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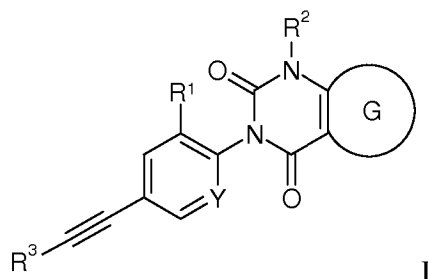
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(56) References cited:
WO-A1-2011/128279 WO-A1-2013/050454

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Description

[0001] The present invention relates to compounds of formula I



wherein

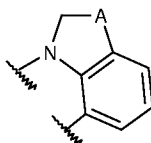
Y is N or C-R¹;

G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, lower alkyl, lower alkoxy, lower alkoxy substituted by halogen or NRR'; R and R' are independently from each other hydrogen or lower alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N-lower alkyl group;

R¹ is hydrogen, halogen or lower alkyl substituted by halogen;

R^{1'} is hydrogen, halogen or lower alkyl substituted by halogen;

R² is hydrogen, lower alkyl, lower alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group



for A being -CH₂-, -CH₂CH₂-, or -C(CH₃)₂-,

R³ is phenyl or pyridinyl, wherein the N atom in the pyridinyl group may be in different positions;

or to a pharmaceutically acceptable salt or acid addition salt, to a racemic mixture, or to its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof.

[0002] It has been surprisingly been found that the compounds of general formula I are positive allosteric modulators (PAMs) of metabotropic glutamate receptor 4 (mGluR4).

Metabotropic glutamate receptor 4 is a protein that in humans is encoded by the *GRM4* gene.

[0003] Together with GRM6, GRM7 and GRM8 it belongs to group III of the Metabotropic glutamate receptor family, and is negatively coupled to adenylate cyclase via activation of the Gai/o protein. It is expressed primarily on presynaptic terminals, functioning as an autoreceptor or heteroreceptor and its activation leads to decreases in transmitter release from presynaptic terminals. mGluR4 is currently receiving much attention based primarily upon its unique distribution and the recent evidence that activation of this receptor plays key modulatory role in many CNS and non-CNS pathways (Celanire S, Campo B, Expert Opinion in Drug Discovery, 2012)

[0004] The similarity in the ligand binding domains of group III mGluRs creates a challenge for identifying selective orthosteric agonists of this receptor, although some progress has been made in this area. However, targeting positive allosteric modulators (PAMs) rather than orthosteric agonists provides a broader opportunity to identify molecules that are exclusively selective between mGluRs.

[0005] mGluR4 PAM is emerging as a promising target for the treatment of motor (and non motor) symptoms as well as a disease-modifying agent in Parkinson's disease through a non-dopaminergic approach.

[0006] Parkinson's disease is a progressive neurodegenerative disease that results in the loss of dopaminergic neurons in the substantia nigra (SN). One consequence of the depletion of dopamine in this disease is a series of movement disorders, including bradykinesia, akinesia, tremor, gait disorders and problems with balance. These motor disturbances form the hallmark of PD, although there are many other non-motor symptoms that are associated with the disease. Early

in the course of the disease, PD symptoms are effectively treated by dopamine replacement or augmentation, with the use of dopamine D2 receptor agonists, levodopa or monoamine oxidase B inhibitors. However, as the disease progresses these agents become less effective in controlling motor symptoms. Additionally, their use is limited by the emergence of adverse effects including dopamine agonist-induced dyskinesias. Consequently, there remains a need for new approaches to the treatment of PD that improve the effectiveness of the control of motor symptoms.

[0007] Activation of metabotropic glutamate receptor 4 (mGluR4) has been proposed as a potential therapeutic approach to Parkinson's disease. A member of the group III mGluRs, mGluR4 is predominantly a presynaptic glutamate receptor that is expressed in several key locations in the basal ganglia circuits that control movement. Activation of mGluR4 with group III-preferring agonists decreases inhibitory and excitatory post synaptic potentials, presumably by decreasing the release of GABA and glutamate respectively.

[0008] The search for novel drugs that relieve motor symptoms of Parkinsonism whilst attenuating the ongoing degeneration of nigrostriatal neurons is of particular interest. Orthosteric mGluR4 agonist L-AP4 has demonstrated neuroprotective effects in a 6-OHDA rodent model of PD and first positive allosteric modulator (-)-PHCCC reduced nigrostriatal degeneration in mice treated with 1-Methyl-4-Phenyl-1,2,3,6-Tetrahydropyridine (MPTP). Those studies provide pre-clinical evidence suggesting that mGluR4 activators constitute a strong approach not only for symptomatic treatments of PD, but also potentially as disease modifiers.

[0009] The neuroprotective effect of selective mGluR4 was also described in Neuroreport, 19(4), 475-8, 2008, Proc.Natl. Acad. Sci, USA, 100(23), 13668-73, 2003 and J.Neurosci. 26(27), 7222-9, 2006 and Mol. Pharmacol. 74(5), 1345-58, 2008..

[0010] Anxiety disorders are among the most prevalent psychiatric disorders in the world, and are co-morbid with Parkinson's disease (Prediger R, et al. Neuropharmacology 2012;62:115-24). Excessive glutamatergic neurotransmission is one important feature of anxiety pathophysiology. Based on presynaptic localization of mGluR4 in brain areas involved in anxiety and mood disorders, and dampening excessive brain excitability, the mGluR4 activators may represent a new generation of anxiolytic therapeutics (Eur. J. Pharmacol., 498(1-3), 153-6, 2004).

[0011] Addex has reported in 2010 that ADX88178 was active in two preclinical rodent models of anxiety: the marble burying test in mice and EPM in mice and rats. ADX88178 also displayed an anxiolytic-like profile in the rat EPM test after oral dosing.

[0012] mGluR4 modulators were also shown to exert anti-depressive actions (Neuropharmacology, 46(2), 151-9, 2004).

[0013] In addition, mGluR4 were also shown to be involved in glucagon secretion inhibition (Diabetes, 53(4), 998-1006, 2004). Therefore, orthosteric or positive allosteric modulators of mGluR4 have potential for the treatment of type 2 diabetes through its hypoglycemic effect.

[0014] Moreover, mGluR4 was shown to be expressed in prostate cancer cell-line (Anticancer Res. 29(1), 371-7, 2009) or colorectal carcinoma (Cli. Cancer Research, 11(9)3288-95, 2005). mGluR4 modulators may therefore have also potential role for the treatment of cancers.

[0015] Other proposed effects of MgluR4 PAM's can be expected for the treatment of emesis, obsessive compulsive disorder and autism.

[0016] Compounds of formula I are distinguished by having valuable therapeutic properties. They can be used in the treatment or prevention of disorders, relating to allosteric modulators for the mGluR4 receptor.

The most preferred indications for compounds which are allosteric modulators for the mGluR4 receptor are Parkinson's disease, anxiety, emesis, obsessive compulsive disorder, autism, neuroprotection, cancer, depression and diabetes type 2.

[0017] The present invention relates to compounds of formula I and to their pharmaceutically acceptable salts, to these compounds as pharmaceutically active substances, to the processes for their production as well as to the compounds for use in the treatment or prevention of disorders, relating to allosteric modulators for the mGluR4 receptor, such as Parkinson's disease, anxiety, emesis, obsessive compulsive disorder, autism, neuroprotection, cancer, depression and diabetes type 2 and to pharmaceutical compositions containing the compounds of formula I..

[0018] Furthermore, the invention includes all racemic mixtures, all their corresponding enantiomers and/or optical isomers.

[0019] The following definitions of the general terms used in the present description apply irrespective of whether the terms in question appear alone or in combination.

[0020] As used herein, the term "lower alkyl" denotes a saturated straight- or branched-chain group containing from 1 to 7 carbon atoms, for example, methyl, ethyl, propyl, isopropyl, n-butyl, i-butyl, 2-butyl, t-butyl and the like. Preferred alkyl groups are groups with 1 - 4 carbon atoms.

[0021] As used herein, the term "lower alkoxy" denotes a lower alkyl group as defined above, which is linked with an O atom.

[0022] The term "cycloalkyl" denotes a saturated ring containing from 3 to 7 carbon atoms, for example cyclopropyl, cyclobutyl, cyclopentyl, cyclopentenyl, cyclohexyl, cyclohexenyl, cycloheptyl or cycloheptenyl.

[0023] The term "heterocycloalkyl" denotes a cycloalkyl ring as defined above, wherein at least one carbon atom is replaced by O, N, or S, for example tetrahydrofuranyl, morpholinyl, piperidinyl or oxetanyl.

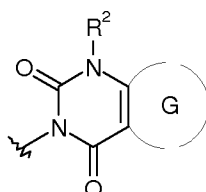
[0024] The term "halogen" denotes chlorine, iodine, fluorine and bromine.

[0025] The term "lower alkyl substituted by halogen" denotes a lower alkyl group as defined above, wherein at least one hydrogen atom is replaced by a halogen atom, for example the following groups: CF_3 , CHF_2 , CH_2F , CH_2CF_3 , CH_2CHF_2 , $\text{CH}_2\text{CH}_2\text{F}$, $\text{CH}_2\text{CH}_2\text{CF}_3$, $\text{CH}_2\text{CH}_2\text{CH}_2\text{CF}_3$, $\text{CH}_2\text{CH}_2\text{Cl}$, $\text{CH}_2\text{CF}_2\text{CF}_3$, $\text{CH}_2\text{CF}_2\text{CHF}_2$, CF_2CHF_2 , $\text{CF}_2\text{CHFCF}_3$, $\text{C}(\text{CH}_3)_2\text{CF}_3$, $\text{CH}(\text{CH}_3)\text{CF}_3$ or $\text{CH}(\text{CH}_2\text{F})\text{CH}_2\text{F}$. The preferred "lower alkyl substituted by halogen" group is CF_3 .

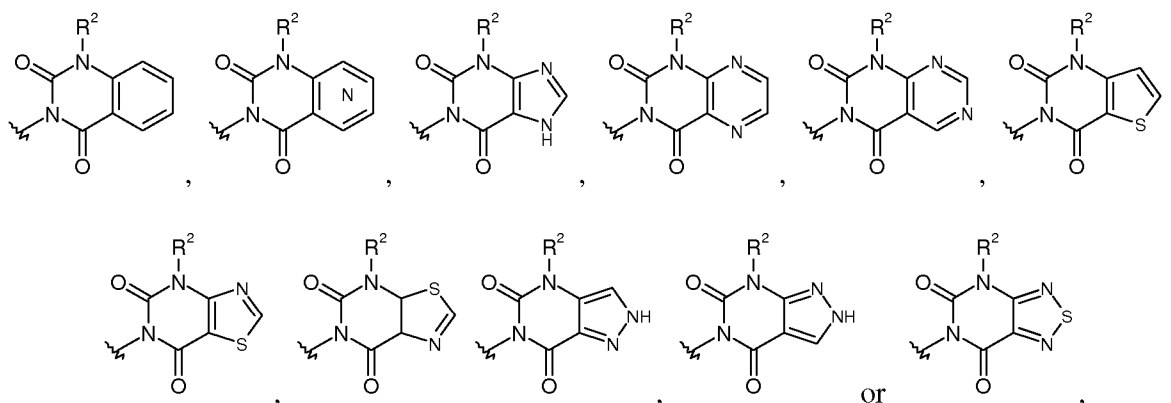
The term "lower alkoxy substituted by halogen" denotes an alkoxy group as defined above, wherein at least one hydrogen atom is replaced by halogen. A preferred group is OCH_2CHF_2 .

[0026] The term "five or six membered saturated heterocyclic group which may contain an additional oxygen, NH or N-lower alkyl group" denotes a piperidine group, a piperazine group or a morpholine group.

The term "lower alkoxyalkyl" denotes an alkyl group as defined above and which is linked with an alkoxy group. The group



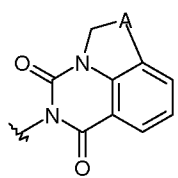
means



wherein these groups may be substituted as described for "G" above, R^2 is defined above and



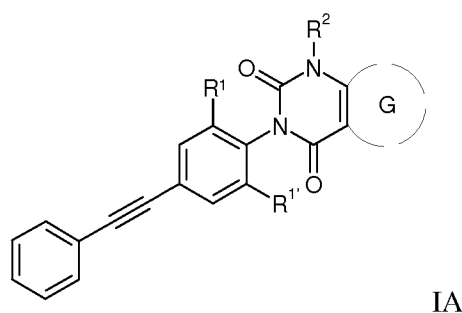
is a pyridine group, wherein the N atom may be in different positions, or means the group



wherein A is as defined above.

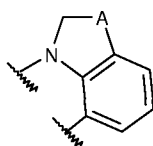
[0027] The term "pharmaceutically acceptable acid addition salts" embraces salts with inorganic and organic acids, such as hydrochloric acid, nitric acid, sulfuric acid, phosphoric acid, citric acid, formic acid, fumaric acid, maleic acid, acetic acid, succinic acid, tartaric acid, methane-sulfonic acid, p-toluenesulfonic acid and the like.

[0028] One embodiment of the invention are compounds of formula IA



wherein

- G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, lower alkyl, lower alkoxy, lower alkoxy substituted by halogen or NRR'; R and R' are independently from each other hydrogen or lower alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N-lower alkyl group;
- R¹ is hydrogen, halogen or lower alkyl substituted by halogen;
- R^{1'} is hydrogen, halogen or lower alkyl substituted by halogen;
- R² is hydrogen, lower alkyl, lower alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group



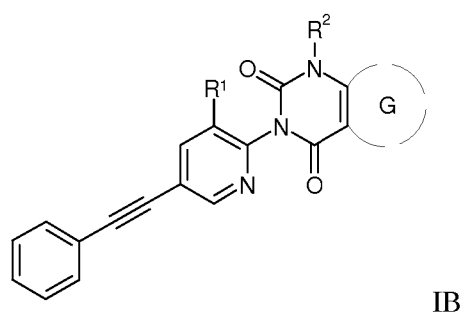
for A being -CH₂-, -CH₂CH₂-, or -C(CH₃)₂-,

or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof, for example the following compounds

- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione
- 8-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione
- 7-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[2,3-d]pyrimidine-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[2,3-d]pyrimidine-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-ethyl-1-methyl-quinazoline-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[4,3-d]pyrimidine-2,4-dione
- 1-[2-Chloro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione
- 2-(2-Chloro-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-quinazoline-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-ethyl-quinazoline-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pteridine-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrimido[4,5-d]pyrimidine-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrimido[4,5-d]pyrimidine-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-6-fluoro-1-methyl-quinazoline-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-7,8-difluoro-1-methyl-quinazoline-2,4-dione
- 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione
 3-(2,6-Difluoro-4-phenylethynyl-phenyl)-1-methyl-1H-pyrido[2,3-d]pyrimidine-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione
 2-(2,6-Difluoro-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4H-thiazolo[4,5-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-methyl-thiazolo[4,5-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-1-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pyrido[2,3-d]pyrimidine-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-pyrido[3,2-d]pyrimidine-2,4-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-methyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,5-dimethyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-(2-methoxyethyl)quinazoline-2,4-dione
 6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4-methyl-2-morpholino-thiazolo[4,5-d]pyrimidine-5,7-dione
 6-[2-Chloro-4-phenylethynyl-phenyl]-4-methyl-4H-thiazolo[4,5-d]pyrimidine-5,7-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-cyclobutyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-isopropyl-1-methyl-quinazoline-2,4-dione
 8-Chloro-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione
 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6-dione
 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6-dione
 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-7-isopropyl-2H-pyrazolo[3,4-d]pyrimidine-4,6-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methyl-quinazoline-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-(oxetan-3-yl)quinazoline-2,4-dione
 6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4-methyl-thiazolo[5,4-d]pyrimidine-5,7-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione
 1-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methyl-quinazoline-2,4-dione
 3-[2-Fluoro-4-(2-phenylethynyl)-6-(trifluoromethyl)phenyl]-1-isopropyl-quinazoline-2,4-dione or
 8-(2,2-Difluoroethoxy)-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione.

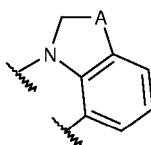
[0029] One embodiment of the invention are further compounds of formula IB,



wherein

G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, lower alkyl, lower alkoxy, lower alkoxy substituted by halogen or NRR'; R and R' are independently from each other hydrogen or lower alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N-lower alkyl group;
 R¹ is hydrogen, halogen or lower alkyl substituted by halogen;

R² is hydrogen, lower alkyl, lower alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group

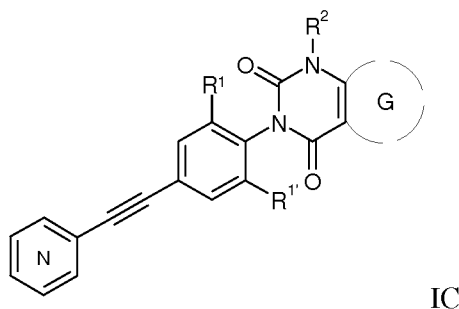


for A being -CH₂-, -CH₂CH₂-, or -C(CH₃)₂-,

or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof, for example the following compounds

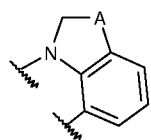
3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1-methyl-quinazoline-2,4-dione
 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1-isopropyl-quinazoline-2,4-dione
 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethyl-quinazoline-2,4-dione
 2-(3-Chloro-5-(phenylethynyl)pyridin-2-yl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 1-Methyl-3-[5-(2-phenylethynyl)-3-(trifluoromethyl)-2-pyridyl]quinazoline-2,4-dione
 3-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethyl-quinazoline-2,4-dione
 1-Isopropyl-3-[5-(2-phenylethynyl)-2-pyridyl]quinazoline-2,4-dione
 6-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione or
 6-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione.

[0030] One embodiment of the invention are compounds of formula IC



wherein

G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, lower alkyl, lower alkoxy, lower alkoxy substituted by halogen or NRR'; R and R' are independently from each other hydrogen or lower alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N-lower alkyl group;
 R¹ is hydrogen, halogen or lower alkyl substituted by halogen;
 R^{1'} is hydrogen, halogen or lower alkyl substituted by halogen;
 R² is hydrogen, lower alkyl, lower alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group

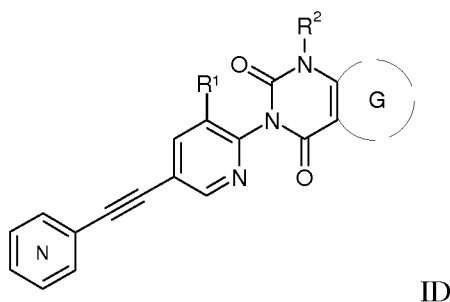


for A being -CH₂-, -CH₂CH₂-, or -C(CH₃)₂-,

or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof, for example the following compounds

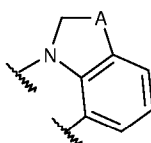
- 3-[2-Chloro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-pteridine-2,4-dione
 3-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione
 2-(2,6-Difluoro-4-(pyridin-3-ylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-quinazoline-2,4-dione
 3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione
 6-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 6-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-(oxetan-3-yl)quinazoline-2,4-dione or
 3-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione.

[0031] One embodiment of the invention are compounds of formula ID,



wherein

- G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, lower alkyl, lower alkoxy, lower alkoxy substituted by halogen or NRR'; R and R' are independently from each other hydrogen or lower alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N-lower alkyl group;
 R¹ is hydrogen, halogen or lower alkyl substituted by halogen;
 R² is hydrogen, lower alkyl, lower alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group



for A being -CH₂-, -CH₂CH₂-, or -C(CH₃)₂-,

or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof, for example the following compound

6-[3-Chloro-5-[2-(3-pyridyl)ethynyl]-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione.

[0032] The preparation of compounds of formula I of the present invention may be carried out in sequential or convergent synthetic routes. Syntheses of the compounds of the invention are shown in the following scheme 1. The skills required for carrying out the reaction and purification of the resulting products are known to those skilled in the art. The substituents and indices used in the following description of the processes have the significance given herein before.

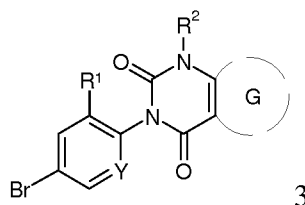
[0033] The compounds of formula I can be manufactured by the methods given below, by the methods given in the examples or by analogous methods. Appropriate reaction conditions for the individual reaction steps are known to a person skilled in the art. The reaction sequence is not limited to the one displayed in the schemes, however, depending on the starting materials and their respective reactivity the sequence of reaction steps can be freely altered. Starting

materials are either commercially available or can be prepared by methods analogous to the methods given below, by methods described in references cited in the description or in the examples, or by methods known in the art.

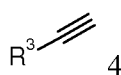
[0034] The present compounds of formula I and their pharmaceutically acceptable salts may be prepared by methods, known in the art, for example by the process variant described below, which process comprises

a) reacting a compound of formula 2

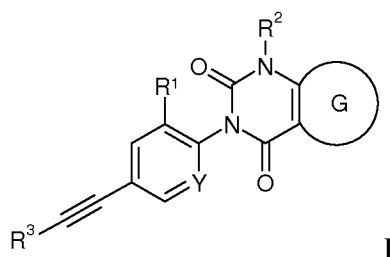
[0035]



with a compound of formula



to a compound of formula I

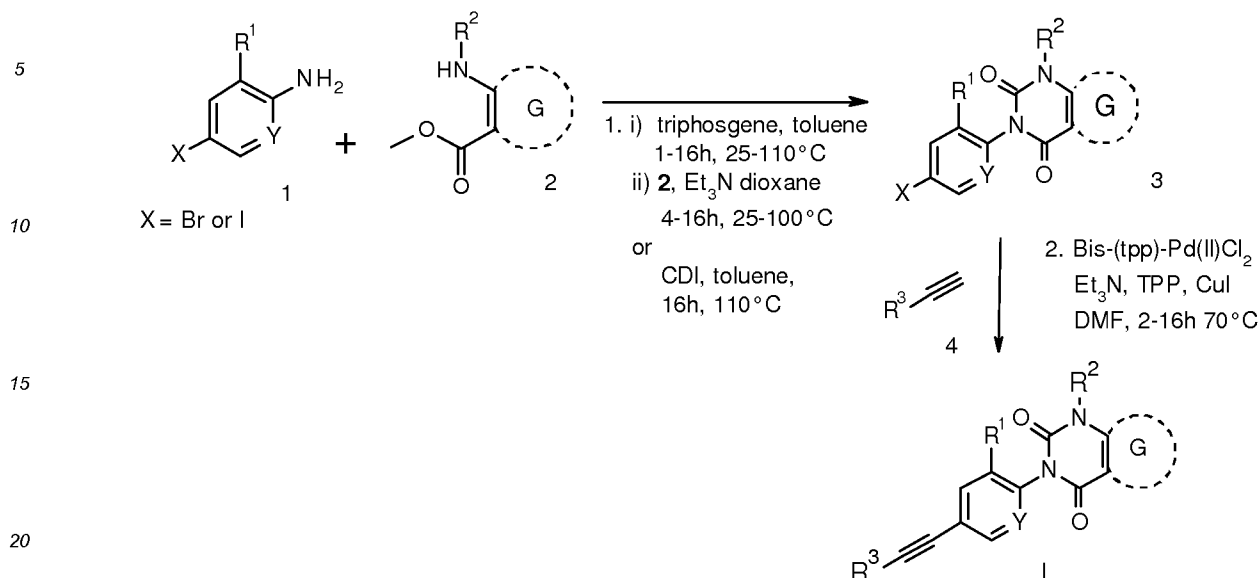


wherein the substituents are described above, or

if desired, converting the compounds obtained into pharmaceutically acceptable acid addition salts.

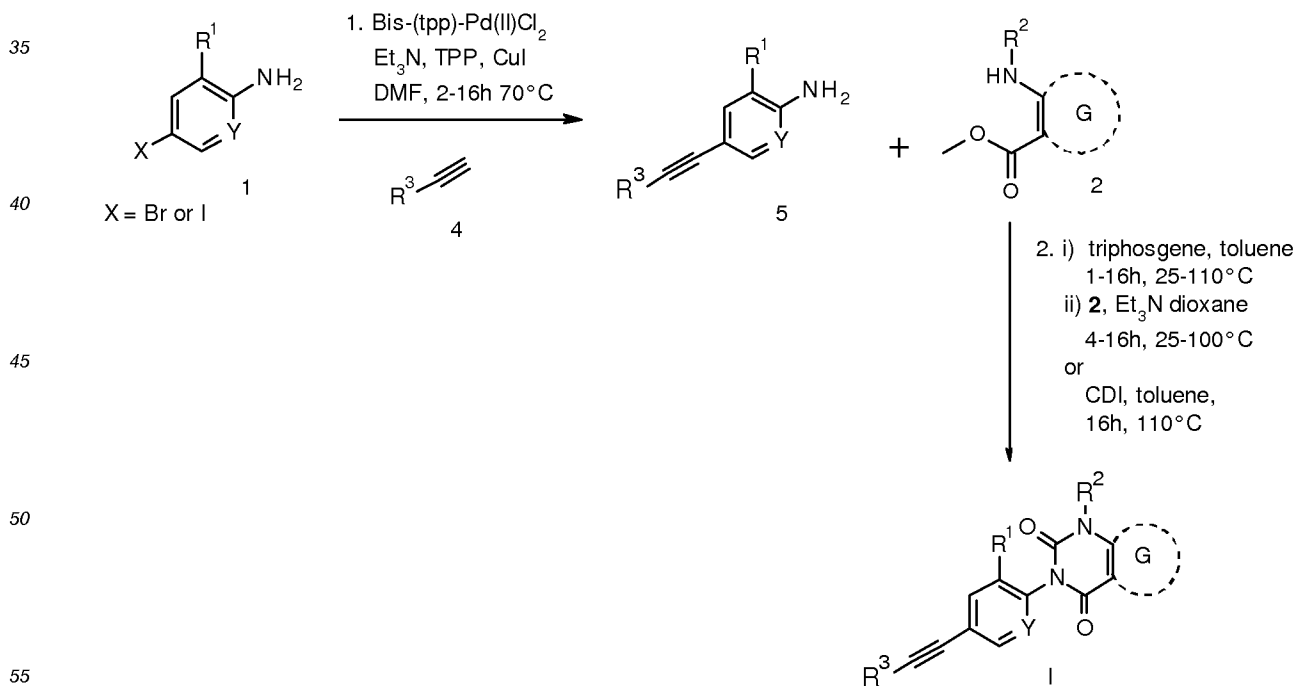
[0036] The preparation of compounds of formula I is further described in more detail in scheme 1 and 2 and in examples 1 -74.

