

1,2,4-TRIAZOLO[4,3-a]PYRIDINE COMPOUNDS AND THEIR USE AS POSITIVE ALLOSTERIC MODULATORS OF MGLUR2 RECEPTORS

5 FIELD OF THE INVENTION

The present invention relates to novel 1,2,4-triazolo[4,3-a]pyridine compounds as positive allosteric modulators (PAMs) of the metabotropic glutamate receptor subtype 2 (“mGluR2”). The invention is also directed to pharmaceutical compositions comprising such compounds, to processes for preparing such compounds and
10 compositions, and to the use of such compounds and compositions for the prevention or treatment of disorders in which mGluR2 subtype of metabotropic receptors is involved.

BACKGROUND OF THE INVENTION

Glutamate is the major amino acid neurotransmitter in the mammalian central
15 nervous system. Glutamate plays a major role in numerous physiological functions, such as learning and memory but also sensory perception, development of synaptic plasticity, motor control, respiration, and regulation of cardiovascular function. Furthermore, glutamate is at the centre of several different neurological and psychiatric diseases, where there is an imbalance in glutamatergic neurotransmission.

20 Glutamate mediates synaptic neurotransmission through the activation of ionotropic glutamate receptor channels (iGluRs), and the NMDA, AMPA and kainate receptors which are responsible for fast excitatory transmission.

In addition, glutamate activates metabotropic glutamate receptors (mGluRs) which have a more modulatory role that contributes to the fine-tuning of synaptic
25 efficacy.

Glutamate activates the mGluRs through binding to the large extracellular amino-terminal domain of the receptor, herein called the orthosteric binding site. This binding induces a conformational change in the receptor which results in the activation of the G-protein and intracellular signaling pathways.

30 The mGluR2 subtype is negatively coupled to adenylate cyclase via activation of G α i-protein, and its activation leads to inhibition of glutamate release in the synapse. In the central nervous system (CNS), mGlu2 receptors are abundant mainly throughout cortex, thalamic regions, accessory olfactory bulb, hippocampus, amygdala, caudate-putamen and nucleus accumbens.

Activating mGluR2 was shown in clinical trials to be efficacious to treat anxiety disorders (for studies with orthosteric mGlu2/3 agonists, see Michelson et al. *Neuropharmacology* 2005, 49(S1), 84-257; Dunayevich et al. *Neuropsychopharmacology* 2008, 33(7), 1603-10), LY354740 had been previously
5 evaluated in non-clinical and clinical model systems predicting utility in the treatment of anxiety disorders beyond generalized anxiety depression (GAD), e.g. panic (see Dunayevich et al. 2008). Non-clinical studies, suggest a role for both mGlu2 and mGlu3 receptors in anxiolysis (Linden et al. *Neuropharmacology* 2005, 49, 120-134) whilst it has been suggested that positive allosteric modulation of the mGluR2 may be
10 sufficient for an anxiolytic effect (Johnson et al. *Psychopharmacology* (Berl) 2005, 179(1), 271-283).
In addition, activating mGluR2 was shown to be potentially efficacious for the treatment of
(a) schizophrenia (Patil et al. *Nat Med* 2007, 13(9), 1102-7); later studies however, do
15 not support treatment of acute exacerbations of schizophrenia with an mGluR2 agonist or allosteric modulator (Adams et al. *BMC Psychiatry* 2013, 13(1), 143; Kinon et al. *J Clin Psychopharmacol.* 2013, 31(3), 349-55; Litman et al. (2013) NCDEU Meeting (abstract)) but do not exclude application for other specific symptom clusters (e.g. negative symptoms (Kent et al. "Safety, tolerability and potential therapeutic efficacy
20 of a novel glutamate modulator as adjunctive treatment in patients with schizophrenia" abstract No. 3160 and poster NR10-47, American Psychiatric Association 166th Annual Meeting 2013 (APA 2013), May 18-22, 2013, San Francisco, California, USA)) or for other phases in the disease (e.g. residual symptoms);
(b) epilepsy, based on acute non-clinical studies with mixed mGlu2/3 receptor agonists
25 (Moldrich et al. *Eur J Pharmacol.* 2003, 476, 3-16; Barton et al. *Epilepsy Research* 2003, 56, 17-26); continued administration of an mGlu2/3 agonist paradoxically induced seizure activity in long-term toxicology studies (Dunayevich et al. (2008), this paradoxical effect may be related to agonist-induced changes in the sensitivity of the receptor systems (tachyphylaxis); positive allosteric modulators, in contrast, modulate
30 ongoing neurotransmission but are not directly stimulatory, thereby reducing the risk for tachyphylaxis;
(c) drug addiction/dependence (Barrett, *Neuropsychopharmacology* 2010, 35, 2007-2008; Foster, *Curr Drug Abuse Rev* 2009, 2, 83-98);
(d) Parkinson's disease (see for example Johnson et al. *CNS Neurol Disord Drug*
35 *Targets* 2009, 8, 475-491; Konieczny et al. *Naunyn Schmiedebergs Arch. Pharmacol.* 1998, 358 (4), 500-502);

(e) pain (Chiechio and Nicoletti, *Curr Opin Pharmacol* 2012, 12, 28-34; Jones et al. *Neuropharmacology* 2005, 49, 206-218; Neugebauer, [Review] *Pain* 2002, 98 (1-2), 1-8; Simmons et al. *Pharmacology, Biochemistry and Behavior* 2002, 73, 419-427);
5 (f) sleep disorders (Ahnaou et al. *European Journal of Pharmacology* 2009, 603, 62-72);

(f) Huntington's disease (based on a potential disease modifying effect (Schiefer et al. *Brain Res* 2004, 1019, 246 – 254) which is to be confirmed further); and

(g) depression (although no efficacy signal was detected on the primary outcome measure, adjunctive administration of JNJ-40411813/ADX71149 in the dose range
10 tested in a multicenter, double-blind, placebo-controlled study in adults with major depressive disorder with anxiety symptoms showed efficacy signals on several secondary outcome measures of both depression and anxiety (Kent et al. "Efficacy and Safety of a Novel mGlu2 Receptor Positive Allosteric Modulator as an Adjunctive Treatment to an SSRI/SNRI in the Treatment of Anxious Depression", Abstract to
15 poster and oral presentation, American Society of Clinical Psychopharmacology (ASCP) 2014 Annual Meeting, June 16-19, 2014 Westin Diplomat, Hollywood, Florida)).

