4 - (3 - chloropyridin - 2 - yl) - N - (4 - fluoro - 3 - trifluoromethylphenyl) - 3 - hydroxymethyl - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
304) (S) - N - (4 - bromo - 3 - chlorophenyl) - 4 - (3 - chloropyridin - 2 - yl) - 3 - hydroxymethyl - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
305) (S) - 4 - (3 - chloropyridin - 2 - yl) - N - (4 - dimethylamino - 3 - trifluoromethylphenyl) - 3 - hydroxymethyl - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
306) (S) - 4 - (3 - chloropyridin - 2 - yl) - N - (4 - isopropoxy - 3 - trifluoromethylphenyl) - 3 - hydroxymethyl - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
307) (S) - 4 - (3 - chloropyridin - 2 - yl) - 3 - hydroxymethyl - N - (4 - piperidino - 3 - trifluoromethylphenyl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
308) (S) - 4 - (3 - chloropyridin - 2 - yl) - N - (4 - ethoxy - 3 - fluorophenyl) - 3 - hydroxymethyl - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
309) (S) - 4 - (3 - chloropyridin - 2 - yl) - 3 - hydroxymethyl - N - (4 - methoxy - 3 - trifluoromethylphenyl) - 3, 4 - dihydro - 2H - benzo[1, 4]oxazine - 8 - carboxamide,
310) (S)-4-[(3-chloro-5-methoxy-pyridin-2-yl)-3-hydroxymethyl N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
311) (S)-4-[(3-chloro-5-methoxy-pyridin-2-yl)-3-hydroxymethyl N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
312) (S)-3-hydroxymethyl-4-[(5-methylpyridin-2-yl)-N-(2-trifluoromethylpyridin-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
313) (S)-3-hydroxymethyl N-(4-methoxy-3-trifluoromethylphenyl)-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
314) (S)-3-hydroxymethyl-4-[(5-methylpyridin-2-yl)-N-(4-piperidino-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
315) (S)-N-(3-fluoro-4-trifluoromethylphenyl)-3-hydroxymethyl-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
316) (S)-N-(3-chloro-4-trifluoromethoxyphenyl)-3-hydroxymethyl-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
317) (S)-N-(3,4-dichlorophenyl)-3-hydroxymethyl-4-[(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
318) (S)-4-[(3-chloropyridin-2-yl)-3-hydroxymethyl N-(4-isobutoxy-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
319) (S)-4-(3-chloropyridin-2-yl)-N-(3,4-dichlorophenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
320) (S)-4-(3-chloropyridin-2-yl)-N-(4-chloro-3-trifluoromethylphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
321) (S)-N-(4-bromo-3-fluorophenyl)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
322) (S)-N-(4-chloro-3-trifluoromethylphenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
323) (S)-3-hydroxymethyl-N-(4-isopropoxy-3-trifluoromethylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
324) (S)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-morpholino-3-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
325) (S)-N-(2-chloro-4-trifluoromethylphenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
326) (S)-3-hydroxymethyl-4-(3-methylpyridin-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
327) (S)-3-hydroxymethyl-4-(3-methylpyridin-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
328) (S)-N-(4-tert-butoxy-3-fluorophenyl)-4-(3-chloropyridin-2-yl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
329) (S)-4-(3-chloropyridin-2-yl)-N-(3-fluoro-4-isobutoxyphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
330) (S)-3-hydroxymethyl-N-(4-isobutoxy-3-trifluoromethylphenyl)-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
331) (S)-N-(3-fluoro-4-isopropoxyphenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
332) (S)-N-(3-fluoro-4-morpholinophenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
333) (S)-N-(4-chlorophenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
334) (S)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-N-(2-morpholino-4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
335) (S)-N-(4-tert-butoxy-3-fluorophenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
336) (S)-N-(4-bromo-3-fluorophenyl)-3-hydroxymethyl-4-(5-methylpyridin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
337)4-(3-chloropyridin-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
338) (S)·4-(3-chloropyridin-2-yl)·N·(3-fluoro-4-trifluoromethylphenyl)·3-hydroxymethyl·3,4-dihydro-2H-benzo[1,4]oxazine·8-carboxamide,
339) (S)·N·(3-fluoro-4-isobutyloxyphenyl)·3-hydroxymethyl·4·(5-methylpyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine·8-carboxamide,