Scheme 1



[0037] An ethynyl-phenyl, ethynyl-pyridyl substituted pyrimidinedione compound of formula I can be obtained for example by reacting an appropriately substituted aniline or aminopyridine **1** with an appropriate aromatic or heteroaromatic orthoaminoester **2** with phosgene or a phosgene equivalent such as triphosgene or carbonyldiimidazole (CDI) in presence or absence of a base such as triethylamine in a solvent such as toluene or dioxane to form the corresponding pyrimidinedione of formula 3. Sonogashira coupling of the bromo or iodo-pyrimidinedione **3** with an appropriately substituted arylacetylene **4** yield the desired ethynyl compounds of general formula I (scheme 1). Introduction of the R² substituent can also be realized at various points in the synthetic sequence via alkylation of the corresponding intermediate where R² = H.

Scheme 2



[0038] Generally speaking, the sequence of steps used to synthesize the compounds of formula I can also be modified in certain cases, for example by first running the Sonogashira coupling with an appropriately substituted bromo or iodo

aniline or aminopyridine **1** with an appropriately substituted arylacetylene **4** to yield the corresponding ethynyl compounds **5**. Reacting **5** with an appropriate aromatic or heteroaromatic orthoaminoester **2** with phosgene or a phosgene equivalent such as triphosgene or carbonyldiimidazole (CDI) in presence or absence of a base such as triethylamine in a solvent such as toluene or dioxane yield the desired ethynyl compounds of general formula **I** (scheme 2). Introduction of the R² substituent can also be realized at various points in the synthetic sequence via alkylation of the corresponding intermediate where R²= H.

Biological Assay and Data:

Determination of EC₅₀ values using a Ca²⁺ mobilization in vitro assay on recombinant human mGlu4 expressed in HEK293 cells:

[0039] A monoclonal HEK-293 cell line stably transfected with a cDNA encoding for the human mGlu4 receptor was generated; for the work with mGlu4 Positive Allosteric Modulators (PAMs), a cell line with low receptor expression levels and low constitutive receptor activity was selected to allow the differentiation of agonistic versus PAM activity. Cells were cultured according to standard protocols (Freshney, 2000) in Dulbecco's Modified Eagle Medium with high glucose supplemented with 1 mM glutamine, 10% (vol/vol) heat-inactivated bovine calf serum, Penicillin/Streptomycin, 50 µg/ml hygromycin and 15 µg/ml blasticidin (all cell culture reagents and antibiotics from Invitrogen, Basel, Switzerland).

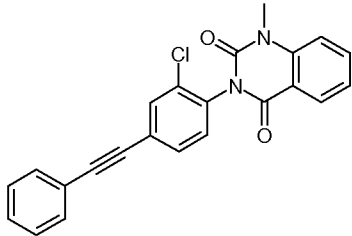
[0040] About 24 hrs before an experiment, 5x10⁴ cells/well were seeded in poly-D-lysine coated, black/clear-bottomed 96-well plates. The cells were loaded with 2.5 µM Fluo-4AM in loading buffer (1xHBSS, 20 mM HEPES) for 1 hr at 37°C and washed five times with loading buffer. The cells were transferred into a Functional Drug Screening System 7000 (Hamamatsu, Paris, France), and 11 half logarithmic serial dilutions of test compound at 37°C were added and the cells were incubated for 10-30 min. with on-line recording of fluorescence. Following this pre-incubation step, the agonist (2S)-2-amino-4-phosphonobutanoic acid (L-AP4) was added to the cells at a concentration corresponding to EC₂₀ with on-line recording of fluorescence; in order to account for day-to-day variations in the responsiveness of cells, the EC₂₀ of L-AP4 was determined immediately ahead of each experiment by recording of a full dose-response curve of L-AP4.

[0041] Responses were measured as peak increase in fluorescence minus basal (i.e. fluorescence without addition of L-AP4), normalized to the maximal stimulatory effect obtained with saturating concentrations of L-AP4. Graphs were plotted with the % maximal stimulatory using XLfit, a curve fitting program that iteratively plots the data using Levenburg Marquardt algorithm. The single site competition analysis equation used was $y = A + ((B-A)/(1+((x/C)^D)))$, where y is the % maximal stimulatory effect, A is the minimum y, B is the maximum y, C is the EC₅₀, x is the log₁₀ of the concentration of the competing compound and D is the slope of the curve (the Hill Coefficient). From these curves the EC₅₀ (drug concentration at which 50% of the maximal receptor activation was achieved), the Hill coefficient as well as the maximal response in % of the maximal stimulatory effect obtained with saturating concentrations of L-AP4 were calculated (see Fig. 1).

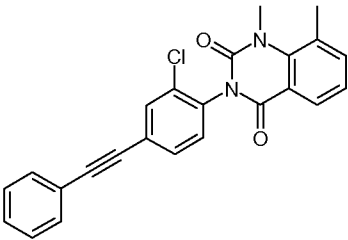
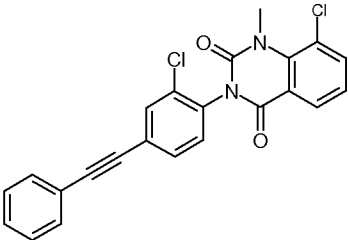
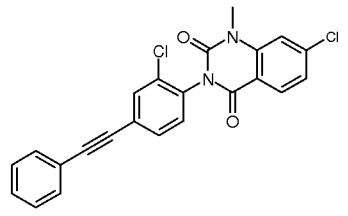
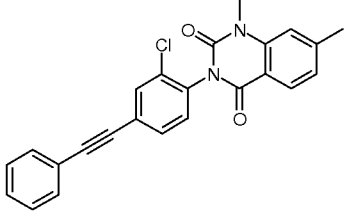
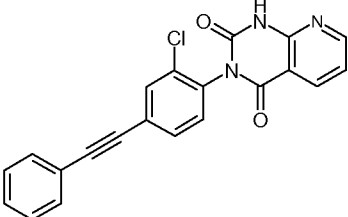
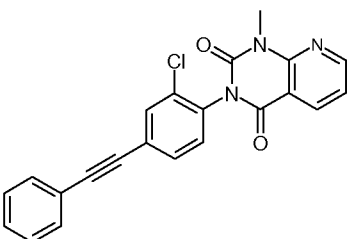
[0042] Positive signals obtained during the pre-incubation with the PAM test compounds (i.e. before application of an EC₂₀ concentration of L-AP4) were indicative of an agonistic activity, the absence of such signals were demonstrating the lack of agonistic activities. A depression of the signal observed after addition of the EC₂₀ concentration of L-AP4 was indicative of an inhibitory activity of the test compound.

[0043] **Fig. 1:** Illustration of the experimental outline for mGlu4 PAM Ca²⁺ mobilization screening assay and the determination of EC₅₀ and % E_{max} values.

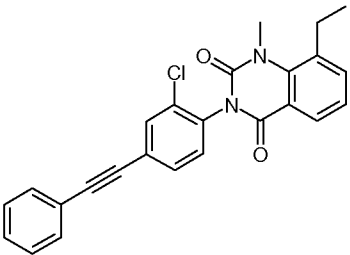
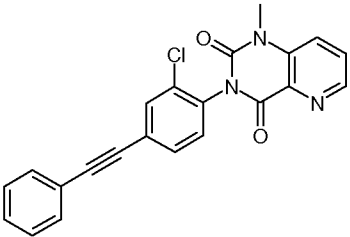
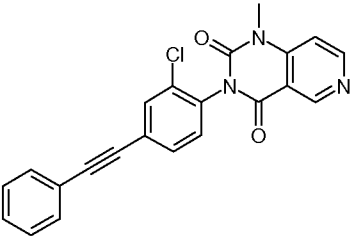
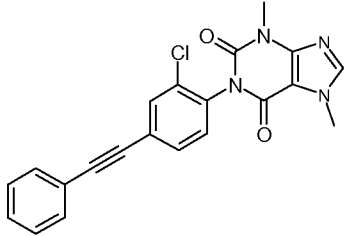
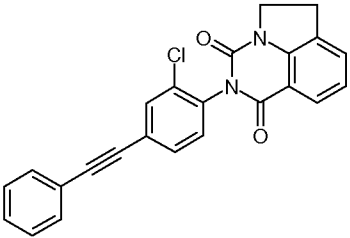
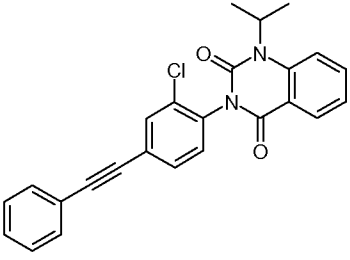
List of Examples and data:

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
1		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione	150	102

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
2		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione	94	99
3		8-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione	137	93
4		7-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione	220	120
5		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione	93	108
6		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[2,3-d]pyrimidine-2,4-dione	179	104
7		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[2,3-d]pyrimidine-2,4-dione	222	101

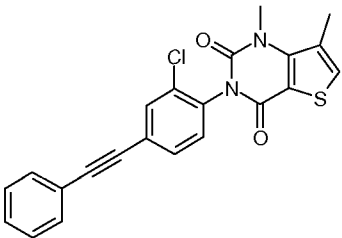
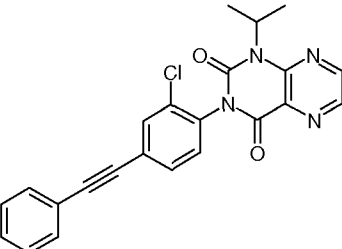
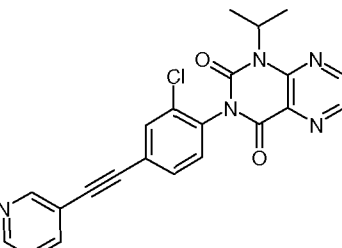
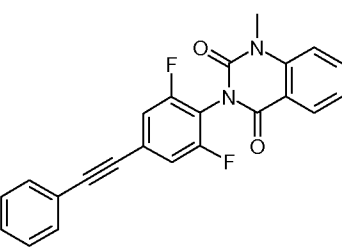
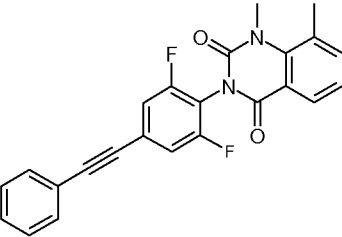
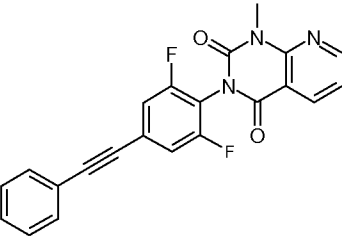
(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
8		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-ethyl-1-methyl-quinazoline-2,4-dione	65	100
9		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione	308	167
10		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[4,3-d]pyrimidine-2,4-dione	399	180
11		1-[2-Chloro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione	175	122
12		2-(2-Chloro-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H) - dione	80	106
13		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-quinazoline-2,4-dione	57	115

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
14		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-ethyl-quinazoline-2,4-dione	172	97
15		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pteridine-2,4-dione	110	111
16		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrimido[4,5-d]pyrimidine-2,4-dione	152	98
17		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrimido[4,5-d]pyrimidine-2,4-dione	143	101
18		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-6-fluoro-1-methyl-quinazoline-2,4-dione	148	86
19		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-7,8-difluoro-1-methyl-quinazoline-2,4-dione	134	73

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
20		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione	50	74
21		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione	52	89
22		3-[2-Chloro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-pteridine-2,4-dione	167	88
23		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione	203	168
24		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione	26	101
25		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[2,3-d]pyrimidine-2,4-dione	89	109

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
26		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 1,7-dimethyl-quinazoline-2,4-dione	115	108
27		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4- dione	16	83
28		3-[2,6-Difluoro-4-[2-(3- pyridyl)ethynyl]phenyl]-1,7-dimethyl- thieno[3,2-d]pyrimidine-2,4-dione	29	86
29		2-(2,6-Difluoro-4-(phenylethynyl)phenyl)- 5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline- 1,3(2H) - dione	52	78
30		2-(2,6-Difluoro-4-(pyridin-3- ylethynyl)phenyl)-5,6-dihydro-1H- pyrrolo[3,2,1-ij]quinazoline-1,3(2H) - dione	91	89
31		6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 4H-thiazolo[4,5-d]pyrimidine-5,7-dione	187	85

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
32		6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 4-methyl-thiazolo[4,5-d]pyrimidine-5,7- dione	33	104
33		6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 2,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7- dione	47	86
34		6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 1,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7- dione	74	85
35		6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 4-isopropyl-2-methyl-pyrazolo[4,3- d]pyrimidine-5,7-dione	30	89
36		6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 4-isopropyl-1-methyl-pyrazolo[4,3- d]pyrimidine-5,7-dione	155	89
37		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 1-isopropyl-pyrido[2,3-d]pyrimidine-2,4- dione	88	73

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
38		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-pyrido[3,2-d]pyrimidine-2,4-dione	108	108
39		6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-methyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione	17	107
40		6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione	23	104
41		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,5-dimethyl-quinazoline-2,4-dione	231	98
42		3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-quinazoline-2,4-dione	83	80
43		3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione	74	107

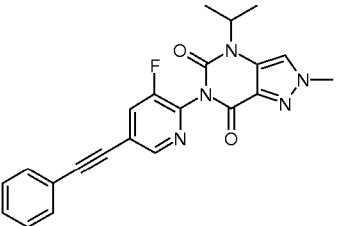
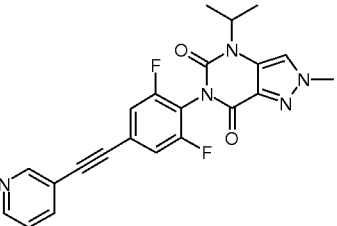
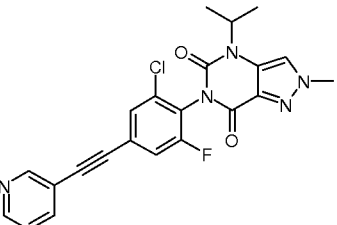
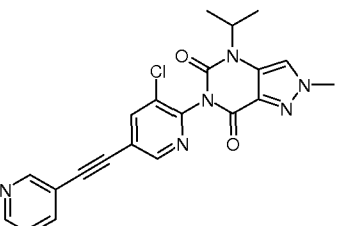
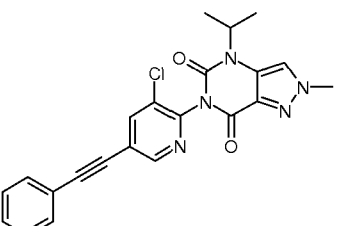
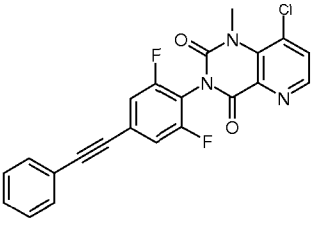
(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
44		3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]- 1-methyl-quinazoline-2,4-dione	372	180
45		3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]- 1-isopropyl-quinazoline-2,4-dione	137	92
46		3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]- 1,8-dimethyl-quinazoline-2,4-dione	141	113
47		2-(3-Chloro-5-(phenylethynyl)pyridin-2-yl)- 5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline- 1,3(2H) - dione	187	110
48		1-Methyl-3-[5-(2-phenylethynyl)- 3-(trifluoromethyl)-2-pyridyl]quinazoline-2,4- dione	175	95
49		3-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]- 1,8-dimethyl-quinazoline-2,4-dione	302	111

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
50		3-[2-Chloro-4-(2-phenylethynyl)phenyl]- 1-(2-methoxyethyl)quinazoline-2,4-dione	499	110
51		6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4- methyl-2-morpholinothiazolo[4,5- d]pyrimidine- 5,7-dione	164	127
52		6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4- methyl-thiazolo[4,5-d]pyrimidine-5,7-dione	200	110
53		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1- cyclobutyl-quinazoline-2,4-dione	102	123
54		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8- isopropyl-1-methyl-quinazoline-2,4-dione	179	97
55		1-Isopropyl-3-[5-(2-phenylethynyl)-2- pyridyl]quinazoline-2,4-dione	766	123

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
56		6-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione	121	120
57		6-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione	65	134
58		6-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione	37	120
59		6-[3-Chloro-5-[2-(3-pyridyl)ethynyl]-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione	169	146
60		6-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione	189	150
61		8-Chloro-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione	91	129

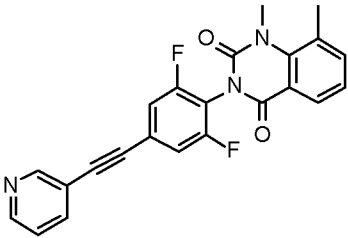
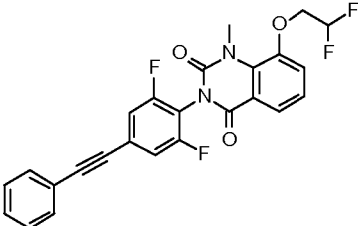
(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
62		5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 2,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6- dione	137	145
63		5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 1,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6- dione	26	146
64		5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 7-isopropyl-2H-pyrazolo[3,4-d]pyrimidine- 4,6-dione	121	120
65		3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8- methoxy-1-methyl-quinazoline-2,4-dione	32	97
66		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 1-(oxetan-3-yl)quinazoline-2,4-dione	75	184
67		6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4- methyl-thiazolo[5,4-d]pyrimidine-5,7-dione	246	188

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
68		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 1-isopropyl-pteridine-2,4-dione	48	225
69		1-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 3,7-dimethyl-purine-2,6-dione	50	113
70		3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]- 8-methoxy-1-methyl-quinazoline-2,4-dione	38	152
71		3-[2-Chloro-6-fluoro-4-[2-(3- pyridyl)ethynyl]phenyl]-1-(oxetan-3- yl)quinazoline-2,4-dione	86	103
72		3-[2-Fluoro-4-(2-phenylethynyl)- 6-(trifluoromethyl)phenyl]-1-isopropyl- quinazoline-2,4-dione	63	134

(continued)

Ex.	Structure	Name	EC ₅₀ (nM) mGlu4PAM	Eff. (%)
73		3-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione	154	197
74		8-(2,2-Difluoroethoxy)-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione	60	122

[0044] The compounds of formula (I) and pharmaceutically acceptable salts thereof can be used as medicaments, e.g. in the form of pharmaceutical preparations. The pharmaceutical preparations can be administered orally, e.g. in the form of tablets, coated tablets, dragées, hard and soft gelatine capsules, solutions, emulsions or suspensions. However, the administration can also be effected rectally, e.g. in the form of suppositories, or parenterally, e.g. in the form of injection solutions.

[0045] The compounds of formula (I) and pharmaceutically acceptable salts thereof can be processed with pharmaceutically inert, inorganic or organic carriers for the production of pharmaceutical preparations. Lactose, corn starch or derivatives thereof, talc, stearic acid or its salts and the like can be used, for example, as such carriers for tablets, coated tablets, dragées and hard gelatin capsules. Suitable carriers for soft gelatin capsules are, for example, vegetable oils, waxes, fats, semi-solid and liquid polyols and the like; depending on the nature of the active substance no carriers are, however, usually required in the case of soft gelatin capsules. Suitable carriers for the production of solutions and syrups are, for example, water, polyols, sucrose, invert sugar, glucose and the like. Adjuvants, such as alcohols, polyols, glycerol, vegetable oils and the like, can be used for aqueous injection solutions of water-soluble salts of compounds of formula (I), but as a rule are not necessary. Suitable carriers for suppositories are, for example, natural or hardened oils, waxes, fats, semi-liquid or liquid polyols and the like.

[0046] In addition, the pharmaceutical preparations can contain preservatives, solubilizers, stabilizers, wetting agents, emulsifiers, sweeteners, colorants, flavorants, salts for varying the osmotic pressure, buffers, masking agents or antioxidants. They can also contain still other therapeutically valuable substances.

[0047] As mentioned earlier, medicaments containing a compound of formula (I) or pharmaceutically acceptable salts thereof and a therapeutically inert excipient are also an object of the present invention, as is a process for the production of such medicaments which comprises bringing one or more compounds of formula I or pharmaceutically acceptable salts thereof and, if desired, one or more other therapeutically valuable substances into a galenical dosage form together with one or more therapeutically inert carriers.

[0048] As further mentioned earlier, the use of the compounds of formula (I) for the preparation of medicaments useful in the prevention and/or the treatment of the above recited diseases is also an object of the present invention.

[0049] The dosage can vary within wide limits and will, of course, be fitted to the individual requirements in each particular case. In general, the effective dosage for oral or parenteral administration is between 0.01-20 mg/kg/day, with a dosage of 0.1-10 mg/kg/day being preferred for all of the indications described. The daily dosage for an adult human being weighing 70 kg accordingly lies between 0.7-1400 mg per day, preferably between 7 and 700 mg per day.

Preparation of pharmaceutical compositions comprising compounds of the invention:

[0050]

Tablets of the following composition are produced in a conventional manner:
mg/Tablet

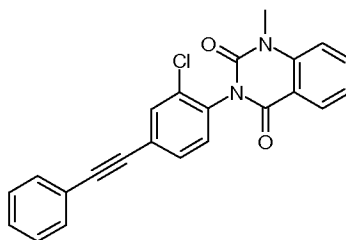
Active ingredient	100
Powdered. lactose	95
White corn starch	35
Polyvinylpyrrolidone	8
Na carboxymethylstarch	10
Magnesium stearate	2
Tablet weight	250

Experimental Section:

Example 1

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione

[0051]



Step 1: 2-Chloro-4-iodo-1-isocyanato-benzene

[0052] 2-Chloro-4-iodoaniline (500 mg, 1.97 mmol) was dissolved in toluene (5 ml) and bis(trichloromethyl) carbonate (234 mg, 0.79 mmol, 0.4 equiv.) was added at room temperature. The mixture was stirred for 16 hours at 110 °C. The reaction mixture was evaporated and the residue was suspended in 5 ml heptane. The suspension was filtered and the filtrate evaporated to dryness. The desired 2-chloro-4-iodo-1-isocyanato-benzene (400 mg, 73 % yield) was obtained as a light yellow solid and used in the next step without further characterization.

Step 2: 3-(2-Chloro-4-iodo-phenyl)-1H-quinazoline-2,4-dione

[0053] Methyl 2-aminobenzoate (100 mg, 0.66 mmol) was dissolved in dioxane (1.5 ml) and 2-chloro-4-iodo-1-isocyanato-benzene (*Example 1, step 1*) (203 mg, 0.73 mmol, 1.1 equiv.) and triethylamine (250 μ l, 1.79 mmol, 2.7 equiv.) were added at room temperature. The mixture was stirred for 16 hours at 85 °C. The reaction mixture was evaporated and the residue was crystallized from heptane and ethyl acetate to yield the desired 3-(2-chloro-4-iodo-phenyl)-1H-quinazoline-2,4-dione (155 mg, 59 % yield) as a white solid, MS: m/e = 399.0/401.0 (M+H⁺).

Step 3: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione

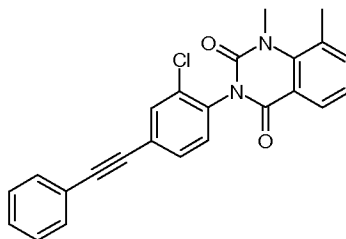
[0054] Bis-(triphenylphosphine)-palladium(II)dichloride (8 mg, 11.7 μ mol, 0.03 equiv.) was dissolved in 2 ml DMF. (155 mg, 389 μ mol) 3-(2-Chloro-4-iodo-phenyl)-1H-quinazoline-2,4-dione (*Example 1, step 2*) and phenylacetylene (79 mg, 85 μ l, 0.778 mmol, 2 equiv.) were added at room temperature. Triethylamine (157 mg, 217 μ l, 1.56 mmol, 4 equiv.), triphenylphosphine (6 mg, 23.4 μ mol, 0.06 equiv.) and copper(I)iodide (2 mg, 7.8 μ mol, 0.02 equiv.) were added and the mixture was stirred for 2 hours at 50°C. The reaction mixture was cooled and extracted with saturated NaHCO₃ solution and two times with ethyl acetate. The organic layers were washed three times with water, dried over sodium sulfate and evaporated to dryness. The crude product was purified by flash chromatography on a silica gel column eluting with an ethyl acetate:heptane gradient 0:100 to 50:50. The desired 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (135 mg, 93 % yield) was obtained as a light yellow solid, MS: m/e = 373.1/375.1 (M+H⁺).