A new avenue for developing selective compounds acting at mGluRs is to identify compounds that act through allosteric mechanisms, modulating the receptor by
20 binding to a site different from the highly conserved orthosteric binding site.

Positive allosteric modulators of mGluRs have emerged recently as novel pharmacological entities offering this attractive alternative.

It was demonstrated that such compounds do not activate the receptor by themselves. Rather, they enable the receptor to produce an increased response to a
25 concentration of glutamate, which by itself induces a minimal response. Mutational analysis has demonstrated unequivocally that the binding of mGluR2 positive allosteric modulators does not occur at the orthosteric site, but instead at an allosteric site situated within the seven transmembrane region of the receptor.

Animal data suggest that positive allosteric modulators of mGluR2 have effects
30 in anxiety and psychosis models similar to those obtained with orthosteric agonists. Allosteric modulators of mGluR2 were shown to be active in fear-potentiated startle (Johnson et al. *J Med Chem* 2003, 46, 3189-3192; Johnson et al. *Psychopharmacology* 2005, 179, 271-283), and in stress-induced hyperthermia models of anxiety (Johnson et al. 2005). Furthermore, such compounds were shown to be active in reversal of
35 ketamine- (Govek et al. *Bioorg Med Chem Lett* 2005, 15(18), 4058-4072) or

amphetamine- (Galici et al. J Pharm Exp Ther 2005, 315(3), 1181-1187) induced hyperlocomotion, and in reversal of amphetamine-induced disruption of prepulse inhibition of the acoustic startle effect (Galici et al. 2005) models of schizophrenia.

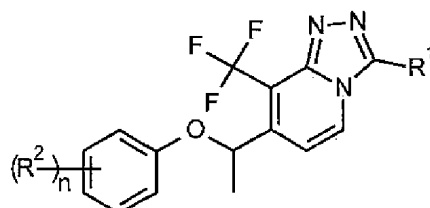
JNJ-40411813/ADX71149, an mGlu2 PAM (which in rat also displays 5-HT_{2A} antagonism activity due to a rat-specific metabolite) has undergone clinical trials for the treatment of schizophrenia, and anxiety-depression (see for instance www.Clinicaltrials.gov). Non-clinical data in the lactate-induced panic model in rodents suggests that it could have potential in the treatment of further anxiety disorders such as panic disorder and phobias, such as agoraphobia (Shekhar et al. Neuropsychopharmacology 2013, 38, S435-S593 (W220)). JNJ-40411813 was also observed to reduce craving and improve smoking cessation-induced deficits in attention and episodic memory versus placebo (Salih et al. Journal of Psychopharmacology, submitted) and showed an efficacy signal in S-ketamine-induced negative symptoms in healthy volunteers and patients with predominant negative symptoms of schizophrenia (De Boer et al. Society of Biological Psychiatry 68th Annual Scientific Convention of Society of Biological Psychiatry, May 16-18, 2013, Hilton Union Square, San Francisco, California, Abstract 2013-P-1060-SOBP).

Positive allosteric modulators enable potentiation of the glutamate response, but they have also been shown to potentiate the response to orthosteric mGluR2 agonists such as LY379268 or DCG-IV. These data provide evidence for yet another novel therapeutic approach to treat the above mentioned neurological and psychiatric diseases involving mGluR2, which would use a combination of a positive allosteric modulator of mGluR2 together with an orthosteric agonist of mGluR2.

Various compounds have been described as mGluR2 positive allosteric modulators. WO2010/130424, WO2010/130423, WO2010/130422, and WO2012/062750, WO2012/062751, and WO2012/062759, published on 18 November 2010 and 18 May 2012, respectively, disclose 1,2,4-triazolo[4,3-a]pyridine derivatives as mGluR2 positive allosteric modulators.

SUMMARY

In one aspect, there is provided a compound of Formula (I)



or a stereochemically isomeric form thereof, wherein

R¹ is selected from the group consisting of C₁₋₆alkyl, (C₃₋₈cycloalkyl)C₁₋₃alkyl, and (C₁₋₃alkyloxy)C₁₋₃alkyl;

each R² is independently selected from F, Cl, C₁₋₃alkyl, C₁₋₃alkyloxy, mono- or
5 polyhaloC₁₋₃alkyl, and mono- or polyhaloC₁₋₃alkyloxy;

n is an integer selected from 1, 2, and 3;

or a pharmaceutically acceptable salt or solvate thereof.

In another aspect, there is provided a pharmaceutical composition comprising a
therapeutically effective amount of a compound as defined herein and a
10 pharmaceutically acceptable carrier or excipient.

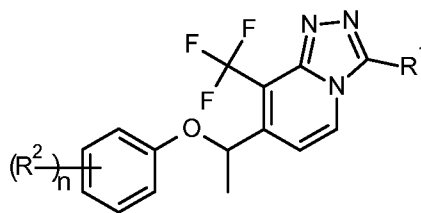
In another aspect, there is provided a compound as defined herein or a pharmaceutical
composition as defined herein, for use as a medicament.

In another aspect, there is provided a compound as defined herein or a pharmaceutical
composition as defined herein for use in the treatment or prevention of a central
15 nervous system disorder selected from the group of anxiety disorders, psychotic
disorders selected from the group consisting of schizophrenia, schizoaffective disorder
and schizophreniform disorder, personality disorders, substance-related disorders,
eating disorders, mood disorders, migraine, epilepsy or convulsive disorders, childhood
disorders, cognitive disorders, neurodegeneration, autistic disorders, neurotoxicity and
20 ischemia.