340) (S)·N·(4-fluoro-3-trifluoromethylphenyl)·3-hydroxymethyl·4·(5-methylpyridin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine·8-carboxamide,
341)4·(3-chloropyridin-2-yl)·3·(1-hydroxyethyl)·N·(4-trifluoromethoxyphenyl)·3,4-dihydro-2H-benzo[1,4]oxazine·8-carboxamide,
342) (R)·9·(3-chloropyridin-2-yl)·7-hydroxy·N·(4-trifluoromethoxyphenyl)·6,7,8,9-tetrahydro-5-oxa-9-azabenzocycloptane·4-carboxamide,
343) (R)·9·(3-chloropyridin-2-yl)·7-hydroxy·N·(4-trifluoromethylphenyl)·6,7,8,9-tetrahydro-5-oxa-9-azabenzocycloptane·4-carboxamide,
344) (S)·9·(3-chloropyridin-2-yl)·7-hydroxy·N·(4-trifluoromethoxyphenyl)·6,7,8,9-tetrahydro-5-oxa-9-azabenzocycloptane·4-carboxamide,
345) N·(4-tert-butylphenyl)·4·(pyrazin-2-yl)·3,4-dihydro-2H-benzo[1,4]oxazine·8-carboxamide,
346) N·(4-chlorophenyl)-4·(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
347) N·(4-ethoxyphenyl)-4·(5-ethylpyrimidin-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
348) 4·(6-chloropyridazin-3-yl)·N·(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
349) 4·(4-methylthiazol-2-yl)·N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
350) 4·(5-methylthiazol-2-yl)·N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
351) (S)·3-hydroxymethyl-4·(5-methylthiazol-2-yl)·N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
352) (S)·3-hydroxymethyl-4·(5-methylthiazol-2-yl)·N·(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
353) (S)·N·(3,4-dichlorophenyl)-3-hydroxymethyl-4·(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
354) (S)·N·(3-fluoro-4-isopropoxyphenyl)-3-hydroxymethyl-4·(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
355) (S)·N·(3-fluoro-4-tert-butoxyphenyl)-3-hydroxymethyl-4·(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
356) (S)-N-(3-fluoro-4-isobutoxyphenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
357) (S)-N-(4-bromo-3-fluorophenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
358) (S)-N-(4-fluoro-3-trifluoromethylphenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
359) (S)-N-(3-fluoro-4-morpholinophenyl)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
360) (S)-3-hydroxymethyl-4-(5-methylthiazol-2-yl)-N-(2-trifluoromethylpyridin-5-yl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
361) (S)-3-hydroxymethyl-4-(5-methyloxazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
362) (S)-3-hydroxymethyl-4-(5-methyloxazol-2-yl)-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
363) (S)-4-(4,5-dimethylthiazol-2-yl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
364) (S)-4-(4,5-dimethylthiazol-2-yl)-3-hydroxymethyl-N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
365) (S)-3-hydroxymethyl-4-(5-methyl[1,3,4]thiadiazol-2-yl) -N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
366) (S)-3-hydroxymethyl-4-(5-methyl[1,3,4]thiadiazol-2-yl) -N-(4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
367) 4-(4,5-dimethylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]thiazine-8-carboxamide,
368) 4-(4,5-dimethylthiazol-2-yl)-N-(4-trifluoromethoxyphenyl)-1-oxo-3,4-tetrahydro-benzo[1,4]thiazine-8-carboxamide,
369) N-(4-tert-butylphenyl)-4-(4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
370) N-(4-isobutoxyphenyl)-4-(4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
371) N-(4-chlorophenyl)-4-(4-methoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
372) (S)-4-(2-chlorophenyl)-3-hydroxymethyl-N-(4-trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
373) (S)-4-(2-chlorophenyl)-3-hydroxymethyl-N-(3-fluoro-4-trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
374) (S)-4-(2-chlorophenyl)-N-(3-fluoro-4-trifluoromethylphenyl)-3-hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide,
375) (S)-4-(4-chlorophenyl)-3-hydroxymethyl-N-(4-
trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-
carboxamide,
376) (S)-3-hydroxymethyl-4-(4-methoxyphenyl)-N-(4-
trifluoromethoxyphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-
carboxamide,
377) (S)-3-hydromethyl-4-(4-methoxyphenyl)-N-(4-
trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-
carboxamide,
378) (S)-4-(4-chlorophenyl)-3-hydroxymethyl-N-(4-
trifluoromethylphenyl)-3,4-dihydro-2H-benzo[1,4]oxazine-8-
carboxamide, or
379) (S)-4-(4-chlorophenyl)-N-(4-chlorophenyl)-3-
hydroxymethyl-3,4-dihydro-2H-benzo[1,4]oxazine-8-carboxamide;
or a pharmaceutically acceptable salt thereof.