Step 4: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione

[0055] (110 mg, 295 μ mol) 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (*Example 1, step 3*) was dissolved in acetone (3 ml) and potassium carbonate (82 mg, 0.59 mmol, 2 equiv.) and iodomethane (251 mg, 111 μ l, 1.77 mmol, 6 equiv.) were added at room temperature. The mixture was stirred for 16 hours at room temperature. The reaction mixture was evaporated to dryness and the crude product was purified directly by flash chromatography on a silica gel column eluting with an ethyl acetate:heptane gradient 0:100 to 50:50. The desired 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione (92 mg, 81 % yield) was obtained as a light yellow solid, MS: m/e = 387.1/389.2 (M+H⁺).

Example 2**3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione**

[0056]

Step 1: 3-(2-Chloro-4-iodo-phenyl)-8-methyl-1H-quinazoline-2,4-dione

[0057] The title compound was obtained as a white solid, MS: m/e = 413.0/415.0 (M+H⁺), using chemistry similar to that described in Example 1, step 2 from 2-chloro-4-iodo-1-isocyanato-benzene (*Example 1, step 1*) and methyl 2-amino-3-methylbenzoate.

Step 2: 3-[2-chloro-4-(2-phenylethynyl)phenyl]-8-methyl-1H-quinazoline-2,4-dione

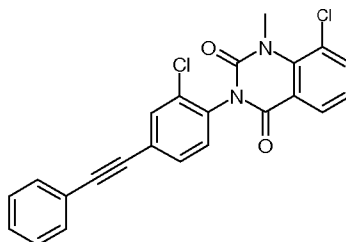
[0058] The title compound was obtained as a light brown solid, MS: m/e = 387.1/389.1 (M+H⁺), using chemistry similar to that described in Example 1, step 3 from 3-(2-chloro-4-iodo-phenyl)-8-methyl-1H-quinazoline-2,4-dione (*Example 2, step 1*) and phenylacetylene.

Step 3: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione

[0059] The title compound was obtained as a light brown solid, MS: m/e = 401.1/403.2 (M+H⁺), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-8-methyl-1H-quinazoline-2,4-dione (*Example 2, step 2*) and iodomethane.

Example 3**8-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione**

[0060]



Step 1: 8-Chloro-3-(2-chloro-4-iodo-phenyl)-1H-quinazoline-2,4-dione

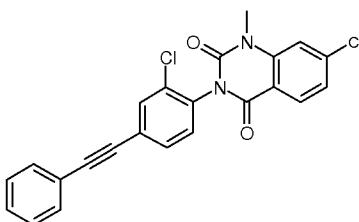
[0061] The title compound was obtained as a light yellow solid, MS: $m/e = 433.0/435.0$ ($M+H^+$), using chemistry similar to that described in Example 1, step 2 from 2-chloro-4-iodo-1-isocyanato-benzene (*Example 1, step 1*) and methyl 2-amino-3-chlorobenzoate.

Step 2: 8-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione

[0062] The title compound was obtained as a light yellow solid, MS: $m/e = 405.2/407.2$ ($M-H^+$), using chemistry similar to that described in Example 1, step 3 from 8-chloro-3-(2-chloro-4-iodophenyl)-1H-quinazoline-2,4-dione (*Example 3, step 1*) and phenylacetylene.

Step 3: 8-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione

[0063] The title compound was obtained as a light brown solid, MS: $m/e = 421.1/423.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 8-chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (*Example 3, step 2*) and iodomethane.

Example 4**7-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione****[0064]**Step 1: 2-Chloro-4-(2-phenylethynyl)aniline

[0065] The title compound was obtained as a yellow solid, MS: $m/e = 228.1/230.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 2-chloro-4-iodoaniline and phenylacetylene.

Step 2: 2-Chloro-1-isocyanato-4-(2-phenylethynyl)benzene

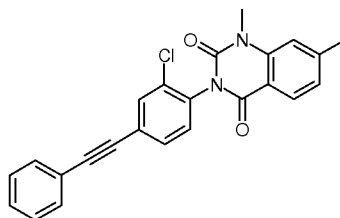
[0066] The title compound was obtained as a dark brown waxy solid, using chemistry similar to that described in Example 1, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*).

Step 3: 7-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione

[0067] The title compound was obtained as a yellow solid, MS: $m/e = 405.2/407.2$ ($M-H^+$), using chemistry similar to that described in Example 1, step 2 from 2-chloro-1-isocyanato-4-(2-phenylethynyl)benzene (*Example 4, step 2*) and methyl 2-amino-4-chlorobenzoate.

Step 3: 7-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione

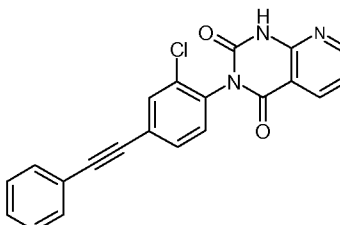
[0068] The title compound was obtained as a yellow solid, MS: $m/e = 421.1/423.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 7-chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (*Example 4, step 3*) and iodomethane.

Example 5**3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione****[0069]**Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-7-methyl-1H-quinazoline-2,4-dione

[0070] The title compound was obtained as a white solid, MS: $m/e = 387.1/389.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 2 from 2-chloro-1-isocyanato-4-(2-phenylethynyl)benzene (*Example 4, step 2*) and methyl 2-amino-4-methylbenzoate.

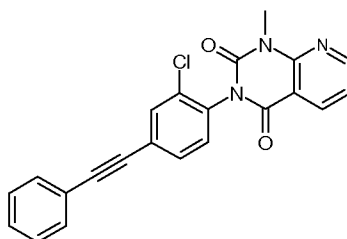
Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione

[0071] The title compound was obtained as a light yellow solid, MS: $m/e = 401.1/403.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-7-methyl-1H-quinazoline-2,4-dione (*Example 5, step 1*) and iodomethane.

Example 6**3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[2,3-d]pyrimidine-2,4-dione****[0072]**

[0073] The title compound was obtained as a light brown solid, MS: $m/e = 374.1/376.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 2 from 2-chloro-1-isocyanato-4-(2-phenylethynyl)benzene (*Example 4, step 2*) and methyl 2-aminonicotinateethyl.

Example 7**3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[2,3-d]pyrimidine-2,4-dione****[0074]**

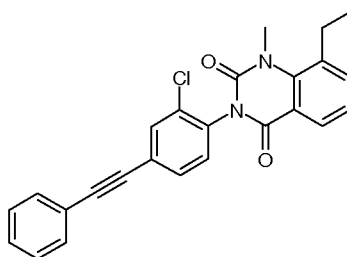


[0075] The title compound was obtained as a light yellow solid, MS: $m/e = 388.1/390.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[2,3-d]pyrimidine-2,4-dione (Example 6) and iodomethane.

Example 8

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-ethyl-1-methyl-quinazoline-2,4-dione

[0076]



Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-ethyl-1H-quinazoline-2,4-dione

[0077] The title compound was obtained as a brown solid, MS: $m/e = 401.2/403.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 2 from 2-chloro-1-isocyanato-4-(2-phenylethynyl)benzene (Example 4, step 2) and methyl 2-amino-3-ethylbenzoate.

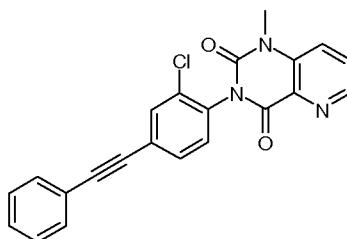
Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-ethyl-1-methyl-quinazoline-2,4-dione

[0078] The title compound was obtained as a light brown solid, MS: $m/e = 415.1/417.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-8-ethyl-1H-quinazoline-2,4-dione (Example 8, step 1) and iodomethane.

Example 9

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione

[0079]

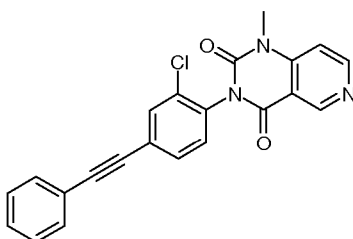


Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[3,2-d]pyrimidine-2,4-dione

[0080] The title compound was obtained as a light brown solid, MS: $m/e = 374.3/376.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 2 from 2-chloro-1-isocyanato-4-(2-phenylethynyl)benzene (*Example 4, step 2*) and ethyl 3-aminopicolinate.

Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione

[0081] The title compound was obtained as a light yellow solid, MS: $m/e = 388.2/390.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[3,2-d]pyrimidine-2,4-dione (*Example 9, step 1*) and iodomethane.

Example 10**3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[4,3-d]pyrimidine-2,4-dione****[0082]**Step 1: 3-(2-Chloro-4-iodo-phenyl)-1H-pyrido[4,3-d]pyrimidine-2,4-dione

[0083] 2-Chloro-4-iodoaniline (200 mg, 0.79 mmol) was dissolved in toluene (2.0 ml) and methyl 4-aminonicotinate (120 mg, 0.79 mmol, 1.0 equiv.) and CDI (154 mg, 0.95 mmol, 1.2 equiv.) were added at room temperature. The mixture was stirred for 16 hours at 110 °C. The reaction mixture was loaded directly onto a silica gel column. The crude product was purified by flash chromatography eluting with an ethyl acetate:heptane gradient 0:100 to 100:0. The desired 3-(2-chloro-4-iodo-phenyl)-1H-pyrido[4,3-d]pyrimidine-2,4-dione (110 mg, 35 % yield) was obtained as a white solid, MS: $m/e = 400.0/402.0$ ($M+H^+$).

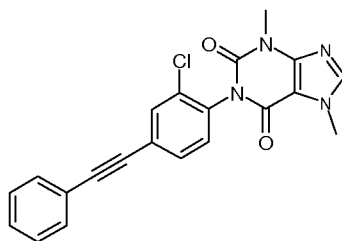
Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[4,3-d]pyrimidine-2,4-dione

[0084] The title compound was obtained as a yellow solid, MS: $m/e = 374.1/376.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 3-(2-chloro-4-iodo-phenyl)-1H-pyrido[4,3-d]pyrimidine-2,4-dione (*Example 10, step 1*) and phenylacetylene.

Step 3: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[4,3-d]pyrimidine-2,4-dione

[0085] The title compound was obtained as a light yellow solid, MS: $m/e = 388.1/390.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[4,3-d]pyrimidine-2,4-dione (*Example 10, step 2*) and iodomethane.

Example 11**1-[2-Chloro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione****[0086]**



Step 1: Ethyl 5-[(2-chloro-4-iodo-phenyl)carbamoylamino]-3-methyl-imidazole-4-carboxylate

[0087] The title compound was obtained as a white solid, MS: $m/e = 449.1/451.1$ ($M-H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-iodoaniline and ethyl 4-amino-1-methyl-1H-imidazole-5-carboxylate.

Step 2: 1-(2-Chloro-4-iodo-phenyl)-7-methyl-3H-purine-2,6-dione

[0088] Ethyl 5-[(2-chloro-4-iodo-phenyl)carbamoylamino]-3-methyl-imidazole-4-carboxylate (*Example 11, step 1*) (330 mg, 0.74 mmol) was dissolved in THF (5.0 ml) and KOtBu (124 mg, 1.1 mmol, 1.5 equiv.) was added at room temperature. The mixture was stirred for 16 hours at room temperature. The reaction mixture was extracted with saturated NaHCO_3 solution and two times with ethyl acetate. The organic layers were washed with saturated NaCl solution, dried over sodium sulfate and evaporated to dryness. The desired 1-(2-chloro-4-iodo-phenyl)-7-methyl-3H-purine-2,6-dione (220 mg, 74 % yield) was obtained as a light yellow solid, MS: $m/e = 403.0/405.0$ ($M+H^+$).

Step 3: 1-[2-Chloro-4-(2-phenylethynyl)phenyl]-7-methyl-3H-purine-2,6-dione

[0089] The title compound was obtained as a light yellow solid, MS: $m/e = 377.1/379.0$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 1-(2-chloro-4-iodo-phenyl)-7-methyl-3H-purine-2,6-dione (*Example 11, step 2*) and phenylacetylene.

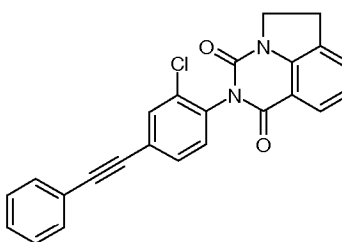
Step 4: 1-[2-Chloro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione

[0090] The title compound was obtained as a yellow solid, MS: $m/e = 391.1/393.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 1-[2-chloro-4-(2-phenylethynyl)phenyl]-7-methyl-3H-purine-2,6-dione (*Example 11, step 3*) and iodomethane.

Example 12

2-(2-Chloro-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione

[0091]

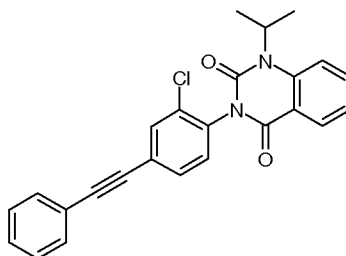


[0092] The title compound was obtained as a white solid, MS: $m/e = 399.1/401.0$ ($M+H^+$), using chemistry similar to that described in Example 1, step 2 from 2-chloro-1-isocyanato-4-(2-phenylethynyl)benzene (*Example 4, step 2*) and methyl indoline-7-carboxylate.

Example 13

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-quinazoline-2,4-dione

[0093]

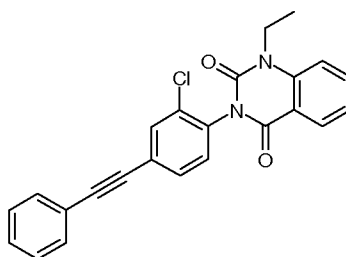


[0094] The title compound was obtained as a yellow oil, MS: $m/e = 415.2/417.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (Example 1, step 3) and 2-iodopropane.

Example 14

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-ethyl-quinazoline-2,4-dione

[0095]

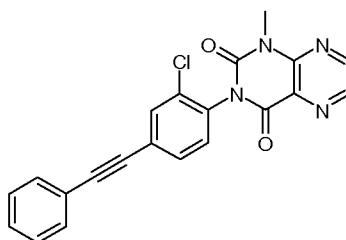


[0096] The title compound was obtained as a yellow oil, MS: $m/e = 401.2/403.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (Example 1, step 3) and iodoethane.

Example 15

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pteridine-2,4-dione

[0097]



Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pteridine-2,4-dione

[0098] The title compound was obtained as a light yellow solid, MS: $m/e = 375.2/377.2$ ($M-H^+$), using chemistry similar

to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 3-aminopyrazine-2-carboxylate.

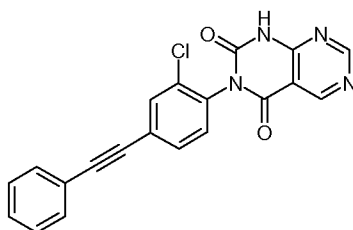
Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pteridine-2,4-dione

[0099] The title compound was obtained as a light yellow solid, MS: $m/e = 389.1/391.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-pteridine-2,4-dione (*Example 15, step 1*) and iodomethane.

Example 16

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrimido[4,5-d]pyrimidine-2,4-dione

[0100]

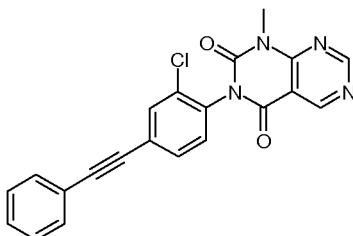


[0101] The title compound was obtained as a yellow solid, MS: $m/e = 375.2/377.1$ ($M-H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 4-aminopyrimidine-5-carboxylate.

Example 17

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrimido[4,5-d]pyrimidine-2,4-dione

[0102]

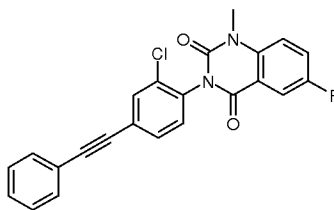


[0103] The title compound was obtained as a light yellow solid, MS: $m/e = 389.2/391.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-pyrimido[4,5-d]pyrimidine-2,4-dione (*Example 16*) and iodomethane.

Example 18

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-6-fluoro-1-methyl-quinazoline-2,4-dione

[0104]



Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-6-fluoro-1H-quinazoline-2,4-dione

[0105] The title compound was obtained as a light brown solid, MS: $m/e = 391.1/393.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 2 from 2-chloro-1-isocyanato-4-(2-phenylethynyl)benzene (*Example 4, step 2*) and methyl 2-amino-5-fluorobenzoate.

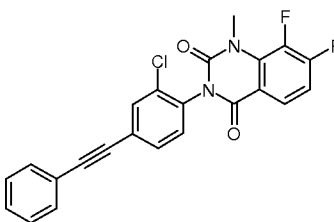
Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-6-fluoro-1-methyl-quinazoline-2,4-dione

[0106] The title compound was obtained as a light yellow solid, MS: $m/e = 405.1/407.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-6-fluoro-1H-quinazoline-2,4-dione (*Example 18, step 1*) and iodomethane.

Example 19

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-7,8-difluoro-1-methyl-quinazoline-2,4-dione

[0107]



Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-7,8-difluoro-1H-quinazoline-2,4-dione

[0108] The title compound was obtained as a yellow solid, MS: $m/e = 407.2/409.2$ ($M-H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 2-amino-3,4-difluorobenzoate.

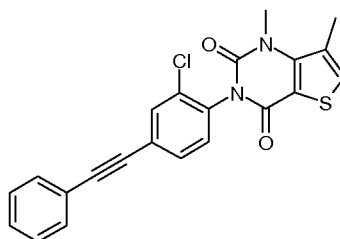
Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-7,8-difluoro-1-methyl-quinazoline-2,4-dione

[0109] The title compound was obtained as a light yellow solid, MS: $m/e = 423.2/425.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-7,8-difluoro-1H-quinazoline-2,4-dione (*Example 19, step 1*) and iodomethane.

Example 20

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione

[0110]



Step 1: 3-(2-Chloro-4-phenylethynyl-phenyl)-7-methyl-1H-thieno[3,2-d]pyrimidine-2,4-dione

[0111] The title compound was obtained as a light brown solid, MS: $m/e = 391.1/393.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 3-amino-4-methylthiophene-2-carboxylate.

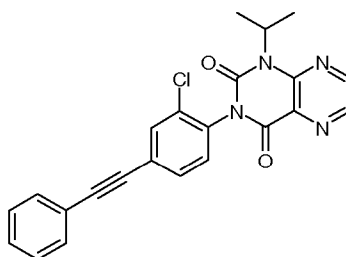
Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione

[0112] The title compound was obtained as an off-white solid, MS: $m/e = 407.2/409.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-(2-chloro-4-phenylethynyl-phenyl)-7-methyl-1H-thieno[3,2-d]pyrimidine-2,4-dione (*Example 20, step 1*) and iodomethane.

Example 21

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione

[0113]



Step 1: 3-(2-Chloro-4-iodo-phenyl)-1H-pteridine-2,4-dione

[0114] The title compound was obtained as a yellow solid, MS: $m/e = 401.1/403.1$ ($M-H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-iodoaniline and methyl 3-aminopyrazine-2-carboxylate.

Step 2: 3-(2-Chloro-4-iodo-phenyl)-1-isopropyl-pteridine-2,4-dione

[0115] The title compound was obtained as a light yellow solid, MS: $m/e = 443.2/445.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-(2-chloro-4-iodo-phenyl)-1H-pteridine-2,4-dione (*Example 21, step 1*) and 2-iodopropane.

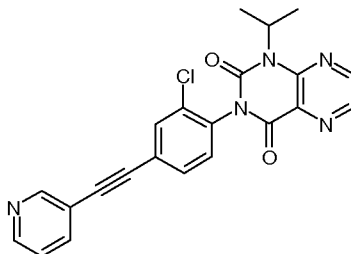
Step 3: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione

[0116] The title compound was obtained as a purple solid, MS: $m/e = 417.3/419.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 3-(2-chloro-4-iodo-phenyl)-1-isopropyl-pteridine-2,4-dione (*Example 21, step 2*) and phenylacetylene.

Example 22

3-[2-Chloro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-pteridine-2,4-dione

[0117]

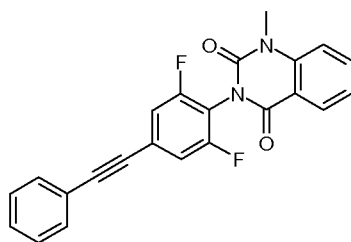


[0118] The title compound was obtained as a white solid, MS: $m/e = 418.3/420.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 3-(2-chloro-4-iodo-phenyl)-1-isopropyl-pteridine-2,4-dione (*Example 21, step 2*) and 3-ethynylpyridine.

Example 23

3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione

[0119]

Step 1: 3-(2,6-Difluoro-4-iodo-phenyl)-1H-quinazoline-2,4-dione

[0120] The title compound was obtained as an off-white solid, MS: $m/e = 401.0$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-iodoaniline and methyl 2-aminobenzoate.

Step 2: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione

[0121] The title compound was obtained as a light brown solid, MS: $m/e = 375.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 3-(2,6-difluoro-4-iodo-phenyl)-1H-quinazoline-2,4-dione (*Example 23, step 1*) and phenylacetylene.

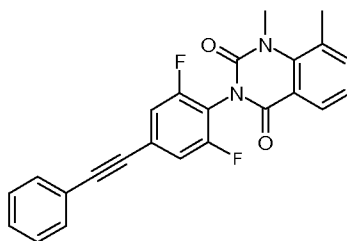
Step 3: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione

[0122] The title compound was obtained as a light brown solid, MS: $m/e = 389.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (*Example 23, step 2*) and iodomethane.

Example 24

3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione

[0123]



Step 1: 3-(2,6-Difluoro-4-iodo-phenyl)-8-methyl-1H-quinazoline-2,4-dione

[0124] The title compound was obtained as an off-white solid, MS: $m/e = 415.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-iodoaniline and methyl 2-amino-3-methylbenzoate.

Step 2: 3-(2,6-Difluoro-4-phenylethynyl-phenyl)-8-methyl-1H-quinazoline-2,4-dione

[0125] The title compound was obtained as a light brown solid, MS: $m/e = 389.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 3-(2,6-difluoro-4-phenylethynyl-phenyl)-8-methyl-1H-quinazoline-2,4-dione (Example 24, step 1) and phenylacetylene.

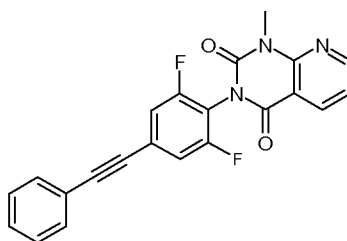
Step 3: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione

[0126] The title compound was obtained as a white solid, MS: $m/e = 403.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-(2,6-difluoro-4-phenylethynyl-phenyl)-8-methyl-1H-quinazoline-2,4-dione (Example 24, step 2) and iodomethane.

Example 25

3-(2,6-Difluoro-4-phenylethynyl-phenyl)-1-methyl-1H-pyrido[2,3-d]pyrimidine-2,4-dione

[0127]



Step 1: 2,6-Difluoro-4-phenylethynyl-phenylamine

[0128] The title compound was obtained as a light brown solid, MS: $m/e = 230.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 2,6-difluoro-4-iodoaniline and phenylacetylene.

Step 2: 3-(2,6-Difluoro-4-phenylethynyl-phenyl)-1H-pyrido[2,3-d]pyrimidine-2,4-dione

[0129] The title compound was obtained as an off-white solid, MS: $m/e = 376.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine and methyl 2-aminonicotinate.

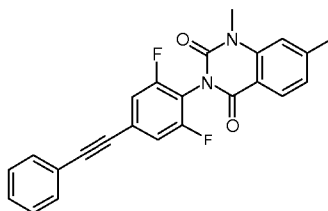
Step 3: 3-(2,6-Difluoro-4-phenylethynyl-phenyl)-1-methyl-1H-pyrido[2,3-d]pyrimidine-2,4-dione

[0130] The title compound was obtained as an off-white solid, MS: $m/e = 390.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-(2,6-difluoro-4-phenylethynyl-phenyl)-1H-pyrido[2,3-d]pyrimidine-2,4-dione (Example 25, step 2) and iodomethane.

Example 26

3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione

[0131]

Step 1: 3-(2,6-Difluoro-4-phenylethynyl-phenyl)-7-methyl-1H-quinazoline-2,4-dione

[0132] The title compound was obtained as an off-white solid, MS: $m/e = 389.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 2-amino-4-methylbenzoate.

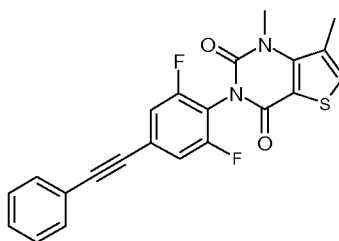
Step 2: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione

[0133] The title compound was obtained as an off-white solid, MS: $m/e = 403.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-(2,6-difluoro-4-phenylethynyl-phenyl)-7-methyl-1H-quinazoline-2,4-dione (*Example 26, step 1*) and iodomethane.