In another aspect, there is provided a product comprising (a) a compound as defined
herein; and (b) an mGluR2 orthosteric agonist, as a combined preparation for
simultaneous, separate or sequential use in the treatment or prevention of a central
nervous system disorder selected from the group of schizophrenia (in particular,
25 negative symptoms or residual symptoms thereof), generalized anxiety disorder, bipolar
disorder (I or II), migraine, behavioral and psychological symptoms of dementia,
epilepsy or convulsive disorders, panic disorder, mixed anxiety and depression, and
agoraphobia.

30 DETAILED DESCRIPTION OF THE INVENTION

The present invention is directed to potent mGluR2 PAM compounds with an
advantageous balance of properties. Thus, the present invention is directed to 1,2,4-
triazolo[4,3-a]pyridine derivatives of Formula (I)



and the stereochemically isomeric forms thereof, wherein

R¹ is selected from the group consisting of C₁₋₆alkyl, (C₃₋₈cycloalkyl)C₁₋₃alkyl, and (C₁₋₃alkyloxy)C₁₋₃alkyl;

- 5 each R² is independently selected from F, Cl, C₁₋₃alkyl, C₁₋₃alkyloxy, mono- or polyhaloC₁₋₃alkyl, and mono- or polyhaloC₁₋₃alkyloxy;
n is an integer selected from 1, 2, and 3;
and the pharmaceutically acceptable salts and the solvates thereof.

- The present invention also relates to a pharmaceutical composition comprising a
10 therapeutically effective amount of a compound of Formula (I) and a pharmaceutically acceptable carrier or excipient.

- Additionally, the invention relates to a compound of Formula (I) for use as a medicament and to a compound of Formula (I) for use as a medicament for the treatment or prevention of neurological and psychiatric disorders in which mGluR2 is
15 involved.

The invention also relates to the use of a compound according to Formula (I) or a pharmaceutical composition according to the invention for the manufacture of a medicament for treating or preventing neurological and psychiatric disorders in which mGluR2 is involved.

- 20 Additionally, the invention relates to the use of a compound of Formula (I) in combination with an additional pharmaceutical agent for the manufacture of a medicament for treating or preventing neurological and psychiatric disorders in which mGluR2 is involved.

- Furthermore, the invention relates to a process for preparing a pharmaceutical
25 composition according to the invention, characterized in that a pharmaceutically acceptable carrier is intimately mixed with a therapeutically effective amount of a compound of Formula (I).

- The invention also relates to a product comprising a compound of Formula (I) and an additional pharmaceutical agent, as a combined preparation for simultaneous, separate
30 or sequential use in the treatment or prevention of neurological or psychiatric disorders and diseases.

The present invention relates in particular to compounds of Formula (I) as defined hereinabove, and stereoisomeric forms thereof, wherein

R^1 is selected from the group consisting of CH_3CH_2 , $CH_3CH_2CH_2$,

(cyclopropyl)methyl, (cyclobutyl)methyl, ethyloxymethyl and methyloxymethyl; and

5 the rest of variables are as defined herein; and the pharmaceutically acceptable salts and the solvates thereof.

In a further embodiment, the present invention relates to compounds of Formula (I) as defined hereinabove, and stereoisomeric forms thereof, wherein

R^1 is selected from the group consisting of CH_3CH_2 , (cyclopropyl)methyl,

10 (cyclobutyl)methyl and methyloxymethyl; and the rest of variables are as defined herein; and the pharmaceutically acceptable salts and the solvates thereof.

In a further embodiment, the present invention relates to compounds of Formula (I) as defined hereinabove, and stereoisomeric forms thereof, wherein

R^1 is selected from the group consisting of CH_3CH_2 , (cyclopropyl)methyl,

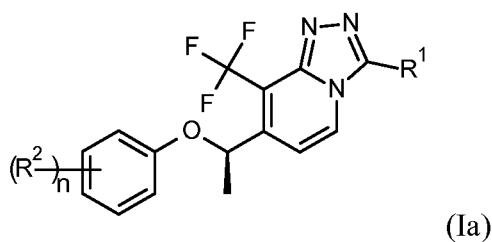
15 (cyclobutyl)methyl and ethyloxymethyl; and the rest of variables are as defined herein; and the pharmaceutically acceptable salts and the solvates thereof.

In an additional embodiment, the invention relates to a compound of Formula (I) as defined hereinabove, and stereoisomeric forms thereof wherein

each R^2 is independently selected from F, Cl, CH_3 , CH_3O and CF_3 ; and the

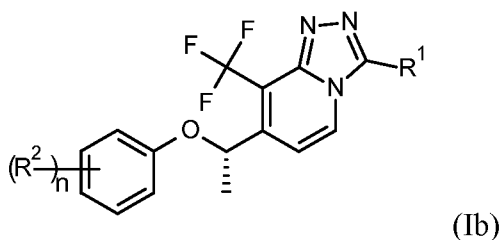
20 pharmaceutically acceptable salts and the solvates thereof.

In a further embodiment, the present invention relates to compounds of Formula (I) as defined herein having the Formula (Ia)



25 wherein the variables are as defined in Formula (I) herein, and the pharmaceutically acceptable salts and the solvates thereof.

In a further embodiment, the present invention relates to compounds of Formula (I) as defined herein having the Formula (Ib)



wherein the variables are as defined in Formula (I) herein, and the pharmaceutically acceptable salts and the solvates thereof.