10. A pharmaceutical composition comprising a condensed
benzamide compound or a pharmaceutically acceptable salt
thereof according to any one of claims 1 to 9 and a
pharmaceutically acceptable carrier.

11. A pharmaceutical composition comprising a condensed
benzamide compound or a pharmaceutically acceptable salt
thereof according to any one of claims 1 to 9 and a
pharmaceutically acceptable carrier for treating and/or
preventing a disease or condition selected from pain,
neurodegenerative disease, cerebral apoplexy, ischemic
symptom, nerve injury, neurogenic skin disorder, inflammatory
disease, pruritus, allergic rhinitis, apoplexy, irritable
bowel syndrome, asthma, chronic obstructive pulmonary
disease, dermatitis, mucositis, stomach and duodenal ulcer,
inflammatory bowel disease, bladder hypersensitivity,
overactive bladder type frequent urination and urinary incontinence.

12. A pharmaceutical composition for treating and/or preventing pain comprising a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9 and a pharmaceutically acceptable carrier.

13. The pharmaceutical composition according to claim 12 wherein the pain is acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy or neurodegenerative disease.

14. An inhibitor of vanilloid receptor subtype 1 (VR1) activity comprising a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9 and a pharmaceutically acceptable carrier.

15. A use of a pharmacologically effective amount of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9 for treating and/or preventing pain, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer, inflammatory bowel disease, bladder
hypersensitivity, overactive bladder type frequent urination or urinary incontinence.

16. A use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9 for treating and/or preventing pain.

17. The use according to claim 16 wherein the pain is acute pain, chronic pain, neuropathic paid, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute posttherpetic neuralgia, posttherpetic neuralgia, chronic posttherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy or neurodegenerative disease.

18. A commercial package comprising a pharmaceutical composition according to any one of claims 10 to 13 and written instructions concerning said pharmaceutical composition stating that said composition can be used or should be used for treating and/or preventing a disease selected from pain, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer, inflammatory bowel disease, bladder hypersensitivity, overactive bladder type frequent urination and urinary incontinence.

19. Use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9 for preparing a pharmaceutical composition according to claim 11 for treating and/or preventing a
disease selected from pain, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritus, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer, inflammatory bowel disease, bladder hypersensitivity, overactive bladder type frequent urination and urinary incontinence.

20. Use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9 for preparing a pharmaceutical composition for treating and/or preventing pain according to claim 12.

21. The use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to claim 20 wherein the pain is acute pain, chronic pain, neuropathic pain, rheumatoid arthritis pain, neuralgia, neuropathy, hyperalgesia, migraine, joint pain, acute postherpetic neuralgia, postherpetic neuralgia, chronic postherpetic neuralgia, postoperative pain, cancer pain, inflammatory pain, interstitial cystitis, posttraumatic neuralgia, diabetic neuropathy or neurodegenerative disease.

22. A drug comprising a combination of a pharmaceutical composition comprising a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9 and a pharmaceutically acceptable carrier with one or more agents selected from the group which consists of an anti-virus agent, an antidepressant, an anticonvulsant, an antiarrhythmic, an anesthetic, a local anesthetic, an N-methyl-D-aspartate receptor antagonist, an adrenal cortical steroid, a nerve block, a nonsteroidal anti-
inflammatory analgesic, a narcotic, an antagonist analgesic, an $\alpha_2$-adrenaline receptor agonist, capsaicin, a calcium channel antagonist, and a potassium channel opener.

23. Use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9 for preparing a drug defined in claim 22.

24. A use of one or more agents selected from the group which consists of an anti-virus agent, an antidepressant, an anticonvulsant, an antiarrhythmic drug, an anesthetic drug, a local anesthetic, an N-methyl-D-aspartate receptor antagonist, adrenal cortical steroid, a nerve block, a nonsteroidal anti-inflammatory analgesic, narcotics, an antagonist analgesic, $\alpha_2$-adrenaline receptor agonist, capsaicin, a calcium channel antagonist, and a potassium channel opener; in combination with a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9, for treating or preventing pain, neurodegenerative disease, cerebral apoplexy, ischemic symptom, nerve injury, neurogenic skin disorder, inflammatory disease, pruritis, allergic rhinitis, apoplexy, irritable bowel syndrome, asthma, chronic obstructive pulmonary disease, dermatitis, mucositis, stomach and duodenal ulcer, inflammatory bowel disease, bladder hypersensitivity, overactive bladder type frequent urination or urinary incontinence.

25. A use of a condensed benzamide compound or a pharmaceutically acceptable salt thereof according to any one of claims 1 to 9, in combination with stimulation-produced analgesia selected from acupuncture, transcutaneous electroacupuncture stimulation therapy, transcutaneous
electrical nerve stimulation therapy, silver spike point (SSP) therapy, peripheral nerve stimulation therapy, spinal cord electrical stimulation therapy, electroconvulsive therapy, laser therapy and low-frequency therapy for treating and/or preventing pain.

26. A use according to claim 15 or claim 16 wherein said pain comprises postoperative neuralgia after a surgical operation selected from cicatrectomy, nerve freezing solidification, peripheral nerve excision, spinal cord dorsal root excision, sympathectomy, spinal cord dorsal root entry zone destruction, cordotomy, and frontal lobe excision.
[Image of a chemical structure with labeled atoms and bonds]