Example 27

3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione

[0134]

Step 1: 3-(2,6-Difluoro-4-phenylethynyl-phenyl)-7-methyl-1H-thieno[3,2-d]pyrimidine-2,4-dione

[0135] The title compound was obtained as an off-white solid, MS: $m/e = 395.0$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 3-amino-4-methylthiophene-2-carboxylate.

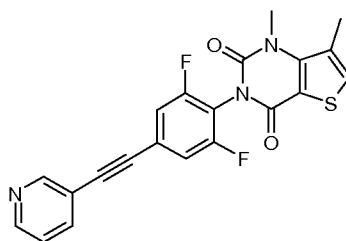
Step 2: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione

[0136] The title compound was obtained as an off-white solid, MS: $m/e = 409.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-(2,6-difluoro-4-phenylethynyl-phenyl)-7-methyl-1H-thieno[3,2-d]pyrimidine-2,4-dione (*Example 27, step 1*) and iodomethane.

Example 28

3-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione

[0137]



Step 1: 2,6-Difluoro-4-pyridin-3-ylethynyl-phenylamine

[0138] The title compound was obtained as a light brown solid, MS: $m/e = 231.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 2,6-difluoro-4-iodoaniline and 3-ethynylpyridine.

Step 2: 3-(2,6-Difluoro-4-pyridin-3-ylethynyl-phenyl)-7-methyl-1H-thieno[3,2-d]pyrimidine-2,4-dione

[0139] The title compound was obtained as an off-white solid, MS: $m/e = 396.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-pyridin-3-ylethynyl-phenylamine (*Example 28, step 1*) and methyl 3-amino-4-methylthiophene-2-carboxylate.

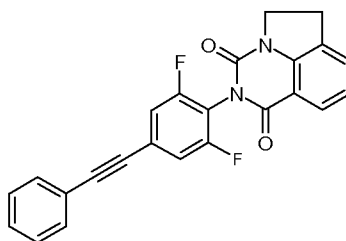
Step 3: 3-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione

[0140] The title compound was obtained as an off-white solid, MS: $m/e = 410.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-(2,6-difluoro-4-pyridin-3-ylethynyl-phenyl)-7-methyl-1H-thieno[3,2-d]pyrimidine-2,4-dione (*Example 28, step 2*) and iodomethane.

Example 29

2-(2,6-Difluoro-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione

[0141]

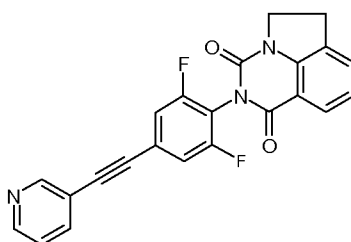


[0142] The title compound was obtained as an off-white solid, MS: $m/e = 401.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methylindoline-7-carboxylate.

Example 30

2-(2,6-Difluoro-4-(pyridin-3-ylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione

[0143]

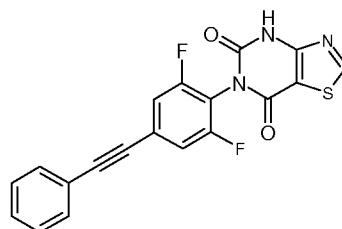


[0144] The title compound was obtained as an off-white solid, MS: $m/e = 402.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-pyridin-3-ylethynyl-phenylamine (*Example 28, step 1*) and methyl indoline-7-carboxylate.

Example 31

6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4H-thiazolo[4,5-d]pyrimidine-5,7-dione

[0145]

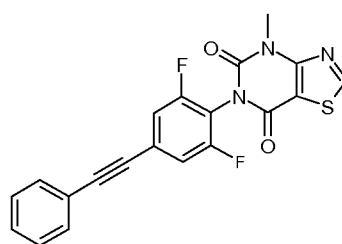


[0146] The title compound was obtained as an off-white solid, MS: $m/e = 380.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 4-aminothiazole-5-carboxylate.

Example 32

6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-methyl-thiazolo[4,5-d]pyrimidine-5,7-dione

[0147]

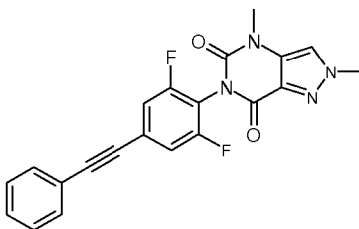


[0148] The title compound was obtained as a white solid, MS: $m/e = 396.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 6-[2,6-difluoro-4-(2 phenylethynyl)-phenyl]-4H-thiazolo[4,5-d]pyrimidine-5,7-dione (*Example 31*) and iodomethane.

Example 33

6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0149]



Step 1: Methyl 4-[[2,6-difluoro-4-(2-phenylethynyl)phenyl]carbamoylamino]-1-methylpyrazole-3-carboxylate

[0150] 2,6-Difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) (200 mg, 0.87 mmol) was dissolved in toluene (6.0 ml) and bis(trichloromethyl) carbonate (104 mg, 0.35 mmol, 0.4 equiv.) was added at room temperature. The mixture was stirred for 1 hour at 110 °C. The mixture was cooled to room temperature and Et₃N (440 mg, 0.61 ml, 4.36 mmol, 5 equiv.) and methyl 4-amino-1-methyl-1H-pyrazole-3-carboxylate (135 mg, 0.87 mmol, 1.0 equiv.) were added at room temperature. The mixture was stirred for 16 hours at 110 °C. The reaction mixture was loaded directly onto a silica gel column. The crude product was purified by flash chromatography eluting with an ethyl acetate:heptane gradient 5:95 to 100:0. The desired methyl 4-[[2,6-difluoro-4-(2-phenylethynyl)phenyl]carbamoylamino]-1-methyl-pyrazole-3-carboxylate (223 mg, 65 % yield) was obtained as a light yellow solid, MS: m/e = 409.4 (M+H⁺).

Step 2: 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2-methyl-4H-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0151] The title compound was obtained as a white solid, MS: m/e = 377.3 (M-H⁺), using chemistry similar to that described in Example 11, step 2 from methyl 4-[[2,6-difluoro-4-(2-phenylethynyl)phenyl]carbamoylamino]-1-methyl-pyrazole-3-carboxylate (*Example 33, step 1*).

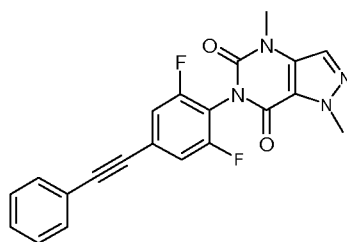
Step 3: 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0152] The title compound was obtained as a yellow solid, MS: m/e = 393.2 (M+H⁺), using chemistry similar to that described in Example 1, step 4 from 6-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-2-methyl-4H-pyrazolo[4,3-d]pyrimidine-5,7-dione (*Example 33, step 2*) and iodomethane.

Example 34

6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0153]



Step 1: 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-4H-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0154] The title compound was obtained as a white solid, MS: m/e = 379.2 (M+H⁺), using chemistry similar to that described in Example 33, step 1 and step 2 starting from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 4-amino-1-methyl-1H-pyrazole-5-carboxylate.

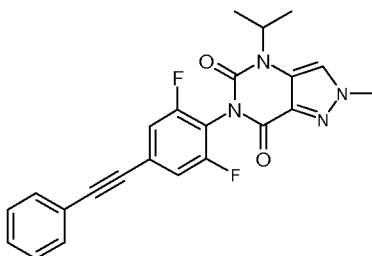
Step 2: 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0155] The title compound was obtained as a white solid, MS: m/e = 393.3 (M+H⁺), using chemistry similar to that described in Example 1, step 4 from 6-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-4H-pyrazolo[4,3-d]pyrimidine-5,7-dione (*Example 34, step 1*) and iodomethane.

Example 35

6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0156]

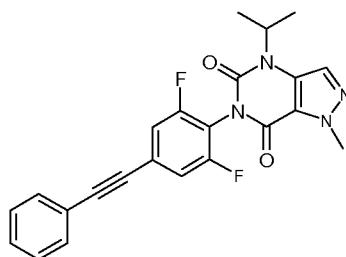


[0157] The title compound was obtained as a white solid, MS: $m/e = 421.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 6-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-2-methyl-4H-pyrazolo[4,3-d]pyrimidine-5,7-dione (Example 33, step 2) and 2-iodopropane.

Example 36

6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-1-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0158]

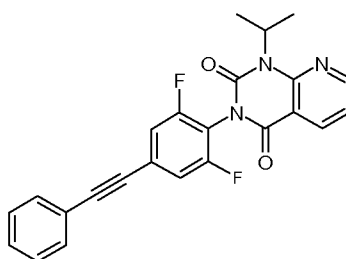


[0159] The title compound was obtained as a yellow solid, MS: $m/e = 421.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 6-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-4H-pyrazolo[4,3-d]pyrimidine-5,7-dione (Example 34, step 1) and 2-iodopropane.

Example 37

3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pyrido[2,3-d]pyrimidine-2,4-dione

[0160]



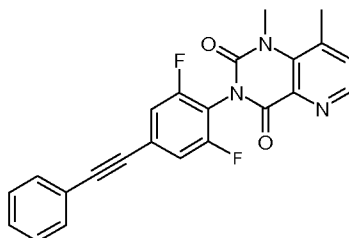
[0161] The title compound was obtained as a white solid, MS: $m/e = 418.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-(2,6-difluoro-4-phenylethynyl-phenyl)-1H-pyrido[2,3-d]pyrimidine-2,4-dione (Ex-

ample 25, step 2) and 2-iodopropane.

Example 38

3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-pyrido[3,2-d]pyrimidine-2,4-dione

[0162]



Step 1: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-8-methyl-1H-pyrido[3,2-d]pyrimidine-2,4-dione

[0163] The title compound was obtained as a yellow solid, MS: $m/e = 388.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (Example 25, step 1) and 3-amino-4-methylpicolinic acid.

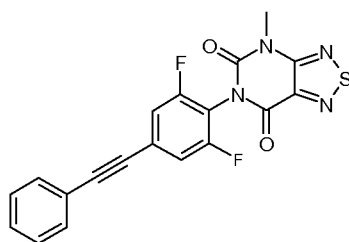
Step 2: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-pyrido[3,2-d]pyrimidine-2,4-dione

[0164] The title compound was obtained as a light yellow solid, MS: $m/e = 404.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-8-methyl-1H-pyrido[3,2-d]pyrimidine-2,4-dione (Example 38, step 1) and iodomethane.

Example 39

6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-methyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione

[0165]

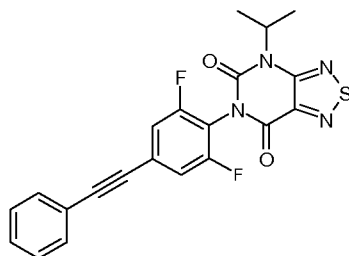


Step 1: 6-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-4H-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione

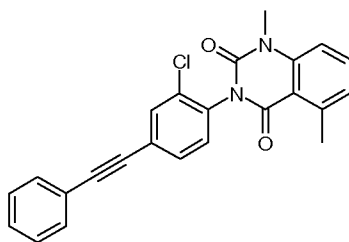
[0166] The title compound was obtained as a yellow solid, MS: $m/e = 381.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (Example 25, step 1) and 4-amino-1,2,5-thiadiazole-3-carboxylic acid.

Step 2: 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-methyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione

[0167] The title compound was obtained as a white solid, MS: $m/e = 397.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 6-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-4H-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione (Example 39, step 1) and iodomethane.

Example 40**6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione****[0168]**

[0169] The title compound was obtained as a white solid, MS: $m/e = 425.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 6-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-4-H-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione (*Example 39, step 1*) and 2-iodopropane.

Example 41**3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,5-dimethyl-quinazoline-2,4-dione****[0170]**

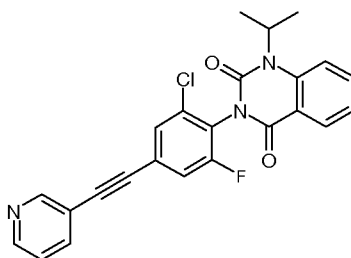
Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-5-methyl-1H-quinazoline-2,4-dione

[0171] The title compound was obtained as a yellow solid, MS: $m/e = 387.1/389.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 2-amino-6-methylbenzoate.

Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,5-dimethyl-quinazoline-2,4-dione

[0172] The title compound was obtained as a light yellow solid, MS: $m/e = 401.2/403.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-5-methyl-1H-quinazoline-2,4-dione (*Example 41, step 1*) and iodomethane.

Example 42**3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-quinazoline-2,4-dione****[0173]**



Step 1: 2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]aniline

[0174] The title compound was obtained as an orange solid, MS: $m/e = 247.1/249.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 2,6-difluoro-4-iodoaniline and 3-ethynylpyridine.

Step 2: Methyl 2-(isopropylamino)benzoate

[0175] Methyl 2-aminobenzoate (2 g, 13.2 mmol) was dissolved in dichloromethane (6.0 ml) and trifluoroacetic acid (2.0 ml, 26.5 mmol, 2.0 equiv.), acetone (2.91 ml, 39.7 mmol, 3 equiv.) and tetramethylammonium triacetoxyborohydride (5.22 g, 19.8 mmol, 1.5 equiv.) were added at room temperature. The mixture was stirred for 16 hour at room temperature. The reaction mixture was extracted with saturated NaHCO_3 solution and twice with dichloromethane. The organic layers were combined and evaporated to dryness. The crude product was purified by flash chromatography on a silica gel column eluting with an ethyl acetate:heptane gradient 0:100 to 20:80. The desired methyl 2-(isopropylamino)benzoate (2.43 g, 90 % yield) was obtained as a colorless oil, MS: $m/e = 194.2$ ($M+H^+$).

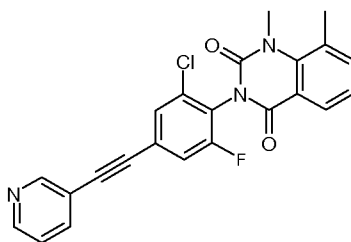
Step 3: 3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-quinazoline-2,4-dione

[0176] The title compound was obtained as a white solid, MS: $m/e = 434.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]aniline (*Example 42, step 1*) and methyl 2-(isopropylamino)benzoate (*Example 42, step 2*).

Example 43

3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione

[0177]

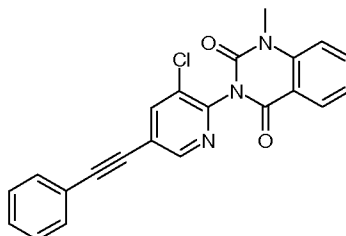


Step 1: 3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-8-methyl-1H-quinazoline-2,4-dione

[0178] The title compound was obtained as a white solid, MS: $m/e = 406.2/408.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]aniline (*Example 42, step 1*) and methyl 2-amino-3-methylbenzoate.

Step 2: 3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione

[0179] The title compound was obtained as a light yellow solid, MS: $m/e = 420.2/422.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-8-methyl-1H-quinazoline-2,4-dione (*Example 43, step 1*) and iodomethane.

Example 44**3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1-methyl-quinazoline-2,4-dione****[0180]**Step 1: 3-Chloro-5-(2-phenylethynyl)pyridin-2-amine

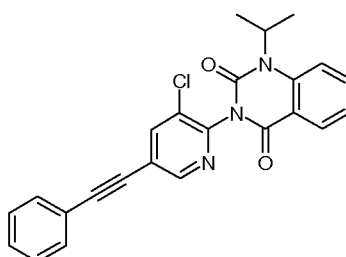
[0181] The title compound was obtained as a yellow solid, MS: $m/e = 229.1/231.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 5-bromo-3-chloropyridin-2-amine and phenylacetylene.

Step 2: 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1H-quinazoline-2,4-dione

[0182] The title compound was obtained as a white solid, MS: $m/e = 374.2/376.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 3-chloro-5-(2-phenylethynyl)pyridin-2-amine (*Example 44, step 1*) and methyl 2-aminobenzoate.

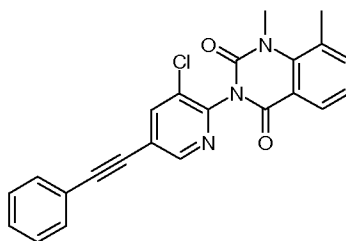
Step 3: 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1-methyl-quinazoline-2,4-dione

[0183] The title compound was obtained as a light yellow solid, MS: $m/e = 388.2/390.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[3-chloro-5-(2-phenylethynyl)-2-pyridyl]-1H-quinazoline-2,4-dione (*Example 44, step 1*) and iodomethane.

Example 45**3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1-isopropyl-quinazoline-2,4-dione****[0184]**

[0185] The title compound was obtained as a white solid, MS: $m/e = 416.3/418.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 3-chloro-5-(2-phenylethynyl)pyridin-2-amine (*Example 44, step 1*) and 2-(isopropylamino)benzoate (*Example 42, step 2*).

Example 46**3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethyl-quinazoline-2,4-dione****[0186]**



Step 1: 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-8-methyl-1H-quinazoline-2,4-dione

[0187] The title compound was obtained as a light yellow solid, MS: $m/e = 388.3/390.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 3-chloro-5-(2-phenylethynyl)pyridin-2-amine (*Example 44, step 1*) and methyl 2-amino-3-methylbenzoate.

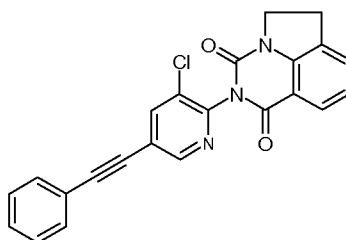
Step 3: 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethyl-quinazoline-2,4-dione

[0188] The title compound was obtained as a light yellow solid, MS: $m/e = 402.2/404.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[3-chloro-5-(2-phenylethynyl)-2-pyridyl]-8-methyl-1H-quinazoline-2,4-dione (*Example 46, step 1*) and iodomethane.

Example 47

2-(3-Chloro-5-(phenylethynyl)pyridin-2-yl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione

[0189]

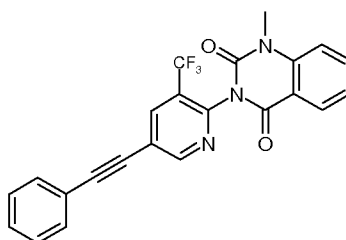


[0190] The title compound was obtained as a light yellow solid, MS: $m/e = 400.1/402.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 3-chloro-5-(2-phenylethynyl)pyridin-2-amine (*Example 44, step 1*) and methyl indoline-7-carboxylate.

Example 48

1-Methyl-3-[5-(2-phenylethynyl)-3-(trifluoromethyl)-2-pyridyl]quinazoline-2,4-dione

[0191]



Step 1: 3-[5-Bromo-3-(trifluoromethyl)-2-pyridyl]-1H-quinazoline-2,4-dione

[0192] The title compound was obtained as a white solid, MS: $m/e = 386.1/388.1$ ($M-H^+$), using chemistry similar to

that described in Example 10, step 1 from 5-bromo-3-(trifluoromethyl)pyridin-2-amine and methyl 2-aminobenzoate.

Step 2: 3-[5-(2-Phenylethynyl)-3-(trifluoromethyl)-2-pyridyl]-1H-quinazoline-2,4-dione

[0193] The title compound was obtained as a light yellow solid, MS: $m/e = 408.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 3-[5-bromo-3-(trifluoromethyl)-2-pyridyl]-1H-quinazoline-2,4-dione (*Example 48, step 1*) and phenylacetylene.

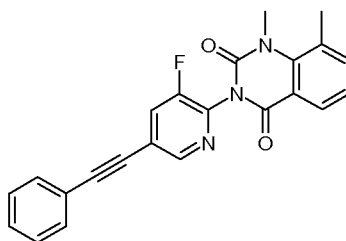
Step 2: 1-Methyl-3-[5-(2-phenylethynyl)-3-(trifluoromethyl)-2-pyridyl]quinazoline-2,4-dione

[0194] The title compound was obtained as a white solid, MS: $m/e = 422.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[5-(2-phenylethynyl)-3-(trifluoromethyl)-2-pyridyl]-1H-quinazoline-2,4-dione (*Example 48, step 2*) and iodomethane.

Example 49

3-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethyl-quinazoline-2,4-dione

[0195]



Step 1: 3-Fluoro-5-(2-phenylethynyl)pyridin-2-amine

[0196] The title compound was obtained as a brown solid, MS: $m/e = 213.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 5-bromo-3-fluoropyridin-2-amine and phenylacetylene.

Step 2: 3-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-8-methyl-1H-quinazoline-2,4-dione

[0197] The title compound was obtained as a light yellow solid, MS: $m/e = 372.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 3-fluoro-5-(2-phenylethynyl)pyridin-2-amine (*Example 49, step 1*) and methyl 2-amino-3-methylbenzoate.

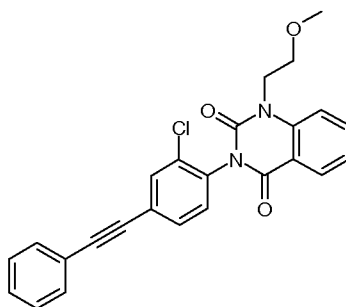
Step 3: 3-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethyl-quinazoline-2,4-dione

[0198] The title compound was obtained as a colorless oil, MS: $m/e = 386.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[3-fluoro-5-(2-phenylethynyl)-2-pyridyl]-8-methyl-1H-quinazoline-2,4-dione (*Example 49, step 1*) and iodomethane.

Example 50

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-(2-methoxyethyl)quinazoline-2,4-dione

[0199]

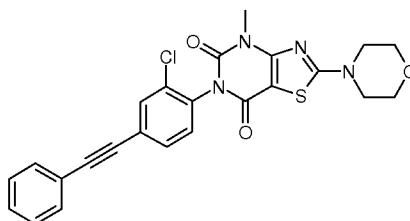


[0200] The title compound was obtained as a yellow solid, MS: $m/e = 431.2/433.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (*Example 1, step 3*) and 1-bromo-2-methoxyethane.

Example 51

6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4-methyl-2-morpholino-thiazolo[4,5-d]pyrimidine-5,7-dione

[0201]



Step 1: Methyl 4-[[2-chloro-4-(2-phenylethynyl)phenyl]carbamoylamino]-2-morpholino-thiazole-5-carboxylate

[0202] The title compound was obtained as a yellow solid, MS: $m/e = 497.3/499.3$ ($M-H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 4-amino-2-morpholinothiazole-5-carboxylate.

Step 2: 6-[2-Chloro-4-(2-phenylethynyl)phenyl]-2-morpholino-4H-thiazolo[4,5-d]pyrimidine-5,7-dione

[0203] The title compound was obtained as a light yellow solid, MS: $m/e = 463.3/465.3$ ($M+H^+$), using chemistry similar to that described in Example 11, step 2 from methyl 4-[[2-chloro-4-(2-phenylethynyl)phenyl]carbamoylamino]-2-morpholino-thiazole-5-carboxylate (*Example 51, step 1*).

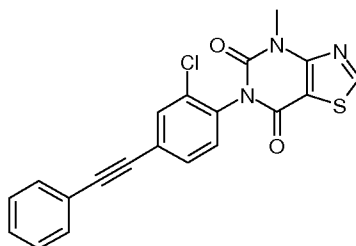
Step 3: 6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4-methyl-2-morpholino-thiazolo[4,5-d]pyrimidine-5,7-dione

[0204] The title compound was obtained as a white solid, MS: $m/e = 479.2/481.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 6-[2-chloro-4-(2-phenylethynyl)phenyl]-2-morpholino-4H-thiazolo[4,5-d]pyrimidine-5,7-dione (*Example 51, step 2*) and iodomethane.

Example 52

6-(2-Chloro-4-phenylethynyl-phenyl)-4-methyl-4H-thiazolo[4,5-d]pyrimidine-5,7-dione

[0205]



Step 1: 6-(2-Chloro-4-phenylethynyl-phenyl)-4H-thiazolo[4,5-d]pyrimidine-5,7-dione

[0206] The title compound was obtained as a light yellow solid, MS: $m/e = 378.2/380.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(phenylethynyl)-aniline (*Example 4, step 1*) and methyl 4-aminothiazole-5-carboxylate.

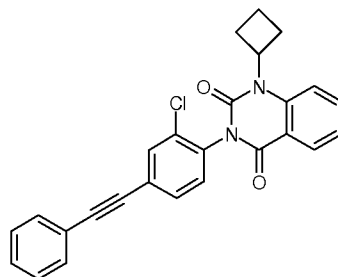
Step 2: 6-(2-Chloro-4-phenylethynyl-phenyl)-4-methyl-4H-thiazolo[4,5-d]pyrimidine-5,7-dione

[0207] The title compound was obtained as an off-white solid, MS: $m/e = 394.1/396.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 6-(2-chloro-4-phenylethynyl-phenyl)-4H-thiazolo[4,5-d]pyrimidine-5,7-dione (*Example 52, step 1*) and iodomethane.

Example 53

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-cyclobutyl-quinazoline-2,4-dione

[0208]

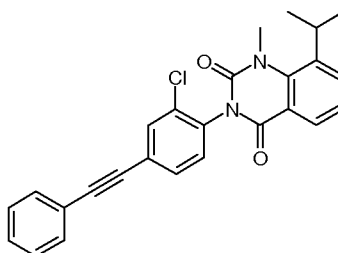


[0209] The title compound was obtained as a white foam, MS: $m/e = 427.2/429.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 2-(cyclobutylamino)benzoate."