Particular compounds may be selected from the group of

- 3-(Cyclopropylmethyl)-7-[1-(4-fluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1*R)-1-(4-fluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1*S)-1-(4-fluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1S)-1-(2,4-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1R)-1-(2,4-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[1-(2,4-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1S)-1-(3,5-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1S)-1-(3,4-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1S)-1-(2,3-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1S)-1-(2,5-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1S)-1-(2,6-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclopropylmethyl)-7-[(1S)-1-(4-fluoro-2-methoxyphenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 3-(Cyclobutylmethyl)-7-[1-(2,4-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;
- 7-[(1S)-1-(2-Chloro-4-methylphenoxy)ethyl]-3-(cyclopropylmethyl)-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

3-(Cyclopropylmethyl)-7-[(1S)-1-(4-fluoro-2-methylphenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

3-(Cyclopropylmethyl)-8-(trifluoromethyl)-7-[(1S)-1-(2,4,6-trifluorophenoxy)ethyl][1,2,4]triazolo[4,3-a]pyridine;

7-[1-(2,4-Difluorophenoxy)ethyl]-3-(ethoxymethyl)-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

3-Ethyl-8-(trifluoromethyl)-7-[1-(2,4,6-trifluorophenoxy)ethyl][1,2,4]triazolo[4,3-a]pyridine;

7-[1-(2,4-Difluorophenoxy)ethyl]-3-ethyl-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

3-(Cyclobutylmethyl)-7-[(1*R)-1-(2,4-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine ;

3-(Cyclobutylmethyl)-7-[(1*S)-1-(2,4-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

3-(Ethoxymethyl)-8-(trifluoromethyl)-7-[(1*R)-1-(2,4,6-trifluorophenoxy)ethyl][1,2,4]triazolo[4,3-a]pyridine;

3-(Ethoxymethyl)-8-(trifluoromethyl)-7-[(1*S)-1-(2,4,6-trifluorophenoxy)ethyl][1,2,4]triazolo[4,3-a]pyridine;

7-[(1*S)-1-(2,4-Difluorophenoxy)ethyl]-3-(ethoxymethyl)-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

7-[(1*R)-1-(2,4-Difluorophenoxy)ethyl]-3-(ethoxymethyl)-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

7-[(1*R)-1-(2,4-Difluorophenoxy)ethyl]-3-ethyl-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

7-[(1*S)-1-(2,4-Difluorophenoxy)ethyl]-3-ethyl-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

7-[1-(2,4-Difluorophenoxy)ethyl]-3-propyl-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine;

3-Ethyl-8-(trifluoromethyl)-7-[(1*R)-1-(2,4,6-trifluorophenoxy)ethyl]-[1,2,4]triazolo[4,3-a]pyridine;

3-Ethyl-8-(trifluoromethyl)-7-[(1*S)-1-(2,4,6-trifluorophenoxy)ethyl]-[1,2,4]triazolo[4,3-a]pyridine;

7-[(1*R)-(2,4-difluorophenoxy)ethyl]-3-propyl-8-(trifluoromethyl)-[1,2,4]triazolo[4,3-a]pyridine; and

7-[(1*S)-(2,4-difluorophenoxy)ethyl]-3-propyl-8-(trifluoromethyl)-[1,2,4]triazolo[4,3-a]pyridine.

Included within the scope of this list are stereoisomeric forms, the pharmaceutically acceptable salts and the solvates thereof.

In an additional embodiment, the compound may be selected from 3-(Cyclopropylmethyl)-7-[(1S)-1-(2,4-difluorophenoxy)ethyl]-8-(trifluoromethyl)[1,2,4]triazolo[4,3-a]pyridine hydrochloride salt.

- 5 The names of the compounds of the present invention were generated according to the nomenclature rules agreed upon by the Chemical Abstracts Service (C.A.S.) using Advanced Chemical Development, Inc., software (ACD/Name product version 10.01.0.14105, October 2006). In case of tautomeric forms, the name of the depicted tautomeric form of the structure was generated. However it should be clear that the
10 other non-depicted tautomeric form is also included within the scope of the present invention.

Definitions

- The notation "C₁₋₃alkyl" or "C₁₋₆alkyl" as used herein alone or as part of another
15 group, defines a saturated, straight or branched, hydrocarbon radical having, unless otherwise stated, from 1 to 3 or 1 to 6 carbon atoms, such as methyl, ethyl, 1-propyl, 1-methylethyl, butyl, 1-methyl-propyl, 2-methyl-1-propyl, 1,1-dimethylethyl, 3-methyl-1-butyl, 1-pentyl, 1-hexyl and the like.

- The notation "C₃₋₈cycloalkyl" as used herein alone or as part of another group, defines
20 a saturated, cyclic hydrocarbon radical having from 3 to 8 carbon atoms, such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and cyclooctyl.

The notation "halogen" or "halo" as used herein alone or as part of another group, refers to fluoro, chloro, bromo or iodo, with fluoro or chloro being preferred.

- The notation "mono- and polyhaloC₁₋₃alkyl" as used herein alone or as part of another
25 group, refers to C₁₋₃alkyl as defined before, substituted with 1, 2, 3 or where possible with more halo atoms as defined before.

- Whenever the term "substituted" is used in the present invention, it is meant, unless otherwise is indicated or is clear from the context, to indicate that one or more hydrogens, preferably from 1 to 3 hydrogens, more preferably from 1 to 2 hydrogens,
30 more preferably 1 hydrogen, on the atom or radical indicated in the expression using "substituted" are replaced with a selection from the indicated group, provided that the normal valency is not exceeded, and that the substitution results in a chemically stable compound, i.e. a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture, and formulation into a therapeutic agent.

The term "subject" as used herein, refers to an animal, preferably a mammal, most preferably a human, who is or has been the object of treatment, observation or experiment.

5 The term "therapeutically effective amount" as used herein, means that amount of active compound or pharmaceutical agent that elicits the biological or medicinal response in a tissue system, animal or human that is being sought by a researcher, veterinarian, medical doctor or other clinician, which includes alleviation of the symptoms of the disease or disorder being treated.

10 As used herein, the term "composition" is intended to encompass a product comprising the specified ingredients in the specified amounts, as well as any product which results, directly or indirectly, from combinations of the specified ingredients in the specified amounts.

15 It will be appreciated that some of the compounds of Formula (I) and their pharmaceutically acceptable addition salts and solvates thereof may contain one or more centres of chirality and exist as stereoisomeric forms.

The term "compounds of the invention" as used herein, is meant to include the compounds of Formula (I), and the salts and solvates thereof.

20 As used herein, any chemical formula with bonds shown only as solid lines and not as solid wedged or hashed wedged bonds, or otherwise indicated as having a particular configuration (e.g. R, S) around one or more atoms, contemplates each possible stereoisomer, or mixture of two or more stereoisomers.

Hereinbefore and hereinafter, the term "compound of Formula (I)" is meant to include the stereoisomers thereof and the tautomeric forms thereof.

25 The terms "stereoisomers", "stereoisomeric forms" or "stereochemically isomeric forms" hereinbefore or hereinafter are used interchangeably.

The invention includes all stereoisomers of the compounds of the invention either as a pure stereoisomer or as a mixture of two or more stereoisomers.

Enantiomers are stereoisomers that are non-superimposable mirror images of each other. A 1:1 mixture of a pair of enantiomers is a racemate or racemic mixture.

30 Diastereomers (or diastereoisomers) are stereoisomers that are not enantiomers, i.e. they are not related as mirror images. If a compound contains a double bond, the substituents may be in the E or the Z configuration. Substituents on bivalent cyclic (partially) saturated radicals may have either the cis- or trans-configuration; for example if a compound contains a disubstituted cycloalkyl group, the substituents may
35 be in the cis or trans configuration.

Therefore, the invention includes enantiomers, diastereomers, racemates, E isomers, Z isomers, cis isomers, trans isomers and mixtures thereof, whenever chemically

possible.

The meaning of all those terms, i.e. enantiomers, diastereomers, racemates, E isomers, Z isomers, cis isomers, trans isomers and mixtures thereof are known to the skilled person.

- 5 The absolute configuration is specified according to the Cahn-Ingold-Prelog system. The configuration at an asymmetric atom is specified by either R or S. Resolved stereoisomers whose absolute configuration is not known can be designated by (+) or (-) depending on the direction in which they rotate plane polarized light. For instance, resolved enantiomers whose absolute configuration is not known can be designated by
- 10 (+) or (-) depending on the direction in which they rotate plane polarized light. When a specific stereoisomer is identified, this means that said stereoisomer is substantially free, i.e. associated with less than 50%, preferably less than 20%, more preferably less than 10%, even more preferably less than 5%, in particular less than 2% and most preferably less than 1%, of the other isomers. Thus, when a compound of
- 15 Formula (I) is for instance specified as (R), this means that the compound is substantially free of the (S) isomer; when a compound of Formula (I) is for instance specified as E, this means that the compound is substantially free of the Z isomer; when a compound of Formula (I) is for instance specified as cis, this means that the compound is substantially free of the trans isomer.
- 20 Some of the compounds according to Formula (I) may also exist in their tautomeric form. Such forms in so far as they may exist, although not explicitly indicated in the above formula are intended to be included within the scope of the present invention. It follows that a single compound may exist in both stereisomeric and tautomeric forms.
- 25 For therapeutic use, salts of the compounds of Formula (I) are those wherein the counterion is pharmaceutically acceptable. However, salts of acids and bases which are non-pharmaceutically acceptable may also find use, for example, in the preparation or purification of a pharmaceutically acceptable compound. All salts, whether pharmaceutically acceptable or not, are included within the ambit of the present
- 30 invention.

The pharmaceutically acceptable acid and base addition salts as mentioned hereinabove or hereinafter are meant to comprise the therapeutically active non-toxic acid and base addition salt forms which the compounds of Formula (I) are able to form. The pharmaceutically acceptable acid addition salts can conveniently be obtained by

35 treating the base form with such appropriate acid. Appropriate acids comprise, for example, inorganic acids such as hydrohalic acids, e.g. hydrochloric or hydrobromic

acid, sulfuric, nitric, phosphoric and the like acids; or organic acids such as, for example, acetic, propanoic, hydroxyacetic, lactic, pyruvic, oxalic (i.e. ethanedioic), malonic, succinic (i.e. butanedioic acid), maleic, fumaric, malic, tartaric, citric, methanesulfonic, ethanesulfonic, benzenesulfonic, p-toluenesulfonic, cyclamic,
5 salicylic, p-aminosalicylic, pamoic and the like acids. Conversely said salt forms can be converted by treatment with an appropriate base into the free base form.

The compounds of Formula (I) containing an acidic proton may also be converted into their non-toxic metal or amine addition salt forms by treatment with appropriate organic and inorganic bases. Appropriate base salt forms comprise, for example, the
10 ammonium salts, the alkali and earth alkaline metal salts, e.g. the lithium, sodium, potassium, magnesium, calcium salts and the like, salts with organic bases, e.g. primary, secondary and tertiary aliphatic and aromatic amines such as methylamine, ethylamine, propylamine, isopropylamine, the four butylamine isomers, dimethylamine, diethylamine, diethanolamine, dipropylamine, diisopropylamine,
15 di-*n*-butylamine, pyrrolidine, piperidine, morpholine, trimethylamine, triethylamine, tripropylamine, quinuclidine, pyridine, quinoline and isoquinoline; the benzathine, *N*-methyl-D-glucamine, hydrabamine salts, and salts with amino acids such as, for example, arginine, lysine and the like. Conversely the salt form can be converted by treatment with acid into the free acid form.

20 The term solvate comprises the solvent addition forms as well as the salts thereof, which the compounds of Formula (I) are able to form. Examples of such solvent addition forms are e.g. hydrates, alcoholates and the like.