Example 54

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-isopropyl-1-methyl-quinazoline-2,4-dione

[0210]

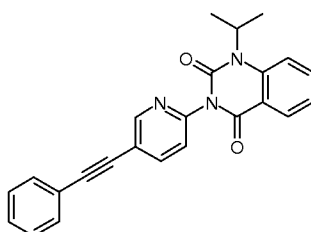


Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-isopropyl-1H-quinazoline-2,4-dione

[0211] The title compound was obtained as a yellow solid, MS: $m/e = 415.2/417.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 2-amino-3-isopropylbenzoate.

Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-isopropyl-1-methyl-quinazoline-2,4-dione

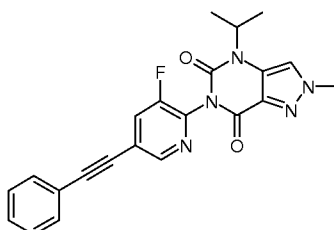
[0212] The title compound was obtained as a light yellow solid, MS: $m/e = 429.2/431.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-8-isopropyl-1H-quinazoline-2,4-dione (*Example 54, step 1*) and iodomethane.

Example 55**1-Isopropyl-3-[5-(2-phenylethynyl)-2-pyridyl]quinazoline-2,4-dione****[0213]**Step 1: 3-(5-Bromo-2-pyridyl)-1-isopropyl-quinazoline-2,4-dione

[0214] The title compound was obtained as a white solid, MS: $m/e = 360.1/362.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 5-bromopyridin-2-amine and methyl 2-(isopropylamino)benzoate (*Example 42, step 2*).

Step 2: 1-Isopropyl-3-[5-(2-phenylethynyl)-2-pyridyl]quinazoline-2,4-dione

[0215] The title compound was obtained as a yellow solid, MS: $m/e = 382.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 3-(5-bromo-2-pyridyl)-1-isopropyl-quinazoline-2,4-dione (*Example 55, step 1*) and phenylacetylene.

Example 56**6-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione****[0216]**Step 1: Methyl 4-(isopropylamino)-1-methyl-pyrazole-3-carboxylate

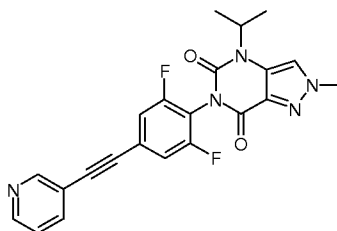
[0217] The title compound was obtained as a light blue oil, MS: $m/e = 198.2$ ($M+H^+$), using chemistry similar to that described in Example 42, step 2 from methyl 4-amino-1-methyl-1H-pyrazole-3-carboxylate.

Step 2: 6-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0218] The title compound was obtained as a white solid, MS: $m/e = 404.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 3-fluoro-5-(2-phenylethynyl)pyridin-2-amine (*Example 49, step 1*) and methyl 4-(isopropylamino)-1-methyl-pyrazole-3-carboxylate (*Example 56, step 1*).

Example 57**6-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione**

[0219]

Step 1: 2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]aniline

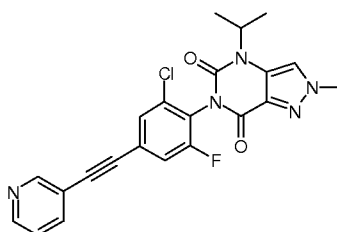
[0220] The title compound was obtained as a yellow solid, MS: $m/e = 231.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 2,6-difluoro-4-iodoaniline and 3-ethynylpyridine.

Step 2: 6-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0221] The title compound was obtained as a white solid, MS: $m/e = 422.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-[2-(3-pyridyl)ethynyl]aniline (*Example 57, step 1*) and methyl 4-(isopropylamino)-1-methyl-pyrazole-3-carboxylate (*Example 56, step 1*).

Example 58**6-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione**

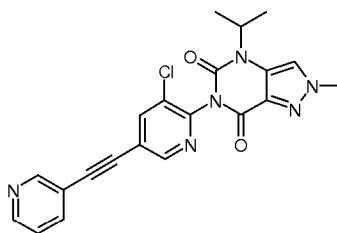
[0222]



[0223] The title compound was obtained as a white solid, MS: $m/e = 438.3/440.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]aniline (*Example 42, step 1*) and 4-(isopropylamino)-1-methyl-pyrazole-3-carboxylate (*Example 56, step 1*).

Example 59**6-[3-Chloro-5-[2-(3-pyridyl)ethynyl]-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione**

[0224]



Step 1: 6-(5-bromo-3-chloro-2-pyridyl)-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0225] The title compound was obtained as a white solid, MS: $m/e = 398.1/400.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 5-bromo-3-chloropyridin-2-amine and 4-(isopropylamino)-1-methyl-pyrazole-3-carboxylate (Example 56, step 1).

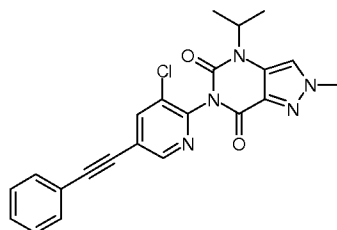
Step 2: 6-[3-Chloro-5-[2-(3-pyridyl)ethynyl]-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0226] The title compound was obtained as a light brown solid, MS: $m/e = 421.2/423.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 6-(5-bromo-3-chloro-2-pyridyl)-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione (Example 59, step 1) and 3-ethynylpyridine.

Example 60

6-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

[0227]

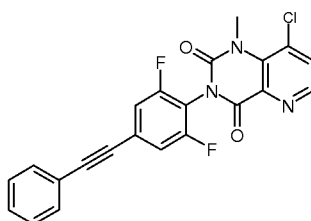


[0228] The title compound was obtained as a light brown solid, MS: $m/e = 420.3/422.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 6-(5-bromo-3-chloro-2-pyridyl)-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione (Example 59, step 1) and phenylacetylene.

Example 61

8-Chloro-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione

[0229]



Step 1: 8-Chloro-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1H-pyrido[3,2-d]pyrimidine-2,4-dione

[0230] The title compound was obtained as a white solid, MS: $m/e = 410.3/412.3$ ($M+H^+$), using chemistry similar to

that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and 3-amino-4-chloropicolinic acid.

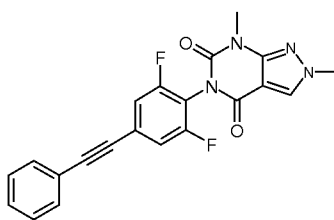
Step 2: 8-Chloro-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione

[0231] The title compound was obtained as a white solid, MS: $m/e = 424.2/426.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 8-chloro-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1H-pyrido[3,2-d]pyrimidine-2,4-dione (*Example 61, step 1*) and iodomethane.

Example 62

5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6-dione

[0232]



Step 1: Methyl 3-[[2,6-difluoro-4-(2-phenylethynyl)phenyl]carbamoylamino]-1-methyl-pyrazole-4-carboxylate

[0233] The title compound was obtained as a light yellow solid, MS: $m/e = 411.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 3-amino-1-methyl-1H-pyrazole-4-carboxylate.

Step 2: 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2-methyl-7H-pyrazolo[3,4-d]pyrimidine-4,6-dione

[0234] The title compound was obtained as a white solid, MS: $m/e = 379.3$ ($M+H^+$), using chemistry similar to that described in Example 11, step 2 from methyl 3-[[2,6-difluoro-4-(2-phenylethynyl)phenyl]carbamoylamino]-1-methyl-pyrazole-4-carboxylate (*Example 62, step 1*).

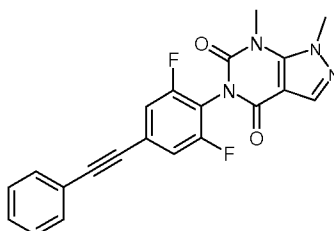
Step 3: 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6-dione

[0235] The title compound was obtained as a white solid, MS: $m/e = 393.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 5-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-2-methyl-7H-pyrazolo[3,4-d]pyrimidine-4,6-dione (*Example 62, step 2*) and iodomethane.

Example 63

5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6-dione

[0236]



Step 1: Methyl 5-[[2,6-difluoro-4-(2-phenylethynyl)phenyl]carbamoylamino]-1-methyl-pyrazole-4-carboxylate

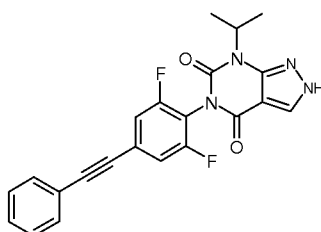
[0237] The title compound was obtained as a light yellow solid, MS: $m/e = 411.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 5-amino-1-methyl-1H-pyrazole-4-carboxylate.

Step 2: 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-7H-pyrazolo[3,4-d]pyrimidine-4,6-dione

[0238] The title compound was obtained as a white solid, MS: $m/e = 379.3$ ($M+H^+$), using chemistry similar to that described in Example 11, step 2 from methyl 5-[[2,6-difluoro-4-(2-phenylethynyl)phenyl]carbamoylamino]-1-methyl-pyrazole-4-carboxylate (*Example 63, step 1*).

Step 3: 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6-dione

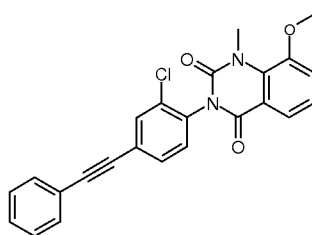
[0239] The title compound was obtained as a white solid, MS: $m/e = 393.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 5-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-7H-pyrazolo[3,4-d]pyrimidine-4,6-dione (*Example 63, step 2*) and iodomethane.

Example 64**5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-7-isopropyl-2H-pyrazolo[3,4-d]pyrimidine-4,6-dione****[0240]**Step 1: Methyl 3-(isopropylamino)-1H-pyrazole-4-carboxylate

[0241] The title compound was obtained as a light blue oil, MS: $m/e = 184.2$ ($M+H^+$), using chemistry similar to that described in Example 42, step 2 from methyl 3-amino-1H-pyrazole-4-carboxylate.

Step 2: 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-7-isopropyl-2H-pyrazolo[3,4-d]pyrimidine-4,6-dione

[0242] The title compound was obtained as a white solid, MS: $m/e = 407.5$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 3-(isopropylamino)-1H-pyrazole-4-carboxylate (*Example 64, step 1*).

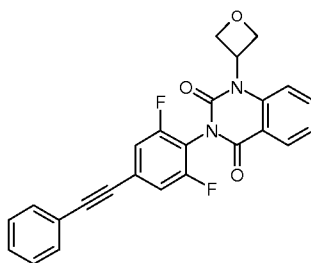
Example 65**3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methyl-quinazoline-2,4-dione****[0243]**

Step 1: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-methoxy-1H-quinazoline-2,4-dione

[0244] The title compound was obtained as a yellow solid, MS: $m/e = 403.2/405.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and methyl 2-amino-3-methoxybenzoate.

Step 2: 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methyl-quinazoline-2,4-dione

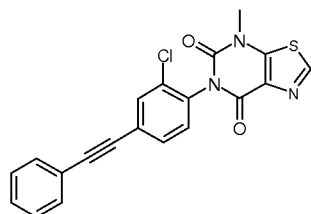
[0245] The title compound was obtained as a light yellow solid, MS: $m/e = 417.2/419.1$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2-chloro-4-(2-phenylethynyl)phenyl]-8-methoxy-1H-quinazoline-2,4-dione (*Example 65, step 1*) and iodomethane.

Example 66**3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-(oxetan-3-yl)quinazoline-2,4-dione****[0246]**Step 1: Methyl 2-(oxetan-3-ylamino)benzoate

[0247] The title compound was obtained as a colorless oil, MS: $m/e = 208.2$ ($M+H^+$), using chemistry similar to that described in Example 42, step 2 from methyl 2-aminobenzoate and oxetan-3-one.

Step 2: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-(oxetan-3-yl)quinazoline-2,4-dione

[0248] The title compound was obtained as a white solid, MS: $m/e = 431.3$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 2-(oxetan-3-ylamino)benzoate (*Example 65, step 1*).

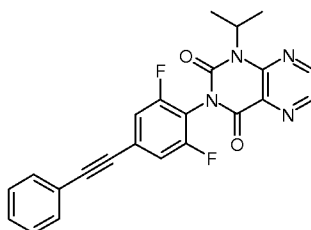
Example 67**6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4-methyl-thiazolo[5,4-d]pyrimidine-5,7-dione****[0249]**

[0250] The title compound was obtained as a light yellow solid, MS: $m/e = 394.1/396.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-4-(2-phenylethynyl)aniline (*Example 4, step 1*) and ethyl 5-(methylamino)thiazole-4-carboxylate.

Example 68

3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione

[0251]



Step 1: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1H-pteridine-2,4-dione

[0252] The title compound was obtained as a white solid, MS: $m/e = 377.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (Example 25, step 1) and methyl 3-aminopyrazine-2-carboxylate.

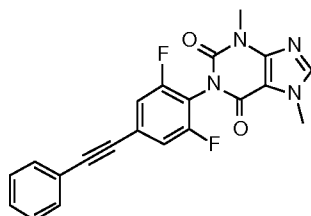
Step 2: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione

[0253] The title compound was obtained as a white solid, MS: $m/e = 419.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1H-pteridine-2,4-dione (Example 68, step 1) and 2-iodopropane.

Example 69

1-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione

[0254]



Step 1: Ethyl 5-[(2,6-difluoro-4-iodo-phenyl)carbamoylamino]-3-methyl-imidazole-4-carboxylate

[0255] The title compound was obtained as a white solid, MS: $m/e = 425.3$ ($M-H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (Example 25, step 1) and ethyl 4-amino-1-methyl-1H-imidazole-5-carboxylate.

Step 2: 1-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-7-methyl-3H-purine-2,6-dione

[0256] The title compound was obtained as a light yellow solid, MS: $m/e = 379.2$ ($M+H^+$), using chemistry similar to that described in Example 11, step 2 from ethyl 5-[(2,6-difluoro-4-iodophenyl)carbamoylamino]-3-methyl-imidazole-4-carboxylate (Example 69, step 1) and by using sodium hydride instead of KOTBu.

Step 3: 1-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione

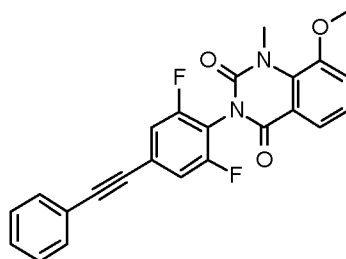
[0257] The title compound was obtained as a yellow solid, MS: $m/e = 393.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 1-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-7-methyl-3H-purine-2,6-dione (Example

69, step 2) and iodomethane.

Example 70

3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methyl-quinazoline-2,4-dione

[0258]



Step 1: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-8-methoxy-1H-quinazoline-2,4-dione

[0259] The title compound was obtained as a white solid, MS: $m/e = 403.2$ ($M-H$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 2-amino-3-methoxybenzoate.

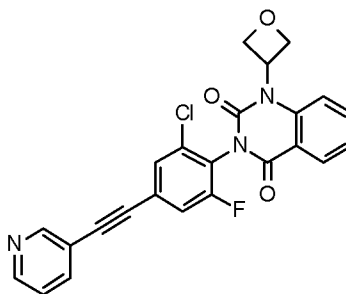
Step 2: 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methyl-quinazoline-2,4-dione

[0260] The title compound was obtained as a white solid, MS: $m/e = 419.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-8-methoxy-1H-quinazoline-2,4-dione (*Example 70, step 1*) and iodomethane.

Example 71

3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-(oxetan-3-yl)quinazoline-2,4-dione

[0261]

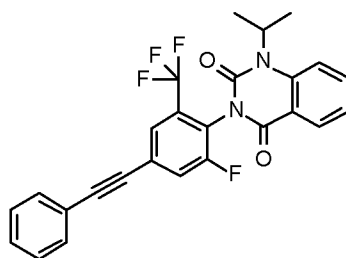


[0262] The title compound was obtained as a white solid, MS: $m/e = 448.2/450.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2-chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]aniline (*Example 42, step 1*) and methyl 2-(oxetan-3-ylamino)benzoate (*Example 65, step 1*).

Example 72

3-[2-Fluoro-4-(2-phenylethynyl)-6-(trifluoromethyl)phenyl]-1-isopropyl-quinazoline-2,4-dione

[0263]



Step 1: 3-[4-Bromo-2-fluoro-6-(trifluoromethyl)phenyl]-1-isopropyl-quinazoline-2,4-dione

[0264] The title compound was obtained as a colorless oil, MS: $m/e = 445.2/447.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 4-bromo-2-fluoro-6-(trifluoromethyl)aniline and methyl 2-(isopropylamino)benzoate (Example 42, step 2).

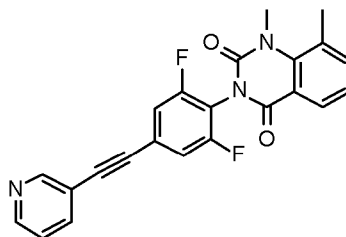
Step 2: 3-[2-Fluoro-4-(2-phenylethynyl)-6-(trifluoromethyl)phenyl]-1-isopropyl-quinazoline-2,4-dione

[0265] The title compound was obtained as a colorless oil, MS: $m/e = 468.3$ ($M+H^+$), using chemistry similar to that described in Example 1, step 3 from 3-[4-bromo-2-fluoro-6-(trifluoromethyl)phenyl]-1-isopropyl-quinazoline-2,4-dione (Example 72, step 1) and phenylacetylene.

Example 73

3-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione

[0266]

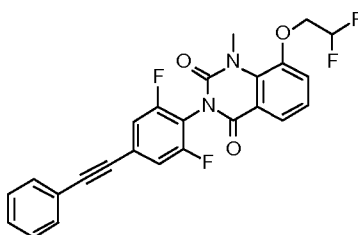


[0267] The title compound was obtained as a white solid, MS: $m/e = 404.2$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-pyridin-3-ylethynyl-phenylamine (Example 28, step 1) and methyl 3-methyl-2-(methylamino)benzoate.

Example 74

8-(2,2-Difluoroethoxy)-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione

[0268]



Step 1: 8-(2,2-Difluoroethoxy)-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione

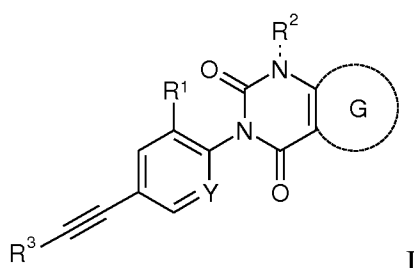
[0269] The title compound was obtained as a yellow solid, MS: $m/e = 455.1$ ($M+H^+$), using chemistry similar to that described in Example 10, step 1 from 2,6-difluoro-4-phenylethynyl-phenylamine (*Example 25, step 1*) and methyl 2-amino-3-(2,2-difluoroethoxy)benzoate.

Step 2: 8-(2,2-Difluoroethoxy)-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methylquinazoline-2,4-dione

[0270] The title compound was obtained as a white solid, MS: $m/e = 469.2$ ($M+H^+$), using chemistry similar to that described in Example 1, step 4 from 8-(2,2-difluoroethoxy)-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1H-quinazoline-2,4-dione (*Example 74, step 1*) and iodomethane.

Claims

1. Compounds of formula I



wherein

Y is N or C-R¹;

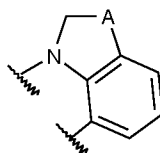
G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, C₁₋₇-alkyl, C₁₋₇-alkoxy, C₁₋₇-alkoxy substituted by halogen or NRR'; R and R' are independently from each other hydrogen or C₁₋₇-alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N-C₁₋₇-alkyl group;

R¹ is hydrogen, halogen or C₁₋₇-alkyl substituted by halogen;

R¹' is hydrogen, halogen or C₁₋₇-alkyl substituted by halogen;

R² is hydrogen, C₁₋₇-alkyl, C₁₋₇-alkoxyalkyl, cycloalkyl or heterocycloalkyl;

or R² may form together with the closest carbon atom in group G a group

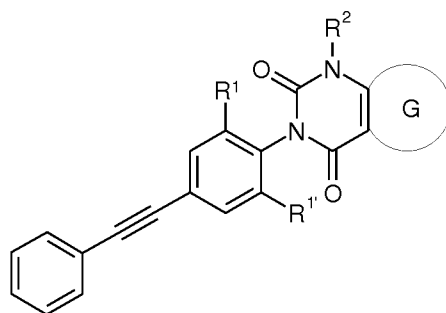


for A being -CH₂-, -CH₂CH₂-, or -C(CH₃)₂-,

R³ is phenyl or pyridinyl, wherein the N atom in the pyridinyl group may be in different positions;

or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof.

2. A compound of formula IA according to claim 1,



IA

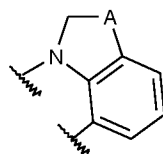
wherein

G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, C₁₋₇-alkyl, C₁₋₇-alkoxy, C₁₋₇-alkoxy substituted by halogen or NRR'; R and R' are independently from each other hydrogen or C₁₋₇-alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N- C₁₋₇-alkyl group;

R¹ is hydrogen, halogen or C₁₋₇-alkyl substituted by halogen;

R^{1'} is hydrogen, halogen or C₁₋₇-alkyl substituted by halogen;

R² is hydrogen, C₁₋₇-alkyl, C₁₋₇-alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group



for A being -CH₂-, -CH₂CH₂-, or -C(CH₃)₂-,

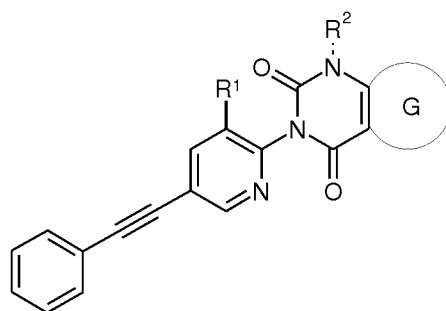
or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof.

3. A compound of formula IA according to claim 2, wherein the compounds are

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione
 8-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione
 7-Chloro-3-[2-chloro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrido[2,3-d]pyrimidine-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[2,3-d]pyrimidine-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-ethyl-1-methyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[4,3-d]pyrimidine-2,4-dione
 1-[2-Chloro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione
 2-(2-Chloro-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-ethyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pteridine-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1H-pyrimido[4,5-d]pyrimidine-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrimido[4,5-d]pyrimidine-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-6-fluoro-1-methyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-7,8-difluoro-1-methyl-quinazoline-2,4-dione

3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-quinazoline-2,4-dione
 5 3-(2,6-Difluoro-4-phenylethynyl-phenyl)-1-methyl-1H-pyrido[2,3-d]pyrimidine-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-quinazoline-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione
 2-(2,6-Difluoro-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4H-thiazolo[4,5-d]pyrimidine-5,7-dione
 10 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-methyl-thiazolo[4,5-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,4-dimethyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-1-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 15 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pyrido[2,3-d]pyrimidine-2,4-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,8-dimethyl-pyrido[3,2-d]pyrimidine-2,4-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-methyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione
 6-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-4-isopropyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1,5-dimethyl-quinazoline-2,4-dione
 20 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-(2-methoxyethyl)quinazoline-2,4-dione
 6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4-methyl-2-morpholino-thiazolo[4,5-d]pyrimidine-5,7-dione
 6-(2-Chloro-4-phenylethynyl-phenyl)-4-methyl-4H-thiazolo[4,5-d]pyrimidine-5,7-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-1-cyclobutyl-quinazoline-2,4-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-isopropyl-1-methyl-quinazoline-2,4-dione
 25 8-Chloro-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-pyrido[3,2-d]pyrimidine-2,4-dione
 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-2,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6-dione
 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1,7-dimethyl-pyrazolo[3,4-d]pyrimidine-4,6-dione
 5-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-7-isopropyl-2H-pyrazolo[3,4-d]pyrimidine-4,6-dione
 3-[2-Chloro-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methyl-quinazoline-2,4-dione
 30 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-(oxetan-3-yl)quinazoline-2,4-dione
 6-[2-Chloro-4-(2-phenylethynyl)phenyl]-4-methyl-thiazolo[5,4-d]pyrimidine-5,7-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-1-isopropyl-pteridine-2,4-dione
 1-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-3,7-dimethyl-purine-2,6-dione
 3-[2,6-Difluoro-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methyl-quinazoline-2,4-dione
 35 3-[2-Fluoro-4-(2-phenylethynyl)-6-(trifluoromethyl)phenyl]-1-isopropyl-quinazoline-2,4-dione or
 8-(2,2-Difluoroethoxy)-3-[2,6-difluoro-4-(2-phenylethynyl)phenyl]-1-methyl-quinazoline-2,4-dione.

4. A compound of formula IB according to claim 1



IB

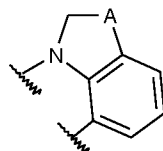
wherein

G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the
 group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl,
 thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the
 group consisting of halogen, C₁₋₇-alkyl, C₁₋₇-alkoxy, C₁₋₇-alkoxy substituted by halogen or NRR'; R and R' are
 independently from each other hydrogen or C₁₋₇-alkyl, or may form together with the N atom to which they are

attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N- C₁₋₇-alkyl group;

R¹ is hydrogen, halogen or C₁₋₇-alkyl substituted by halogen;

R² is hydrogen, C₁₋₇-alkyl, C₁₋₇-alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group



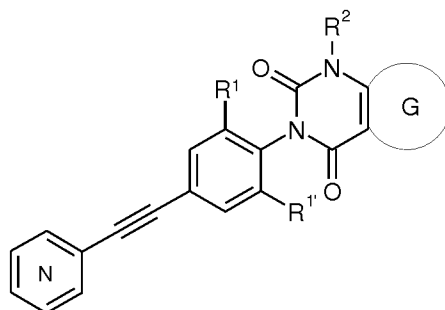
for A being -CH₂-, -CH₂CH₂-, or -C(CH₃)₂-,

or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof.