In the framework of this application, an element, in particular when mentioned in relation to a compound according to Formula (I), comprises all isotopes and isotopic
25 mixtures of this element, either naturally occurring or synthetically produced, either with natural abundance or in an isotopically enriched form. Radiolabelled compounds of Formula (I) may comprise a radioactive isotope selected from the group of ^3H , ^{11}C , ^{18}F , ^{122}I , ^{123}I , ^{125}I , ^{131}I , ^{75}Br , ^{76}Br , ^{77}Br and ^{82}Br . Preferably, the radioactive isotope is selected from the group of ^3H , ^{11}C and ^{18}F .

30

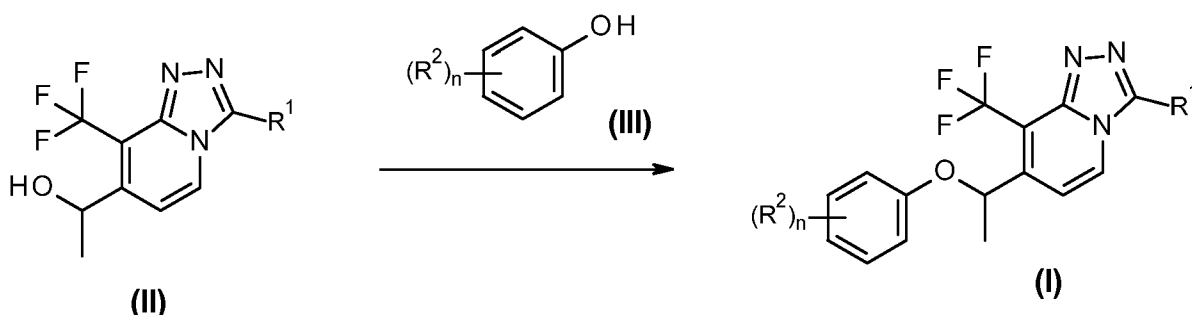
Preparation

The compounds according to the invention can generally be prepared by a succession of steps, each of which is known to the skilled person. In particular, the compounds can be prepared according to the following synthesis methods.

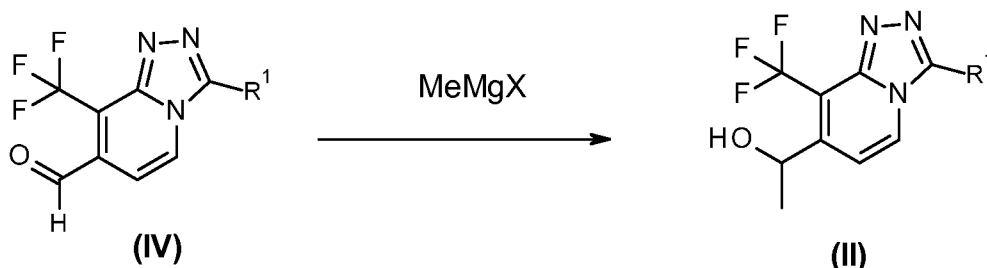
- 5 The compounds of Formula (I) may be synthesized in the form of racemic mixtures of enantiomers which can be separated from one another following art-known resolution procedures. The racemic compounds of Formula (I) may be converted into the corresponding diastereomeric salt forms by reaction with a suitable chiral acid. Said diastereomeric salt forms are subsequently separated, for example, by selective or
- 10 fractional crystallization and the enantiomers are liberated therefrom by alkali. An alternative manner of separating the enantiomeric forms of the compounds of Formula (I) involves liquid chromatography or supercritical fluid chromatography (SFC) using a chiral stationary phase. Said pure stereochemically isomeric forms may also be derived from the corresponding pure stereochemically isomeric forms of the appropriate
- 15 starting materials, provided that the reaction occurs stereospecifically.

A. Preparation of the final compounds

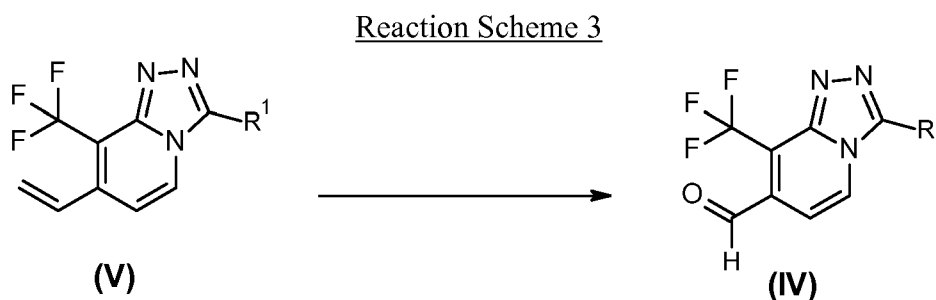
- Final compounds according to Formula (I), can be prepared by reacting an intermediate compound of Formula (II) with a compound of Formula (III) according to
- 20 reaction scheme (1), a reaction that is performed under classical Mitsunobu conditions. The reaction is preferably conducted with a phosphine and an azodicarboxylic ester or amide in tetrahydrofuran, 1,4-dioxane, diethyl ether, toluene, benzene, dichloromethane, or mixtures thereof, at -30 to 150°C, under thermal heating or microwave irradiation. Phosphines often used are triphenylphosphine and
- 25 tributylphosphine which are usually combined with dimethyl azodicarboxylate, diethyl azodicarboxylate, diisopropyl azodicarboxylate, di-(4-chlorobenzyl) azodicarboxylate, dibenzyl azodicarboxylate, di-tert-butyl azodicarboxylate, azodicarboxylic acid bis-(dimethylamide), azodicarboxylic acid dipiperidide, or azodicarboxylic acid dimorpholide. In reaction scheme (1), all variables are as defined in Formula (I)

Reaction Scheme 1B. Preparation of the intermediates5 Experimental procedure 2

Intermediate compounds according to Formula (II) can be prepared by
 10 subjecting an intermediate of Formula (IV) to conditions that are known to those skilled
 in the art. This is illustrated in reaction scheme (2) wherein all variables are defined as
 mentioned hereabove. Methods accomplishing these transformations are well known to
 those skilled in the art. Treatment of the aldehyde of formula (IV) with an
 organometallic such as methyl lithium or methyl magnesium bromide gives a
 compound of formula (II). A suitable solvent for this reaction is an ether such as
 tetrahydrofuran and the reaction is usually carried out at a temperature between -78°C
 and 40°C. In reaction scheme (2), all variables are defined as in Formula (I).