5. A compound of formula IB according to claim 4, which compounds are

- 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1-methyl-quinazoline-2,4-dione
- 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1-isopropyl-quinazoline-2,4-dione
- 3-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethyl-quinazoline-2,4-dione
- 2-(3-Chloro-5-(phenylethynyl)pyridin-2-yl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
- 1-Methyl-3-[5-(2-phenylethynyl)-3-(trifluoromethyl)-2-pyridyl]quinazoline-2,4-dione
- 3-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethyl-quinazoline-2,4-dione
- 1-Isopropyl-3-[5-(2-phenylethynyl)-2-pyridyl]quinazoline-2,4-dione
- 6-[3-Fluoro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione or
- 6-[3-Chloro-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione.

6. A compound of formula IC according to claim 1,



IC,

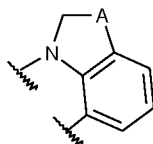
wherein

G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, C₁₋₇-alkyl, C₁₋₇-alkoxy, C₁₋₇-alkoxy substituted by halogen or NRR'; R and R' are independently from each other hydrogen or C₁₋₇-alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N- C₁₋₇-alkyl group;

R¹ is hydrogen, halogen or C₁₋₇-alkyl substituted by halogen;

R^{1'} is hydrogen, halogen or C₁₋₇-alkyl substituted by halogen;

R² is hydrogen, C₁₋₇-alkyl, C₁₋₇-alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group



for A being $-\text{CH}_2-$, $-\text{CH}_2\text{CH}_2-$, or $-\text{C}(\text{CH}_3)_2-$,

or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof.

7. A compound of formula IC according to claim 6, wherein the compounds are

3-[2-Chloro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-pteridine-2,4-dione

3-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,7-dimethyl-thieno[3,2-d]pyrimidine-2,4-dione

2-(2,6-Difluoro-4-(pyridin-3-ylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione

3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropyl-quinazoline-2,4-dione

3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione

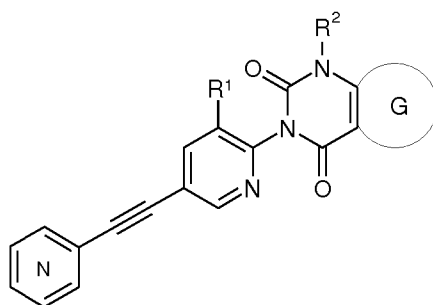
6-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

6-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione

3-[2-Chloro-6-fluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1-(oxetan-3-yl)quinazoline-2,4-dione or

3-[2,6-Difluoro-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethyl-quinazoline-2,4-dione.

8. A compound of formula ID according to claim 1,



ID

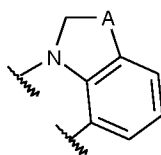
wherein

G is a 5 or 6-membered aromatic or heteroaromatic ring containing 0, 1, 2 or 3 heteroatoms, selected from the group consisting of phenyl, pyridinyl with different N-positions, imidazolyl, pyrazinyl, pyrimidinyl, thiophenyl, thiazolyl, pyrazolyl or thiadiazolyl, which are optionally substituted by 1, 2 or 3 substituents, selected from the group consisting of halogen, C_{1-7} -alkyl, C_{1-7} -alkoxy, C_{1-7} -alkoxy substituted by halogen or NRR' ;

R and R' are independently from each other hydrogen or C_{1-7} -alkyl, or may form together with the N atom to which they are attached a five or six membered saturated heterocyclic group which may contain an additional oxygen, NH, or N-C_{1-7} -alkyl group;

R¹ is hydrogen, halogen or C_{1-7} -alkyl substituted by halogen;

R² is hydrogen, C_{1-7} -alkyl, C_{1-7} -alkoxyalkyl, cycloalkyl or heterocycloalkyl; or R² may form together with the closest carbon atom in group G a group

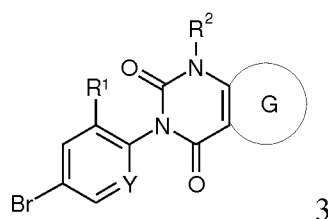


for A being $-\text{CH}_2-$, $-\text{CH}_2\text{CH}_2-$, or $-\text{C}(\text{CH}_3)_2-$,

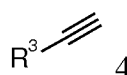
or a pharmaceutically acceptable salt or acid addition salt, a racemic mixture, or its corresponding enantiomer and/or optical isomer and/or stereoisomer thereof.

9. A compound of formula ID according to claim 8, wherein the compound is 6-[3-Chloro-5-[2-(3-pyridyl)ethynyl]-2-pyridyl]-4-isopropyl-2-methyl-pyrazolo[4,3-d]pyrimidine-5,7-dione.

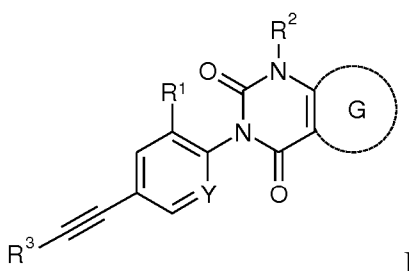
10. A process for the manufacture of a compound of formula I as defined in any one of claims 1 to 9, which process comprises
reacting a compound of formula 3



with a compound of formula



to a compound of formula I



wherein the substituents are described in claim 1, or
if desired, converting the compounds obtained into pharmaceutically acceptable acid addition salts.

11. A compound of formula I according to any one of claims 1 - 9 for use as therapeutically active substances.

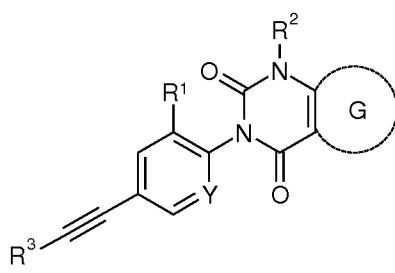
12. A compound of formula I according to any one of claims 1- 9 for use in the treatment of Parkinson's disease, anxiety, emesis, obsessive compulsive disorder, autism, neuroprotection, cancer, depression and diabetes type 2.

13. A pharmaceutical composition comprising a compound of formula I as claimed in any one of claim 1 - 9 and pharmaceutically acceptable excipients.

14. The use of a compound of formula I according to any one of claims 1 - 9 for the preparation of a medicament for the treatment of Parkinson's disease, anxiety, emesis, obsessive compulsive disorder, autism, neuroprotection, cancer, depression and diabetes type 2,

Patentansprüche

1. Verbindungen der Formel I,



wobei

Y N oder C-R¹ ist;

G ein 5- oder 6-gliedriger aromatischer oder heteroaromatischer Ring ist, enthaltend 0, 1, 2 oder 3 Heteroatome, die aus der Gruppe ausgewählt sind, bestehend aus Phenyl, Pyridinyl mit unterschiedlichen N-Positionen, Imidazolyl, Pyrazinyl, Pyrimidinyl, Thiophenyl, Thiazolyl, Pyrazolyl oder Thiadiazolyl, die gegebenenfalls durch 1, 2 oder 3 Substituenten substituiert sind, die aus der Gruppe ausgewählt sind, bestehend aus Halogen, C₁-C₇-Alkyl, C₁-C₇-Alkoxy, C₁-C₇-Alkoxy substituiert durch Halogen oder NRR';

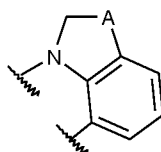
R und R' unabhängig voneinander Wasserstoff oder C₁-C₇-Alkyl sind, oder zusammen mit dem N-Atom, an das sie gebunden sind, eine fünf- oder sechsgliedrige gesättigte heterocyclische Gruppe bilden können, die eine/n zusätzliche/n Sauerstoff, NH, oder N-C₁-C₇-Alkylgruppe enthalten kann;

R¹ Wasserstoff, Halogen oder C₁-C₇-Alkyl substituiert durch Halogen ist;

R¹' Wasserstoff, Halogen oder C₁-C₇-Alkyl substituiert durch Halogen ist;

R² Wasserstoff, C₁-C₇-Alkyl, C₁-C₇-Alkoxyalkyl, Cycloalkyl oder Heterocycloalkyl ist;

oder R² zusammen mit dem nächstgelegenen Kohlenstoffatom in Gruppe G eine Gruppe bilden kann,

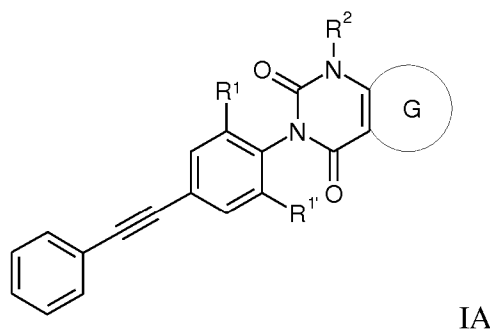


wobei A -CH₂-, -CH₂CH₂-, oder -C(CH₃)₂- ist,

R³ Phenyl oder Pyridinyl ist, wobei sich das N-Atom in der Pyridinylgruppe in verschiedenen Positionen befinden kann;

oder ein pharmazeutisch annehmbares Salz oder Säureadditionssalz, ein racemisches Gemisch, oder sein entsprechendes Enantiomer und/oder optisches Isomer und/oder Stereoisomer davon.

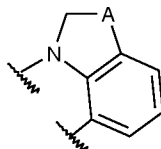
2. Verbindung der Formel IA nach Anspruch 1,



wobei

G ein 5- oder 6-gliedriger aromatischer oder heteroaromatischer Ring ist, enthaltend 0, 1, 2 oder 3 Heteroatome, die aus der Gruppe ausgewählt sind, bestehend aus Phenyl, Pyridinyl mit unterschiedlichen N-Positionen, Imidazolyl, Pyrazinyl, Pyrimidinyl, Thiophenyl, Thiazolyl, Pyrazolyl oder Thiadiazolyl, die gegebenenfalls durch

1, 2 oder 3 Substituenten substituiert sind, die aus der Gruppe ausgewählt sind, bestehend aus Halogen, C₁-C₇-Alkyl, C₁-C₇-Alkoxy, C₁-C₇-Alkoxy substituiert durch Halogen oder NRR';
 R und R' unabhängig voneinander Wasserstoff oder C₁-C₇-Alkyl sind, oder zusammen mit dem N-Atom, an das sie gebunden sind, eine fünf- oder sechsgliedrige gesättigte heterocyclische Gruppe bilden können, die
 5 eine/n zusätzliche/n Sauerstoff, NH, oder N-C₁-C₇-Alkylgruppe enthalten kann;
 R¹ Wasserstoff, Halogen oder C₁-C₇-Alkyl substituiert durch Halogen ist;
 R¹ Wasserstoff, Halogen oder C₁-C₇-Alkyl substituiert durch Halogen ist;
 R² Wasserstoff, C₁-C₇-Alkyl, C₁-C₇-Alkoxyalkyl, Cycloalkyl oder Heterocycloalkyl ist; oder R² zusammen mit dem nächstgelegenen Kohlenstoffatom in Gruppe G eine Gruppe bilden kann,



wobei A -CH₂-, -CH₂CH₂-, oder -C(CH₃)₂- ist,

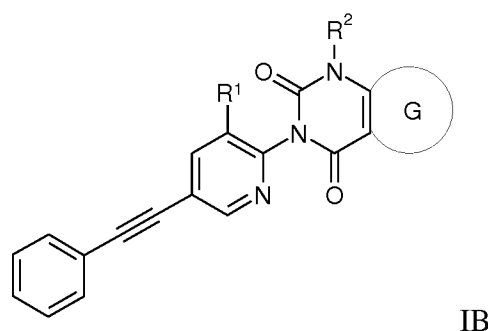
oder ein pharmazeutisch annehmbares Salz oder Säureadditionssalz, ein racemisches Gemisch oder ihr entsprechendes Enantiomer und/oder optisches Isomer und/oder Stereoisomer davon.

3. Verbindung der Formel IA nach Anspruch 2, wobei die Verbindungen

3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-methylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1,8-dimethylchinazolin-2,4-dion
 8-Chlor-3-[2-chlor-4-(2-phenylethynyl)phenyl]-1-methylchinazolin-2,4-dion
 7-Chlor-3-[2-chlor-4-(2-phenylethynyl)phenyl]-1-methylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1,7-dimethylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1H-pyrido[2,3-d]pyrimidin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-methylpyrido[2,3-d]pyrimidin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-8-ethyl-1-methylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-methylpyrido[3,2-d]pyrimidin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-methylpyrido[4,3-d]pyrimidin-2,4-dion
 1-[2-Chlor-4-(2-phenylethynyl)phenyl]-3,7-dimethylpurin-2,6-dion
 2-(2-Chlor-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]chinazolin-1,3(2H)-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-isopropylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-ethylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-methylpteridin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1H-pyrimido[4,5-d]pyrimidin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-methylpyrimido[4,5-d]pyrimidin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-6-fluor-1-methylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-7,8-difluor-1-methylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1,7-dimethylthieno[3,2-d]pyrimidin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-isopropylpteridin-2,4-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1-methylchinazolin-2,4-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1,8-dimethylchinazolin-2,4-dion
 3-(2,6-Difluor-4-phenylethynylphenyl)-1-methyl-1H-pyrido[2,3-d]pyrimidin-2,4-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1,7-dimethylchinazolin-2,4-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1,7-dimethylthieno[3,2-d]pyrimidin-2,4-dion
 2-(2,6-Difluor-4-(phenylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]chinazolin-1,3(2H)-dion
 6-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-4H-thiazolo[4,5-d]pyrimidin-5,7-dion
 6-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-4-methylthiazolo[4,5-d]pyrimidin-5,7-dion
 6-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-2,4-dimethylpyrazolo[4,3-d]pyrimidin-5,7-dion
 6-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1,4-dimethylpyrazolo[4,3-d]pyrimidin-5,7-dion
 6-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-4-isopropyl-2-methylpyrazolo[4,3-d]pyrimidin-5,7-dion
 6-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-4-isopropyl-1-methylpyrazolo[4,3-d]pyrimidin-5,7-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1-isopropylpyrido[2,3-d]pyrimidin-2,4-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1,8-dimethylpyrido[3,2-d]pyrimidin-2,4-dion

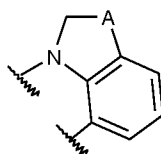
6-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-4-methyl-[1,2,5]thiadiazolo[3,4-d]pyrimidin-5,7-dion
 6-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-4-isopropyl-[1,2,5]thiadiazolo[3,4-d]pyrimidin-5,7-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1,5-dimethyl-chinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-(2-methoxyethyl)chinazolin-2,4-dion
 5 6-[2-Chlor-4-(2-phenylethynyl)phenyl]-4-methyl-2-morpholinothiazolo[4,5-d]pyrimidin-5,7-dion
 6-(2-Chlor-4-phenylethynylphenyl)-4-methyl-4H-thiazolo[4,5-d]pyrimidin-5,7-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-1-cyclobutylchinazolin-2,4-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-8-isopropyl-1-methylchinazolin-2,4-dion
 8-Chlor-3-[2,6-difluor-4-(2-phenylethynyl)phenyl]-1-methylpyrido[3,2-d]pyrimidin-2,4-dion
 10 5-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-2,7-dimethylpyrazolo[3,4-d]pyrimidin-4,6-dion
 5-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1,7-dimethylpyrazolo[3,4-d]pyrimidin-4,6-dion
 5-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-7-isopropyl-2H-pyrazolo[3,4-d]pyrimidin-4,6-dion
 3-[2-Chlor-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methylchinazolin-2,4-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1-(oxetan-3-yl)chinazolin-2,4-dion
 15 6-[2-Chlor-4-(2-phenylethynyl)phenyl]-4-methylthiazolo [5,4-d]pyrimidin-5,7-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-1-isopropylpteridin-2,4-dion
 1-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-3,7-dimethylpurin-2,6-dion
 3-[2,6-Difluor-4-(2-phenylethynyl)phenyl]-8-methoxy-1-methylchinazolin-2,4-dion
 3-[2-Fluor-4-(2-phenylethynyl)-6-(trifluormethyl)phenyl]-1-isopropylchinazolin-2,4-dion oder
 20 8-(2,2-Difluorethoxy)-3-[2,6-difluor-4-(2-phenylethynyl)phenyl]-1-methylchinazolin-2,4-dion sind.

4. Verbindung der Formel IB nach Anspruch 1



wobei

G ein 5- oder 6-gliedriger aromatischer oder heteroaromatischer Ring ist, enthaltend 0, 1, 2 oder 3 Heteroatome, die aus der Gruppe ausgewählt sind, bestehend aus Phenyl, Pyridinyl mit unterschiedlichen N-Positionen, Imidazolyl, Pyrazinyl, Pyrimidinyl, Thiophenyl, Thiazolyl, Pyrazolyl oder Thiadiazolyl, die gegebenenfalls durch 1, 2 oder 3 Substituenten substituiert sind, die aus der Gruppe ausgewählt sind, bestehend aus Halogen, C₁-C₇-Alkyl, C₁-C₇-Alkoxy, C₁-C₇-Alkoxy substituiert durch Halogen oder NRR';
 R und R' unabhängig voneinander Wasserstoff oder C₁-C₇-Alkyl sind, oder zusammen mit dem N-Atom, an das sie gebunden sind, eine fünf- oder sechsgliedrige gesättigte heterocyclische Gruppe bilden können, die eine/n zusätzliche/n Sauerstoff, NH, oder N-C₁-C₇-Alkylgruppe enthalten kann;
 R¹ Wasserstoff, Halogen oder C₁-C₇-Alkyl substituiert durch Halogen ist;
 R² Wasserstoff, C₁-C₇-Alkyl, C₁-C₇-Alkoxyalkyl, Cycloalkyl oder Heterocycloalkyl ist;
 oder R² zusammen mit dem nächstgelegenen Kohlenstoffatom in Gruppe G eine Gruppe bilden kann,



wobei A -CH₂-, -CH₂CH₂-, oder -C(CH₃)₂- ist,

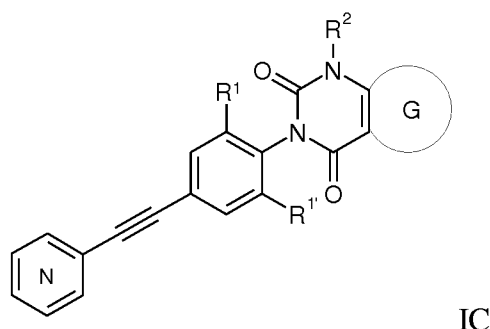
oder ein pharmazeutisch annehmbares Salz oder Säureadditionssalz, ein racemisches Gemisch oder sein entspre-

chendes Enantiomer und/oder optisches Isomer und/oder Stereoisomer davon.

5. Verbindung der Formel IB nach Anspruch 4, wobei die Verbindungen

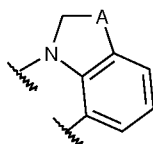
- 3-[3-Chlor-5-(2-phenylethynyl)-2-pyridyl]-1-methylchinazolin-2,4-dion
 3-[3-Chlor-5-(2-phenylethynyl)-2-pyridyl]-1-isopropylchinazolin-2,4-dion
 3-[3-Chlor-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethylchinazolin-2,4-dion
 2-(3-Chlor-5-(phenylethynyl)pyridin-2-yl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]chinazolin-1,3(2H)-dion
 1-Methyl-3-[5-(2-phenylethynyl)-3-(trifluormethyl)-2-pyridyl]chinazolin-2,4-dion
 3-[3-Fluor-5-(2-phenylethynyl)-2-pyridyl]-1,8-dimethylchinazolin-2,4-dion
 1-Isopropyl-3-[5-(2-phenylethynyl)-2-pyridyl]chinazolin-2,4-dion
 6-[3-Fluor-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methylpyrazolo[4,3-d]pyrimidin-5,7-dion oder
 6-[3-Chlor-5-(2-phenylethynyl)-2-pyridyl]-4-isopropyl-2-methylpyrazolo[4,3-d]pyrimidin-5,7-dion sind.

6. Verbindung der Formel IC nach Anspruch 1,



wobei

G ein 5- oder 6-gliedriger aromatischer oder heteroaromatischer Ring ist, enthaltend 0, 1, 2 oder 3 Heteroatome, die aus der Gruppe ausgewählt sind, bestehend aus Phenyl, Pyridinyl mit unterschiedlichen N-Positionen, Imidazolyl, Pyrazinyl, Pyrimidinyl, Thiophenyl, Thiazolyl, Pyrazolyl oder Thiadiazolyl, die gegebenenfalls durch 1, 2 oder 3 Substituenten substituiert sind, die aus der Gruppe ausgewählt sind, bestehend aus Halogen, C₁-C₇-Alkyl, C₁-C₇-Alkoxy, C₁-C₇-Alkoxy substituiert durch Halogen oder NRR';
 R und R' unabhängig voneinander Wasserstoff oder C₁-C₇-Alkyl sind, oder zusammen mit dem N-Atom, an das sie gebunden sind, eine fünf- oder sechsgliedrige gesättigte heterocyclische Gruppe bilden können, die eine/n zusätzliche/n Sauerstoff, NH, oder N-C₁-C₇-Alkylgruppe enthalten kann;
 R¹ Wasserstoff, Halogen oder C₁-C₇-Alkyl substituiert durch Halogen ist;
 R^{1'} Wasserstoff, Halogen oder C₁-C₇-Alkyl substituiert durch Halogen ist;
 R² Wasserstoff, C₁-C₇-Alkyl, C₁-C₇-Alkoxyalkyl, Cycloalkyl oder Heterocycloalkyl ist;
 oder R² zusammen mit dem nächstgelegenen Kohlenstoffatom in Gruppe G eine Gruppe bilden kann,



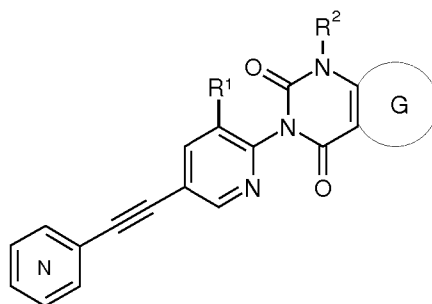
wobei A -CH₂-, -CH₂CH₂-, oder -C(CH₃)₂- ist,

oder ein pharmazeutisch annehmbares Salz oder Säureadditionssalz, ein racemisches Gemisch oder sein entsprechendes Enantiomer und/oder optisches Isomer und/oder Stereoisomer davon.

7. Verbindung der Formel IC nach Anspruch 6, wobei die Verbindungen

3-[2-Chlor-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropylpteridin-2,4-dion
 3-[2,6-Difluor-4-[2-(3-pyridyl)ethynyl]phenyl]-1,7-dimethylthieno[3,2-d]pyrimidin-2,4-dion
 2-(2,6-Difluor-4-(pyridin-3-ylethynyl)phenyl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]chinazofin-1,3(2H)-dion
 3-[2-Chlor-6-fluor-4-[2-(3-pyridyl)ethynyl]phenyl]-1-isopropylchinazolin-2,4-dion
 3-[2-Chlor-6-fluor-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethylchinazolin-2,4-dion
 6-[2,6-Difluor-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methylpyrazolo[4,3-d]pyrimidin-5,7-dion
 6-[2-Chlor-6-fluor-4-[2-(3-pyridyl)ethynyl]phenyl]-4-isopropyl-2-methylpyrazolo[4,3-d]pyrimidin-5,7-dion
 3-[2-Chlor-6-fluor-4-[2-(3-pyridyl)ethynyl]phenyl]-1-(oxetan-3-yl)chinazolin-2,4-dion oder
 3-[2,6-Difluor-4-[2-(3-pyridyl)ethynyl]phenyl]-1,8-dimethylchinazolin-2,4-dion sind.

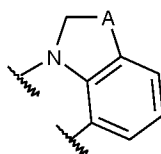
8. Verbindung der Formel ID nach Anspruch 1,



ID

wobei

G ein 5- oder 6-gliedriger aromatischer oder heteroaromatischer Ring ist, enthaltend 0, 1, 2 oder 3 Heteroatome, die aus der Gruppe ausgewählt sind, bestehend aus Phenyl, Pyridinyl mit unterschiedlichen N-Positionen, Imidazolyl, Pyrazinyl, Pyrimidinyl, Thiophenyl, Thiazolyl, Pyrazolyl oder Thiadiazolyl, die gegebenenfalls durch 1, 2 oder 3 Substituenten substituiert sind, die aus der Gruppe ausgewählt sind, bestehend aus Halogen, C₁-C₇-Alkyl, C₁-C₇-Alkoxy, C₁-C₇-Alkoxy substituiert durch Halogen oder NRR';
 R und R' unabhängig voneinander Wasserstoff oder C₁-C₇-Alkyl sind, oder zusammen mit dem N-Atom, an das sie gebunden sind, eine fünf- oder sechsgliedrige gesättigte heterocyclische Gruppe bilden können, die eine/n zusätzliche/n Sauerstoff, NH, oder N-C₁-C₇-Alkylgruppe enthalten kann;
 R¹ Wasserstoff, Halogen oder C₁-C₇-Alkyl substituiert durch Halogen ist;
 R² Wasserstoff, C₁-C₇-Alkyl, C₁-C₇-Alkoxyalkyl, Cycloalkyl oder Heterocycloalkyl ist;
 oder R² zusammen mit dem nächstgelegenen Kohlenstoffatom in Gruppe G eine Gruppe bilden kann,

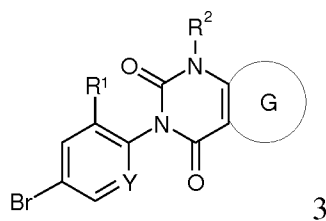


wobei A -CH₂-, -CH₂CH₂, oder -C(CH₃)₂- ist,

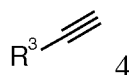
oder ein pharmazeutisch annehmbares Salz oder Säureadditionssalz, ein racemisches Gemisch oder sein entsprechendes Enantiomer und/oder optisches Isomer und/oder Stereoisomer davon.