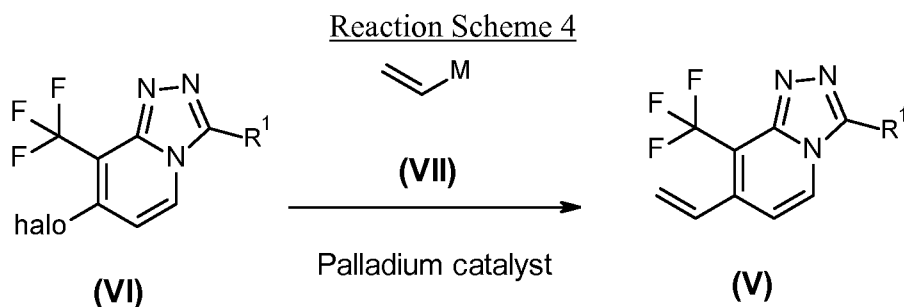
Reaction Scheme 2Experimental procedure 3

Intermediate compounds according to Formula (IV) can be prepared by reacting
 20 an intermediate of Formula (V) under dihydroxylation and oxidative cleavage
 conditions that are known to those skilled in the art and can be realized for example
 with oxone, osmium tetroxide. The process may be carried out optionally in a solvent
 such as 1,4-dioxane, water and generally at temperatures between about -100°C and
 about 100°C. A summary of such methods is found in "Comprehensive Organic
 25 Transformations", VCH Publishers, (1989), R.C.Larock, pp.595-596. This is illustrated
 in reaction scheme (3) wherein all variables are defined as mentioned hereabove.



Experimental procedure 4

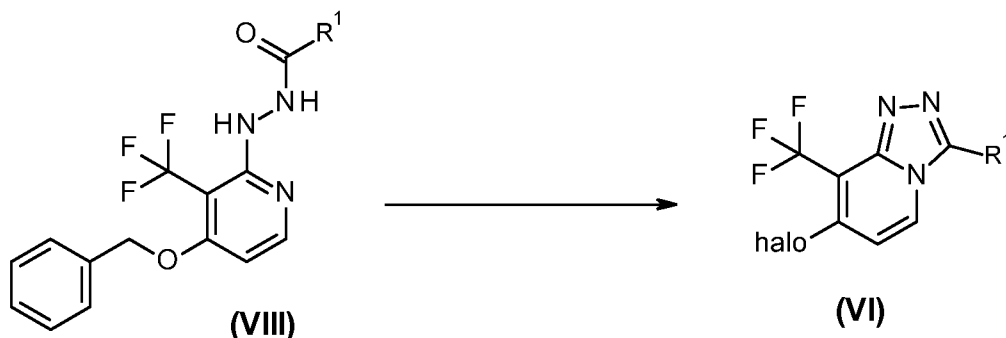
- 5 Intermediate compounds according to Formula (V) can be prepared by coupling reactions, such as Stille or Suzuki reactions of an intermediate of Formula (VI) with a compound of Formula (VII) under conditions that are known to those skilled in the art. The process may be carried out optionally in a solvent such as 1,4-dioxane, water and generally at temperatures between about r.t and about 200°C in the presence of a base.
- 10 This is illustrated in reaction scheme (4) wherein all variables are defined as mentioned hereabove, wherein M is trialkyltin, boronic acid or boronate ester, and a palladium catalyst and halo is chloro, bromo or iodo.



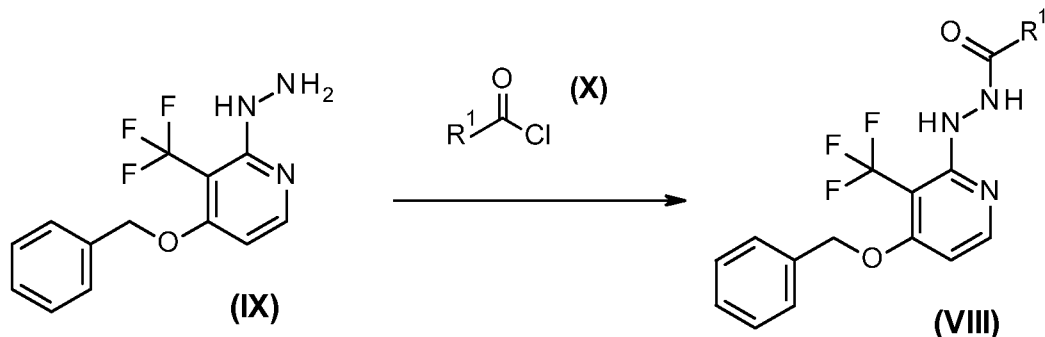
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Experimental procedure 5

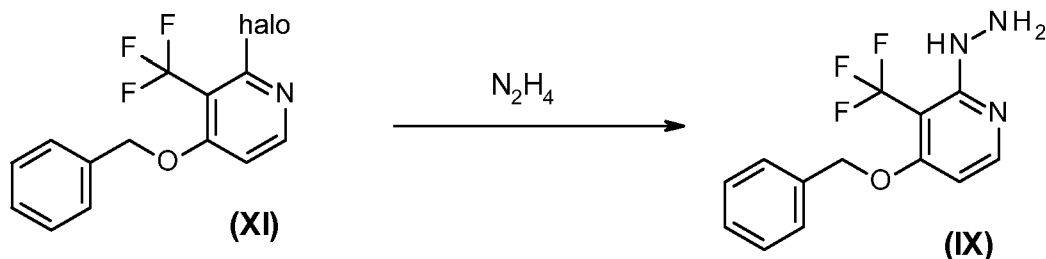
- Intermediate compounds according to Formula (VI) can be prepared following art known procedures by cyclization of an intermediate compound of Formula (VIII) in the presence of a halogenating agent such as for example phosphorus (V) oxychloride (POCl₃) in a suitable solvent such as, for example, dichloroethane, stirred under
- 20 microwave irradiation, for a suitable period of time that allows the completion of the reaction, as for example 5 min at a temperature between 140-200 °C. In reaction scheme (5), R¹ is defined as in Formula (I) and halo is chloro, bromo or iodo.