9. Verbindung der Formel ID nach Anspruch 8, wobei die Verbindung 6-[3-Chlor-5-[2-(3-pyridyl)ethynyl]-2-pyridyl]-4-isopropyl-2-methylpyrazolo[4,3-d]pyrimidin-5,7-dion ist.

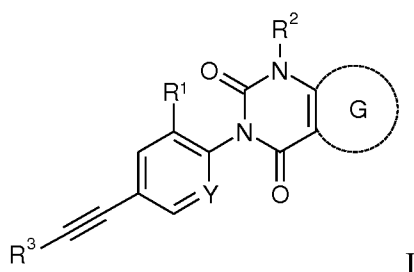
10. Verfahren zur Herstellung einer Verbindung der Formel I, wie in einem der Ansprüche 1 bis 9 definiert, wobei das Verfahren umfasst, dass eine Verbindung der Formel 3



mit einer Verbindung der Formel



zu einer Verbindung der Formel I

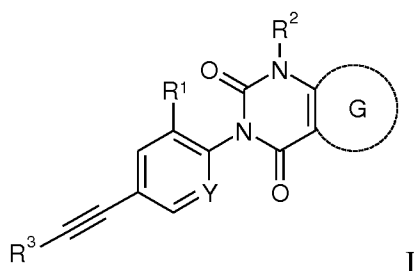


umgesetzt wird, wobei die Substituenten in Anspruch 1 beschrieben sind, oder, falls erwünscht, die erhaltenen Verbindungen in pharmazeutisch annehmbare Säureadditionssalze umgewandelt werden.

11. Verbindung der Formel I nach einem der Ansprüche 1-9 zur Verwendung als therapeutisch wirksame Substanzen.
12. Verbindung der Formel I nach einem der Ansprüche 1-9 zur Verwendung bei der Behandlung von Morbus Parkinson, Angstzuständen, Erbrechen, Zwangsstörungen, Autismus, Neuroprotektion, Krebs, Depression und Diabetes Typ 2.
13. Pharmazeutische Zusammensetzung, umfassend eine Verbindung der Formel I, wie in einem der Ansprüche 1-9 beansprucht, und pharmazeutisch annehmbare Exzipienten.
14. Verwendung einer Verbindung der Formel I nach einem der Ansprüche 1-9 zur Herstellung eines Arzneimittels für die Behandlung von Morbus Parkinson, Angstzuständen, Erbrechen, Zwangsstörungen, Autismus, Neuroprotektion, Krebs, Depression und Diabetes Typ 2.

Revendications

1. Composés de formule I



dans laquelle

Y représente N ou C-R^{1'} ;

G représente un cycle aromatique ou hétéroaromatique à 5 ou 6 chaînons contenant 0, 1, 2 ou 3 hétéroatomes, choisi dans le groupe comprenant un groupe phényle, pyridinyle avec différentes positions de N, imidazolyle, pyrazinyle, pyrimidinyle, thiophényle, thiazolyle, pyrazolyle ou thiadiazolyle, qui est éventuellement substitué par 1, 2 ou 3 substituants, choisis dans le groupe comprenant un atome d'halogène, un groupe alkyle en C₁ à C₇, alcoxy en C₁ à C₇, alcoxy en C₁ à C₇ substitué par un atome d'halogène ou NRR' ;

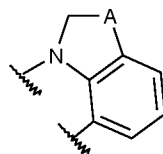
R et R' représentent indépendamment l'un de l'autre un atome d'hydrogène ou un groupe alkyle en C₁ à C₇, ou peuvent former, ensemble avec l'atome d'azote auquel ils sont liés, un groupe hétérocyclique saturé à cinq ou six chaînons qui peut contenir un oxygène supplémentaire, NH, ou un groupe N-alkyle en C₁ à C₇ ;

R¹ représente un atome d'hydrogène, un atome d'halogène ou un groupe alkyle en C₁ à C₇ substitué par un atome d'halogène ;

R^{1'} représente un atome d'hydrogène, un atome d'halogène ou un groupe alkyle en C₁ à C₇ substitué par un atome d'halogène ;

R² représente un atome d'hydrogène, un groupe alkyle en C₁ à C₇, alcoxyalkyle en C₁ à C₇, cycloalkyle ou hétérocycloalkyle ;

ou R² peut former, ensemble avec l'atome de carbone le plus proche dans le groupe G, un groupe

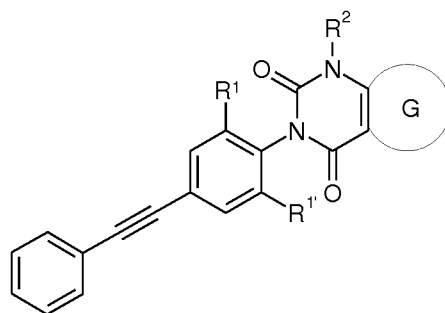


A représentant -CH₂-, -CH₂CH₂-, ou -C(CH₃)₂-;

R³ représente un groupe phényle ou pyridinyle, dans lequel l'atome N peut se trouver à différentes positions dans le groupe pyridinyle ;

ou un sel ou un sel d'addition d'acide pharmaceutiquement acceptable, un mélange racémique, ou ses énantiomères et/ou isomères optiques et/ou stéréoisomères correspondants.

2. Composé de formule IA selon la revendication 1,



IA

dans laquelle

G représente un cycle aromatique ou hétéroaromatique à 5 ou 6 chaînons contenant 0, 1, 2 ou 3 hétéroatomes, choisi dans le groupe comprenant un groupe phényle, pyridinyle avec différentes positions de N, imidazolyle, pyrazinyle, pyrimidinyle, thiophényle, thiazolyle, pyrazolyle ou thiadiazolyle, qui est éventuellement substitué par 1, 2 ou 3 substituants, choisis dans le groupe comprenant un atome d'halogène, un groupe alkyle en C₁ à C₇, alcoxy en C₁ à C₇, alcoxy en C₁ à C₇ substitué par un atome d'halogène ou NRR' ;

R et R' représentent indépendamment l'un de l'autre un atome d'hydrogène ou un groupe alkyle en C₁ à C₇, ou peuvent former, ensemble avec l'atome d'azote auquel ils sont liés, un groupe hétérocyclique saturé à cinq ou six chaînons qui peut contenir un oxygène supplémentaire, NH, ou un groupe N-alkyle en C₁ à C₇ ;

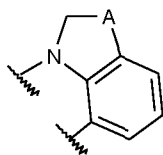
R¹ représente un atome d'hydrogène, un atome d'halogène ou un groupe alkyle en C₁ à C₇ substitué par un atome d'halogène ;

R^{1'} représente un atome d'hydrogène, un atome d'halogène ou un groupe alkyle en C₁ à C₇ substitué par un

atome d'halogène ;

R² représente un atome d'hydrogène, un groupe alkyle en C₁ à C₇, alcoxyalkyle en C₁ à C₇, cycloalkyle ou hétérocycloalkyle ;

ou R² peut former, ensemble avec l'atome de carbone le plus proche dans le groupe G, un groupe



A représentant -CH₂-, -CH₂CH₂-, ou -C(CH₃)₂-,

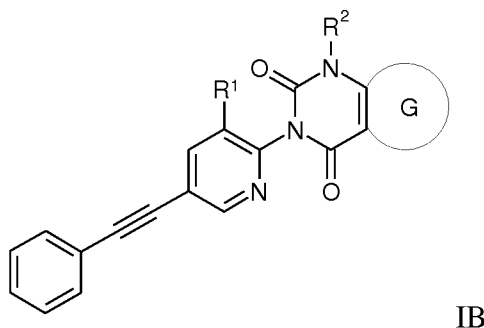
ou un sel ou un sel d'addition d'acide pharmaceutiquement acceptable, un mélange racémique, ou ses énantiomères et/ou isomères optiques et/ou stéréoisomères correspondants.

3. Composé de formule IA selon la revendication 2, ledit composé étant :

la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-méthyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1,8-diméthyl-quinazoline-2,4-dione
 la 8-chloro-3-[2-chloro-4-(2-phényléthynyl)phényl]-1-méthyl-quinazoline-2,4-dione
 la 7-chloro-3-[2-chloro-4-(2-phényléthynyl)phényl]-1-méthyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1,7-diméthyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1H-pyrido[2,3-d]pyrimidine-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-méthyl-pyrido[2,3-d]pyrimidine-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-8-éthyl-1-méthyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-méthyl-pyrido[3,2-d]pyrimidine-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-méthyl-pyrido[4,3-d]pyrimidine-2,4-dione
 la 1-[2-chloro-4-(2-phényléthynyl)phényl]-3,7-diméthyl-purine-2,6-dione
 la 2-(2-chloro-4-(2-phényléthynyl)phényl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-isopropyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-éthyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-méthyl-ptéridine-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1H-pyrimido[4,5-d]pyrimidine-2,4-dione la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-méthyl-pyrimido[4,5-d]pyrimidine-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-6-fluoro-1-méthyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-7,8-difluoro-1-méthyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1,7-diméthyl-thiéno[3,2-d]pyrimidine-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-isopropyl-ptéridine-2,4-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1-méthyl-quinazoline-2,4-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1,8-diméthyl-quinazoline-2,4-dione
 la 3-(2,6-difluoro-4-phényléthynyl-phényl)-1-méthyl-1H-pyrido[2,3-d]pyrimidine-2,4-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1,7-diméthyl-quinazoline-2,4-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1,7-diméthyl-thiéno[3,2-d]pyrimidine-2,4-dione
 la 2-(2,6-difluoro-4-(2-phényléthynyl)phényl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 la 6-[2,6-difluoro-4-(2-phényléthynyl)phényl]-4H-thiazolo[4,5-d]pyrimidine-5,7-dione
 la 6-[2,6-difluoro-4-(2-phényléthynyl)phényl]-4-méthyl-thiazolo[4,5-d]pyrimidine-5,7-dione
 la 6-[2,6-difluoro-4-(2-phényléthynyl)phényl]-2,4-diméthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 la 6-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1,4-diméthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 la 6-[2,6-difluoro-4-(2-phényléthynyl)phényl]-4-isopropyl-2-méthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 la 6-[2,6-difluoro-4-(2-phényléthynyl)phényl]-4-isopropyl-1-méthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1-isopropyl-pyrido[2,3-d]pyrimidine-2,4-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1,8-diméthyl-pyrido[3,2-d]pyrimidine-2,4-dione
 la 6-[2,6-difluoro-4-(2-phényléthynyl)phényl]-4-méthyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione
 la 6-[2,6-difluoro-4-(2-phényléthynyl)phényl]-4-isopropyl-[1,2,5]thiadiazolo[3,4-d]pyrimidine-5,7-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1,5-diméthyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-(2-méthoxyéthyl)quinazoline-2,4-dione
 la 6-[2-chloro-4-(2-phényléthynyl)phényl]-4-méthyl-2-morpholino-thiazolo[4,5-d]pyrimidine-5,7-dione

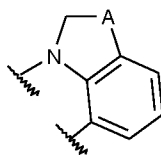
la 6-(2-chloro-4-phényléthynyl-phényl)-4-méthyl-4H-thiazolo[4,5-d]pyrimidine-5,7-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-1-cyclobutyl-quinazoline-2,4-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-8-isopropyl-1-méthyl-quinazoline-2,4-dione
 la 8-chloro-3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1-méthyl-pyrido[3,2-d]pyrimidine-2,4-dione
 la 5-[2,6-difluoro-4-(2-phényléthynyl)phényl]-2,7-diméthyl-pyrazolo[3,4-d]pyrimidine-4,6-dione
 la 5-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1,7-diméthyl-pyrazolo[3,4-d]pyrimidine-4,6-dione
 la 5-[2,6-difluoro-4-(2-phényléthynyl)phényl]-7-isopropyl-2H-pyrazolo[3,4-d]pyrimidine-4,6-dione
 la 3-[2-chloro-4-(2-phényléthynyl)phényl]-8-méthoxy-1-méthyl-quinazoline-2,4-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1-(oxétan-3-yl)quinazoline-2,4-dione
 la 6-[2-chloro-4-(2-phényléthynyl)phényl]-4-méthyl-thiazolo[5,4-d]pyrimidine-5,7-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1-isopropyl-ptéridine-2,4-dione
 la 1-[2,6-difluoro-4-(2-phényléthynyl)phényl]-3,7-diméthyl-purine-2,6-dione
 la 3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-8-méthoxy-1-méthyl-quinazoline-2,4-dione
 la 3-[2-fluoro-4-(2-phényléthynyl)-6-(trifluorométhyl)phényl]-1-isopropyl-quinazoline-2,4-dione ou
 la 8-(2,2-difluoroéthoxy)-3-[2,6-difluoro-4-(2-phényléthynyl)phényl]-1-méthyl-quinazoline-2,4-dione.

4. Composé de formule IB selon la revendication 1,



dans laquelle

G représente un cycle aromatique ou hétéroaromatique à 5 ou 6 chaînons contenant 0, 1, 2 ou 3 hétéroatomes, choisi dans le groupe comprenant un groupe phényle, pyridinyle avec différentes positions de N, imidazole, pyrazinyle, pyrimidinyle, thiophényle, thiazole, pyrazole ou thiadiazole, qui est éventuellement substitué par 1, 2 ou 3 substituants, choisis dans le groupe comprenant un atome d'halogène, un groupe alkyle en C₁ à C₇, alcoxy en C₁ à C₇, alcoxy en C₁ à C₇ substitué par un atome d'halogène ou NRR' ;
 R et R' représentent indépendamment l'un de l'autre un atome d'hydrogène ou un groupe alkyle en C₁ à C₇, ou peuvent former, ensemble avec l'atome d'azote auquel ils sont liés, un groupe hétérocyclique saturé à cinq ou six chaînons qui peut contenir un oxygène supplémentaire, NH, ou un groupe N-alkyle en C₁ à C₇ ;
 R¹ représente un atome d'hydrogène, un atome d'halogène ou un groupe alkyle en C₁ à C₇ substitué par un atome d'halogène ;
 R² représente un atome d'hydrogène, un groupe alkyle en C₁ à C₇, alcoxyalkyle en C₁ à C₇, cycloalkyle ou hétérocycloalkyle ;
 ou R² peut former, ensemble avec l'atome de carbone le plus proche dans le groupe G, un groupe



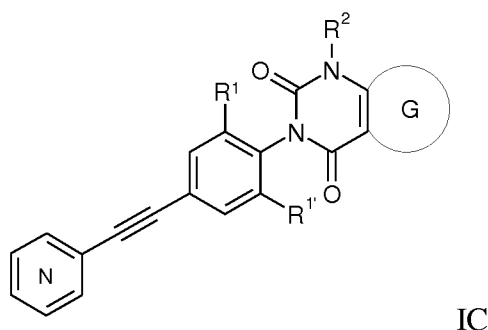
A représentant -CH₂-, -CH₂CH₂-, ou -C(CH₃)₂-,

ou un sel ou un sel d'addition d'acide pharmaceutiquement acceptable, un mélange racémique, ou ses énantiomères et/ou isomères optiques et/ou stéréoisomères correspondants.

5. Composé de formule IB selon la revendication 4, ledit composé étant :

la 3-[3-chloro-5-(2-phényléthynyl)-2-pyridyl]-1-méthyl-quinazoline-2,4-dione
 la 3-[3-chloro-5-(2-phényléthynyl)-2-pyridyl]-1-isopropyl-quinazoline-2,4-dione
 la 3-[3-chloro-5-(2-phényléthynyl)-2-pyridyl]-1,8-diméthyl-quinazoline-2,4-dione
 la 2-(3-chloro-5-(phényléthynyl)pyridin-2-yl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 la 1-méthyl-3-[5-(2-phényléthynyl)-3-(trifluorométhyl)-2-pyridyl]quinazoline-2,4-dione
 la 3-[3-fluoro-5-(2-phényléthynyl)-2-pyridyl]-1,8-diméthyl-quinazoline-2,4-dione
 la 1-isopropyl-3-[5-(2-phényléthynyl)-2-pyridyl]quinazoline-2,4-dione
 la 6-[3-fluoro-5-(2-phényléthynyl)-2-pyridyl]-4-isopropyl-2-méthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione ou
 la 6-[3-chloro-5-(2-phényléthynyl)-2-pyridyl]-4-isopropyl-2-méthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione.

6. Composé de formule IC selon la revendication 1,



dans laquelle

G représente un cycle aromatique ou hétéroaromatique à 5 ou 6 chaînons contenant 0, 1, 2 ou 3 hétéroatomes, choisi dans le groupe comprenant un groupe phényle, pyridinyle avec différentes positions de N, imidazolyle, pyrazinyle, pyrimidinyle, thiophényle, thiazolyle, pyrazolyle ou thiadiazolyle, qui est éventuellement substitué par 1, 2 ou 3 substituants, choisis dans le groupe comprenant un atome d'halogène, un groupe alkyle en C₁ à C₇, alcoxy en C₁ à C₇, alcoxy en C₁ à C₇ substitué par un atome d'halogène ou NRR' ;

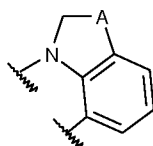
R et R' représentent indépendamment l'un de l'autre un atome d'hydrogène ou un groupe alkyle en C₁ à C₇, ou peuvent former, ensemble avec l'atome d'azote auquel ils sont liés, un groupe hétérocyclique saturé à cinq ou six chaînons qui peut contenir un oxygène supplémentaire, NH, ou un groupe N-alkyle en C₁ à C₇ ;

R¹ représente un atome d'hydrogène, un atome d'halogène ou un groupe alkyle en C₁ à C₇ substitué par un atome d'halogène ;

R^{1'} représente un atome d'hydrogène, un atome d'halogène ou un groupe alkyle en C₁ à C₇ substitué par un atome d'halogène ;

R² représente un atome d'hydrogène, Un groupe alkyle en C₁ à C₇, alcoxyalkyle en C₁ à C₇, cycloalkyle ou hétérocycloalkyle ;

ou R² peut former, ensemble avec l'atome de carbone le plus proche dans le groupe G, un groupe



A représentant -CH₂-, -CH₂CH₂-, ou -C(CH₃)₂-,

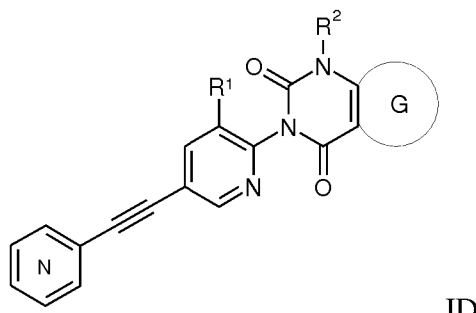
ou un sel ou un sel d'addition d'acide pharmaceutiquement acceptable, un mélange racémique, ou ses énantiomères et/ou isomères optiques et/ou stéréoisomères correspondants.

7. Composé de formule IC selon la revendication 6, le composé étant :

la 3-[2-chloro-4-[2-(3-pyridyl)éthynyl]phényl]-1-isopropyl-ptéridine-2,4-dione

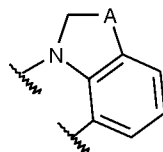
la 3-[2,6-difluoro-4-[2-(3-pyridyl)éthynyl]phényl]-1,7-diméthyl-thiéno[3,2-d]pyrimidine-2,4-dione
 la 2-(2,6-difluoro-4-(pyridin-3-yléthynyl)phényl)-5,6-dihydro-1H-pyrrolo[3,2,1-ij]quinazoline-1,3(2H)-dione
 la 3-[2-chloro-6-fluoro-4-[2-(3-pyridyl)éthynyl]phényl]-1-isopropyl-quinazoline-2,4-dione
 la 3-[2-chloro-6-fluoro-4-[2-(3-pyridyl)éthynyl]phényl]-1,8-diméthyl-quinazoline-2,4-dione
 la 6-[2,6-difluoro-4-[2-(3-pyridyl)éthynyl]phényl]-4-isopropyl-2-méthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 la 6-[2-chloro-6-fluoro-4-[2-(3-pyridyl)éthynyl]phényl]-4-isopropyl-2-méthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione
 la 3-[2-chloro-6-fluoro-4-[2-(3-pyridyl)éthynyl]phényl]-1-(oxétan-3-yl)quinazoline-2,4-dione ou
 la 3-[2,6-difluoro-4-[2-(3-pyridyl)éthynyl]phényl]-1,8-diméthyl-quinazoline-2,4-dione.

8. Composé de formule ID selon la revendication 1,



dans laquelle

G représente un cycle aromatique ou hétéroaromatique à 5 ou 6 chaînons contenant 0, 1, 2 ou 3 hétéroatomes, choisi dans le groupe comprenant un groupe phényle, pyridinyle avec différentes positions de N, imidazolyle, pyrazinyle, pyrimidinyle, thiophényle, thiazolyle, pyrazolyle ou thiadiazolyle, qui est éventuellement substitué par 1, 2 ou 3 substituants, choisis dans le groupe comprenant un atome d'halogène, un groupe alkyle en C₁ à C₇, alcoxy en C₁ à C₇, alcoxy en C₁ à C₇ substitué par un atome d'halogène ou NRR' ;
 R et R' représentent indépendamment l'un de l'autre un atome d'hydrogène ou un groupe alkyle en C₁ à C₇, ou peuvent former, ensemble avec l'atome d'azote auquel ils sont liés, un groupe hétérocyclique saturé à cinq ou six chaînons qui peut contenir un oxygène supplémentaire, NH, ou un groupe N-alkyle en C₁ à C₇ ;
 R¹ représente un atome d'hydrogène, un atome d'halogène ou un groupe alkyle en C₁ à C₇ substitué par un atome d'halogène ;
 R² représente un atome d'hydrogène, un groupe alkyle en C₁ à C₇, alcoxyalkyle en C₁ à C₇, cycloalkyle ou hétérocycloalkyle ;
 ou R² peut former, ensemble avec l'atome de carbone le plus proche dans le groupe G, un groupe



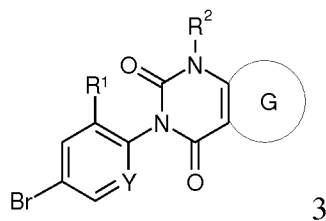
A représentant -CH₂-, -CH₂CH₂-, ou -C(CH₃)₂-,

ou un sel ou un sel d'addition d'acide pharmaceutiquement acceptable, un mélange racémique, ou ses énantiomères et/ou isomères optiques et/ou stéréoisomères correspondants.

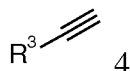
9. Composé de formule ID selon la revendication 8, ledit composé étant :

la 6-[3-chloro-5-[2-(3-pyridyl)éthynyl]-2-pyridyl]-4-isopropyl-2-méthyl-pyrazolo[4,3-d]pyrimidine-5,7-dione.

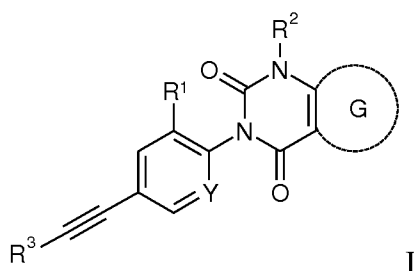
10. Procédé de fabrication d'un composé de formule I tel que défini dans l'une quelconque des revendications 1 à 9, ledit procédé comprenant la mise en réaction d'un composé de formule 3



avec un composé de formule :



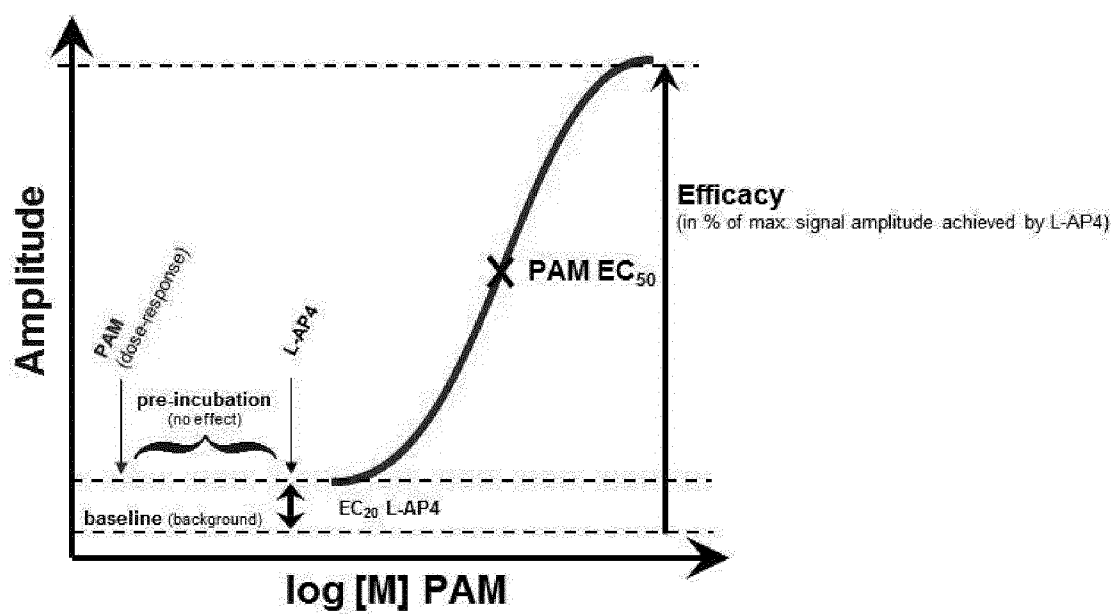
afin d'obtenir un composé de formule I



dans lesquelles les substituants sont tels que décrits dans la revendication 1, ou si cela est souhaité, la conversion des composés obtenus en des sels d'addition d'acide pharmaceutiquement acceptables.

11. Composé de formule I selon l'une quelconque des revendications 1 à 9 destiné à être utilisé comme substances thérapeutiquement actives.
12. Composé de formule I selon l'une quelconque des revendications 1 à 9 destiné à être utilisé dans le traitement de la maladie de Parkinson, de l'anxiété, des vomissements, des troubles obsessionnels-compulsifs, de l'autisme, de la neuroprotection, d'un cancer, de la dépression et du diabète de type 2.
13. Composition pharmaceutique comprenant un composé de formule I selon l'une quelconque des revendications 1 à 9 et des excipients pharmaceutiquement acceptables.
14. Utilisation d'un composé de formule I selon l'une quelconque des revendications 1 à 9 pour la préparation d'un médicament destiné au traitement de la maladie de Parkinson, de l'anxiété, des vomissements, des troubles obsessionnels-compulsifs, de l'autisme, de la neuroprotection, d'un cancer, de la dépression et du diabète de type 2.

Figure 1



REFERENCES CITED IN THE DESCRIPTION

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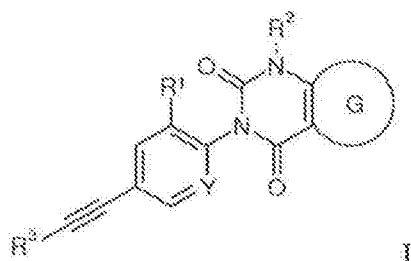
Non-patent literature cited in the description

- **CELANIRE S ; CAMPO B.** *Expert Opinion in Drug Discovery*, 2012 **[0003]**
- *Neuroreport*, 2008, vol. 19 (4), 475-8 **[0009]**
- *Proc.Natl. Acad. Sci*, 2003, vol. 100 (23), 13668-73 **[0009]**
- *J.Neurosci.*, 2006, vol. 26 (27), 7222-9 **[0009]**
- *Mol. Pharmacol.*, 2008, vol. 74 (5), 1345-58 **[0009]**
- **PREDIGER R et al.** *Neuropharmacology*, 2012, vol. 62, 115-24 **[0010]**
- *Eur. J. Pharmacol.*, 2004, vol. 498 (1-3), 153-6 **[0010]**
- *Neuropharmacology*, 2004, vol. 46 (2), 151-9 **[0012]**
- *Diabetes*, 2004, vol. 53 (4), 998-1006 **[0013]**
- *Anticancer Res.*, 2009, vol. 29 (1), 371-7 **[0014]**
- *Cli. Cancer Research*, 2005, vol. 11 (9), 3288-95 **[0014]**

Etinil származékok

Szabadalmi igénypontok

1. Az I képlet szerinti vegyület,



amelyben

Y jelentése N vagy C-R¹;

G jelentése 5- vagy 6-tagú aromás vagy 0, 1, 2 vagy 3 heteroatomot tartalmazó heteroaromás gyűrű a fenil, különböző nitrogén helyzetű piridinilek, imidazolil, pirazinil, pirimidinil, tiofenil, tiazolil, pirazolil vagy tiadiazolil csoportból kiválasztva, amely opcionálisan a halogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxi, halogénatommal vagy NRR'-vel szubsztituált 1–7 szénatomos alkoxi csoportból kiválasztott 1, 2 vagy 3 szubsztituenssel szubsztituált; R és R¹ jelentése egymástól függetlenül hidrogénatom vagy 1–7 szénatomos alkil, vagy együttesen azzal a nitrogénatommal, amelyhez csatlakoznak, öt- vagy hattagú telített heterociklusos csoportot alkotnak, amely további oxigénatomot, NH- vagy N-(1–7 szénatomos alkil)-csoportot tartalmazhat;

R¹ jelentése hidrogénatom, halogénatom vagy 1–7 szénatomos, halogénatommal szubsztituált alkil;

R¹⁷ jelentése hidrogénatom, halogénatom vagy 1–7 szénatomos, halogénatommal szubsztituált alkil;

R² jelentése hidrogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxi, cikloalkil vagy heterocikloalkil;

vagy R² együttesen a G csoport legközelebbi szénatomjával csoportot alkothat,



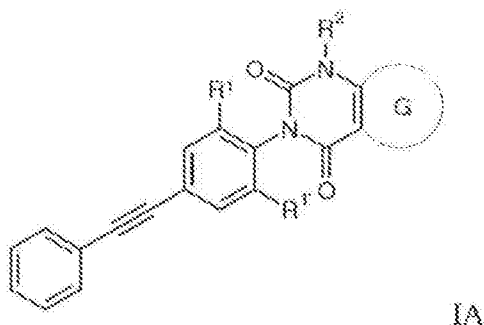
amelyben A jelentése -CH₂-, -CH₂CH₂ vagy -C(CH₃)₂-,

R³ jelentése fenil vagy piridinil, amelyben a piridinil-csoport nitrogénatomja

különböző pozíciókban lehet;

vagy ezek gyógyászatilag elfogadható sója vagy savaddíciós sója, racém elegye vagy annak megfelelő enantiomerje és/vagy optikai izomerje és/vagy sztereoizomerje.

2. Az 1. igénypont szerinti IA képletű vegyület,



amelyben

G jelentése 5- vagy 6-tagú aromás vagy 0, 1, 2 vagy 3 heteroatomot tartalmazó heteroaromás gyűrű a fenil, különböző nitrogén helyzetű piridinilek, imidazolil, pirazinil, pirimidinil, tiofenil, tiazolil, pirazolil vagy tiadiazolil csoportból kiválasztva, amely opcionálisan a halogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxi, halogénatommal vagy NRR'-vel szubsztituált 1–7 szénatomos alkoxi csoportból kiválasztott 1, 2 vagy 3 szubsztituenssel szubsztituált;

R és R' jelentése egymástól függetlenül hidrogénatom vagy 1–7 szénatomos alkil, vagy együttesen azzal a nitrogénatommal, amelyhez csatlakoznak, öt- vagy hattagú telített heterociklusos csoportot alkotnak, amely további oxigénatomot, NH- vagy N-(1–7 szénatomos alkil)-csoportot tartalmazhat;

Rⁱ jelentése hidrogénatom, halogénatom vagy 1–7 szénatomos, halogénatommal szubsztituált alkil;

R^{i'} jelentése hidrogénatom, halogénatom vagy 1–7 szénatomos, halogénatommal szubsztituált alkil;

R² jelentése hidrogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxialkil, cikloalkil vagy heterocikloalkil;

vagy R² együttesen a G csoport legközelebbi szénatomjával csoportot alkothat,



amelyben A jelentése $-\text{CH}_2-$, $-\text{CH}_2\text{CH}_2$ vagy $-\text{C}(\text{CH}_3)_2-$,

vagy ezek gyógyászatilag elfogadható sója vagy savaddíciós sója, racém elegye vagy annak megfelelő enantiomerje és/vagy optikai izomerje és/vagy sztereoizomerje.

3. Az 2. igénypont szerinti IA képletű vegyület, amely vegyület:

3-[2-klór-4-(2-feniletinil)fenil]-1-metil-kinazolin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1,8-dimetil-kinazolin-2,4-dion

8-klór-3-[2-klór-4-(2-feniletinil)fenil]-1-metil-kinazolin-2,4-dion

7-klór-3-[2-klór-4-(2-feniletinil)fenil]-1-metil-kinazolin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1,7-dimetil-kinazolin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1H-pirido[2,3-d]pirimidin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1-metil-pirido[2,3-d]pirimidin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-8-etil-1-metil-kinazolin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1-metil-pirido[3,2-d]pirimidin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1-metil-pirido[4,3-d]pirimidin-2,4-dion

1-[2-klór-4-(2-feniletinil)fenil]-3,7-dimetil-purin-2,6-dion

2-(2-klór-4-(2-feniletinil)fenil)-5,6-dihidro-1H-pirrolo[3,2,1-ij]kinazolin-1,3(2H)-dion

3-[2-klór-4-(2-feniletinil)fenil]-1-izopropil-kinazolin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1-etil-kinazolin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1-metil-pteridin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1H-pirimido[4,5-d]pirimidin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1-metil-pirimido[4,5-d]pirimidin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-6-fluor-1-metil-kinazolin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-7,8-difluor-1-metil-kinazolin-2,4-dion

3-[2-klór-4-(2-feniletinil)fenil]-1,7-dimetil-tieno[3,2-d]pirimidin-2,4-dion

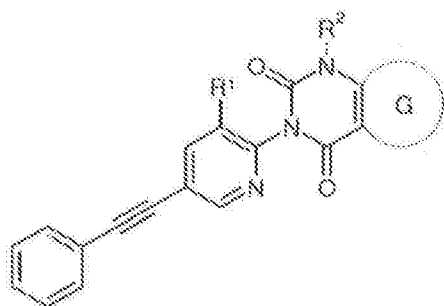
3-[2-klór-4-(2-feniletinil)fenil]-1-izopropil-pteridin-2,4-dion

3-[2,6-difluor-4-(2-feniletinil)fenil]-1-metil-kinazolin-2,4-dion

3-[2,6-difluor-4-(2-feniletinil)fenil]-1,8-dimetil-kinazolin-2,4-dion

3-(2,6-difluor-4-feniletinil-fenil)-1-metil-1H-pirido[2,3-d]pirimidin-2,4-dion
 3-[2,6-difluor-4-(2-feniletinil)fenil]-1,7-dimetil-kinazolin-2,4-dion
 3-[2,6-difluor-4-(2-feniletinil)fenil]-1,7-dimetil-tieno[3,2-d]pirimidin-2,4-dion
 2-(2,6-difluor-4-(feniletinil)fenil)-5,6-dihidro-1H-pirrolo[3,2,1-il]kinazolin-1,3(2H)-dion
 6-[2,6-difluor-4-(2-feniletinil)fenil]-4H-tiazolo[4,5-d]pirimidin-5,7-dion
 6-[2,6-difluor-4-(2-feniletinil)fenil]-4-metil-tiazolo[4,5-d]pirimidin-5,7-dion
 6-[2,6-difluor-4-(2-feniletinil)fenil]-2,4-dimetil-pirazolo[4,3-d]pirimidin-5,7-dion
 6-[2,6-difluor-4-(2-feniletinil)fenil]-1,4-dimetil-pirazolo[4,3-d]pirimidin-5,7-dion
 6-[2,6-difluor-4-(2-feniletinil)fenil]-4-izopropil-2-metil-pirazolo[4,3-d]pirimidin-5,7-dion
 6-[2,6-difluor-4-(2-feniletinil)fenil]-4-izopropil-1-metil-pirazolo[4,3-d]pirimidin-5,7-dion
 3-[2,6-difluor-4-(2-feniletinil)fenil]-1-izopropil-pirido[2,3-d]pirimidin-2,4-dion
 3-[2,6-difluor-4-(2-feniletinil)fenil]-1,8-dimetil-pirido[3,2-d]pirimidin-2,4-dion
 6-[2,6-difluor-4-(2-feniletinil)fenil]-4-metil-[1,2,5]tiadiazolo[3,4-d]pirimidin-5,7-dion
 6-[2,6-difluor-4-(2-feniletinil)fenil]-4-izopropil-[1,2,5]tiadiazolo[3,4-d]pirimidin-5,7-dion
 3-[2-klór-4-(2-feniletinil)fenil]-1,5-dimetil-kinazolin-2,4-dion
 3-[2-klór-4-(2-feniletinil)fenil]-1-(2-metoxietil)kinazolin-2,4-dion
 6-[2-klór-4-(2-feniletinil)fenil]-4-metil-2-morfolino-tiazolo[4,5-d]pirimidin-dion
 6-(2-klór-4-feniletinil-fenil)-4-metil-4H-tiazolo[4,5-d]pirimidin-5,7-dion
 3-[2-klór-4-(2-feniletinil)fenil]-1-ciklobutil-kinazolin-2,4-dion
 3-[2-klór-4-(2-feniletinil)fenil]-8-izopropil-1-metil-kinazolin-2,4-dion
 8-klór-3-[2,6-difluor-4-(2-feniletinil)fenil]-1-metil-pirido[3,2-d]pirimidin-2,4-dion
 5-[2,6-difluor-4-(2-feniletinil)fenil]-2,7-dimetil-pirazolo[3,4-d]pirimidin-4,6-dion
 5-[2,6-difluor-4-(2-feniletinil)fenil]-1,7-dimetil-pirazolo[3,4-d]pirimidin-4,6-dion
 5-[2,6-difluor-4-(2-feniletinil)fenil]-7-izopropil-2H-pirazolo[3,4-d]pirimidin-4,6-dion
 3-[2-klór-4-(2-feniletinil)fenil]-8-metoxi-1-metil-kinazolin-2,4-dion
 3-[2,6-difluor-4-(2-feniletinil)fenil]-1-(oxetán-3-il)kinazolin-2,4-dion
 6-[2-klór-4-(2-feniletinil)fenil]-4-metil-tiazolo[5,4-d]pirimidin-5,7-dion
 3-[2,6-difluor-4-(2-feniletinil)fenil]-1-izopropil-pteridin-2,4-dion
 1-[2,6-difluor-4-(2-feniletinil)fenil]-3,7-dimetil-purin-2,6-dion
 3-[2,6-difluor-4-(2-feniletinil)fenil]-8-metoxi-1-metil-kinazolin-2,4-dion
 3-[2-fluor-4-(2-feniletinil)-6-(trifluormetil)fenil]-1-izopropil-kinazolin-2,4-dion vagy
 8-(2,2-difluoretoxi)-3-[2,6-difluor-4-(2-feniletinil)fenil]-1-metil-kinazolin-2,4-dion.

4. Az 1. igénypont szerinti IB képletű vegyület,



IB

amelyben

G jelentése 5- vagy 6-tagú aromás vagy 0, 1, 2 vagy 3 heteroatomot tartalmazó heteroaromás gyűrű a fenil, különböző nitrogén helyzetű piridinilek, imidazolil, pirazinil, pirimidinil, tiofenil, tiazolil, pirazolil vagy tiadiazolil csoportból kiválasztva, amely opcionálisan a halogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxi, halogénatommal vagy NRR'-vel szubsztituált 1–7 szénatomos alkoxi csoportból kiválasztott 1, 2 vagy 3 szubsztituenssel szubsztituált;

R és R' jelentése egymástól függetlenül hidrogénatom vagy 1–7 szénatomos alkil-csoport, vagy együttesen azzal a nitrogénatommal, amelyhez csatlakoznak, öt- vagy hattagú telített heterociklusos csoportot alkotnak, amely további oxigénatomot, NH- vagy N-(1–7 szénatomos alkil)-csoportot tartalmazhat;

R¹ jelentése hidrogénatom, halogénatom vagy 1–7 szénatomos, halogénatommal szubsztituált alkil;

R² jelentése hidrogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxialkil, cikloalkil vagy heterocikloalkil;

vagy R² együttesen a G csoport legközelebbi szénatomjával csoportot alkothat,



amelyben A jelentése -CH₂-, -CH₂CH₂ vagy -C(CH₃)₂-,

vagy ezek gyógyászatilag elfogadható sója vagy savaddíciós sója, racém elegye vagy annak megfelelő enantiomerje és/vagy optikai izomerje és/vagy sztereoizomerje.

5. A 4. igénypont szerinti IB képletű vegyület, amely vegyület:

3-[3-klór-5-(2-feniletinil)-2-piridil]-1-metil-kinazolin-2,4-dion

3-[3-klór-5-(2-feniletinil)-2-piridil]-1-izopropil-kinazolin-2,4-dion

3-[3-klór-5-(2-feniletinil)-2-piridil]-1,8-dimetil-kinazolin-2,4-dion

2-(3-klór-5-(2-feniletinil)piridin-2-il)-5,6-dihidro-1H-pirrolo[3,2,1-ij]kinazolin-1,3(2H)-dion

1-metil-3-[5-(2-feniletinil)-3-(trifluormetil)-2-piridil]kinazolin-2,4-dion

3-[3-fluor-5-(2-feniletinil)-2-piridil]-1,8-dimetil-kinazolin-2,4-dion

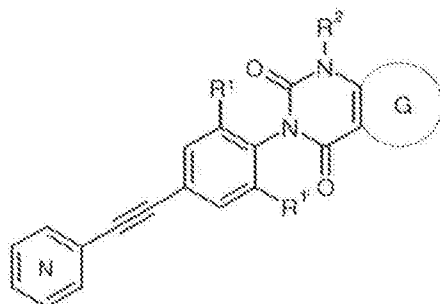
1-izopropil-3-[5-(2-feniletinil)-2-piridil]kinazolin-2,4-dion

6-[3-fluor-5-(2-feniletinil)-2-piridil]-4-izopropil-2-metil-pirazolo[4,3-d]pirimidin-5,7-dion

vagy

6-[3-klór-5-(2-feniletinil)-2-piridil]-4-izopropil-2-metil-pirazolo[4,3-d]pirimidin-5,7-dion.

6. Az 1. igénypont szerinti IC képletű vegyület,



IC,

amelyben

G jelentése 5- vagy 6-tagú aromás vagy 0, 1, 2 vagy 3 heteroatomot tartalmazó heteroaromás gyűrű a fenil, különböző nitrogén helyzetű piridinilek, imidazolil, pirazinil, pirimidinil, tiofenil, tiazolil, pirazolil vagy tiadiazolil csoportból kiválasztva, amely opcionálisan a halogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxi, halogénatommal vagy NRR'-vel szubsztituált 1–7 szénatomos alkoxi csoportból kiválasztott 1, 2 vagy 3 szubsztituenssel szubsztituált;

R és R' jelentése egymástól függetlenül hidrogénatom vagy 1–7 szénatomos alkil-csoport, vagy együttesen azzal a nitrogénatommal, amelyhez csatlakoznak, öt- vagy hattagú telített heterociklusos csoportot alkotnak, amely további oxigénatomot, NH- vagy N-(1–7 szénatomos alkil)-csoportot tartalmazhat;

R^1 jelentése hidrogénatom, halogénatom vagy 1–7 szénatomos, halogénatommal szubsztituált alkil;

$R^{1'}$ jelentése hidrogénatom, halogénatom vagy 1–7 szénatomos, halogénatommal szubsztituált alkil;

R^2 jelentése hidrogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoalkil, cikloalkil vagy heterocikloalkil;

vagy R^2 együttesen a G csoport legközelebbi szénatomjával csoportot alkothat,



amelyben A jelentése $-\text{CH}_2-$, $-\text{CH}_2\text{CH}_2$ vagy $-\text{C}(\text{CH}_3)_2-$,

vagy ezek gyógyászatilag elfogadható sója vagy savaddíciós sója, racém elegye vagy annak megfelelő enantiomerje és/vagy optikai izomerje és/vagy sztereoizomerje.

7. Az 6. igénypont szerinti IC képletű vegyület, amely vegyület:

3-[2-klór-4-[2-(3-piridil)etinil]fenil]-1-izopropil-pteridin-2,4-dion

3-[2,6-difluor-4-[2-(3-piridil)etinil]fenil]-1,7-dimetil-tieno[3,2-d]pirimidin-2,4-dion

2-(2,6-difluor-4-(piridin-3-iletinil)fenil)-5,6-dihidro-1H-pirroló[3,2,1-ij]kinazolin-1,3(2H)-dion

3-[2-klór-6-fluor-4-[2-(3-piridil)etinil]fenil]-1-izopropil-kinazolin-2,4-dion

3-[2-klór-6-fluor-4-[2-(3-piridil)etinil]fenil]-1,8-dimetil-kinazolin-2,4-dion

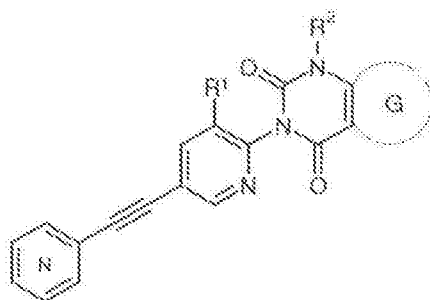
6-[2,6-difluor-4-[2-(3-piridil)etinil]fenil]-4-izopropil-2-metil-pirazolo[4,3-d]pirimidini-5,7-dion

6-[2-klór-6-fluor-4-[2-(3-piridil)etinil]fenil]-4-izopropil-2-metil-pirazolo[4,3-d]pirimidin-5,7-dion

3-[2-klór-6-fluor-4-[2-(3-piridil)etinil]fenil]-1-(oxetán-3-il)kinazolin-2,4-dion vagy

3-[2,6-difluor-4-[2-(3-piridil)etinil]fenil]-1,8-dimetil-kinazolin-2,4-dion.

8. Az 1. igénypont szerinti ID képletű vegyület,



ID

amelyben

G jelentése 5- vagy 6-tagú aromás vagy 0, 1, 2 vagy 3 heteroatomot tartalmazó heteroaromás gyűrű a fenil, különböző nitrogén helyzetű piridinilek, imidazolil, pirazinil, pirimidinil, tiofenil, tiazolil, pirazolil vagy tiadiazolil csoportból kiválasztva, amely opcionálisan a halogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxi, halogénatommal vagy NRR^1 -vel szubsztituált 1–7 szénatomos alkoxi csoportból kiválasztott 1, 2 vagy 3 szubsztituenssel szubsztituált;

R és R^1 jelentése egymástól függetlenül hidrogénatom vagy 1–7 szénatomos alkil-csoport, vagy együttesen azzal a nitrogénatommal, amelyhez csatlakoznak, öt- vagy hattagú telített heterociklusos csoportot alkotnak, amely további oxigénatomot, NH - vagy N -(1–7 szénatomos alkil)-csoportot tartalmazhat;

R^1 jelentése hidrogénatom, halogénatom vagy 1–7 szénatomos, halogénatommal szubsztituált alkil;

R^2 jelentése hidrogénatom, 1–7 szénatomos alkil, 1–7 szénatomos alkoxialkil, cikloalkil vagy heterocikloalkil;

vagy R együttesen a G csoport legközelebbi szénatomjával csoportot alkothat,



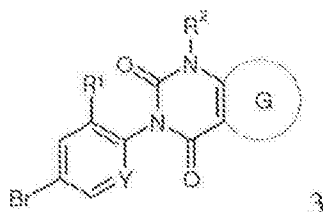
amelyben A jelentése $-\text{CH}_2-$, $-\text{CH}_2\text{CH}_2$ vagy $-\text{C}(\text{CH}_3)_2-$,

vagy ezek gyógyászatilag elfogadható sója vagy savaddíciós sója, racém elegye vagy annak megfelelő enantiomerje és/vagy optikai izomerje és/vagy sztereoizomerje.

9. A 8. igénypont szerinti ID képletű vegyület, amely vegyület a 6-[3-klór-5-[2-(3-piridil)etilil]-2-piridil]-4-izopropil-2-metil-pirazolo[4,3-d] pirimidin-5,7-dion.

10. Eljárás az 1–9. igénypontok bármelyike szerinti I általános képletű vegyület előállítására, amely eljárás tartalmazza

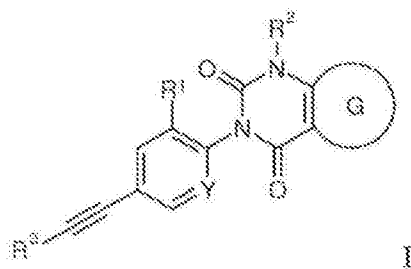
a 3 képletű vegyület reagáltatását



az alábbi képletű vegyülettel



az I képletű vegyület létrehozására,



ahol a szubsztituenseket az 1. igénypont írja le, vagy

kívánt esetben a kapott vegyületek gyógyszerészetileg elfogadható savaddíciós sókká történő átalakítását.

11. Az 1–9. igénypontok bármelyike szerinti I képletű vegyület terápiásan aktív anyagként történő alkalmazásra.

12. Az 1–9. igénypontok bármelyike szerinti I képletű vegyület Parkinson-kór, szorongás, hányás, rögeszmés kényszerbetegség, autizmus, idegvédelem, rák, depresszió és 2-es típusú cukorbetegség kezelésében történő alkalmazásra.

13. Az I általános képletű, az 1–9. igénypontok bármelyike szerinti vegyületet és

gyógyászati lag elfogadható excipienst tartalmazó gyógyászati készítmény.

14. Az 1–9. igénypontok bármelyike szerinti, I általános képletű vegyület alkalmazása Parkinson-kór, szorongás, hányás, rögeszmés kényszerbetegség, autizmus, idegvédelem, rák, depresszió és 2-es típusú cukorbetegség kezelésére szolgáló gyógyszer előállítására.