Reaction Scheme 5Experimental procedure 6

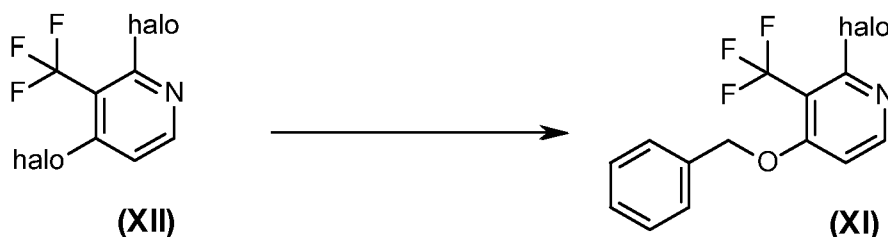
- 5 Intermediate compounds according to Formula (VIII) can be prepared by art known procedures by reaction of a hydrazine intermediate of Formula (IX) with acid halides of Formula (X). The reaction can be carried out using an inert-solvent, such as for example DCM, in the presence of a base such as for example triethylamine, for example at r.t. for a suitable period of time that allows completion of the reaction, for
- 10 example 20 min. In reaction scheme (6), R¹ is defined as in Formula (I).

Reaction Scheme 6Experimental procedure 7

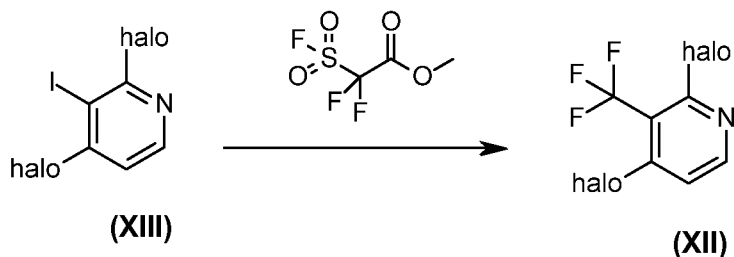
- 15 Intermediate compounds according to Formula (IX) can be prepared by reacting an intermediate compound of Formula (XI) with hydrazine according to reaction scheme (7), a reaction that is performed in a suitable reaction-inert solvent, such as, for example, ethanol, THF or 1,4-dioxane under thermal conditions such as, for example, heating the reaction mixture for example at 160 °C under microwave irradiation for 30
- 20 min or classical thermal heating at 70°C for 16 h. In reaction scheme (7), halo is chloro, bromo or iodo.

Reaction Scheme 7Experimental procedure 8

- 5 Intermediate compounds according to Formula (XI) can be prepared by reacting an intermediate compound of Formula (XII) with benzyl alcohol according to reaction scheme (8), a reaction that is performed in a suitable reaction-inert solvent, such as, for example, *N,N*-dimethylformamide in the presence of a suitable base, such as for example sodium hydride at r.t. for a suitable period of time that allows the completion
- 10 of the reaction, such as for example 1 h. In reaction scheme (8), halo is chloro, bromo or iodo.

Reaction Scheme 815 Experimental procedure 9

- Intermediate compounds of Formula (XII), can be prepared by reacting an intermediate of Formula (XIII), with a suitable trifluoromethylating agent, such as for example fluorosulfonyl(difluoro)acetic acid methyl ester, according to reaction scheme (9). This reaction is performed in a suitable reaction-inert solvent such as, for example,
- 20 *N,N*-dimethylformamide in the presence of a suitable coupling agent such as for example, copper(I) iodide, under thermal conditions such as, for example, heating the reaction mixture for example at 160 °C under microwave irradiation for 45 min. In reaction scheme (9), halo is chloro, bromo or iodo.

Reaction Scheme 9

The starting materials according to Formulae (II), (VII) (X) or (XIII) are compounds
5 that are either commercially available or may be prepared according to conventional
reaction procedures generally known to those skilled in the art.

Pharmacology

The compounds provided in this invention are positive allosteric modulators
10 (PAMs) of metabotropic glutamate receptors, in particular they are positive allosteric
modulators of mGluR2. The compounds of the present invention do not appear to bind
to the glutamate recognition site, the orthosteric ligand site, but instead to an allosteric
site within the seven transmembrane region of the receptor. In the presence of
glutamate or an agonist of mGluR2, the compounds of this invention increase the
15 mGluR2 response. The compounds provided in this invention are expected to have
their effect at mGluR2 by virtue of their ability to increase the response of such
receptors to glutamate or mGluR2 agonists.

As used herein, the term “treatment” is intended to refer to all processes,
wherein there may be a slowing, interrupting, arresting or stopping of the progression
20 of a disease or an alleviation of symptoms, but does not necessarily indicate a total
elimination of all symptoms.

Hence, the present invention relates to a compound according to the general
Formula (I), the stereoisomeric forms and the tautomers thereof and the
pharmaceutically acceptable acid or base addition salts and the solvates thereof, for use
25 as a medicament.

The invention also relates to the use of a compound according to the general
Formula (I), the stereoisomeric forms and the tautomers thereof and the
pharmaceutically acceptable acid or base addition salts and the solvates thereof, or a
pharmaceutical composition according to the invention for the manufacture of a
30 medicament.

The invention also relates to a compound according to the general Formula (I),
the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable

acid or base addition salts and the solvates thereof, or a pharmaceutical composition according to the invention for use in the treatment or prevention of, in particular treatment of, a condition in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of allosteric modulators of mGluR2, in particular positive allosteric modulators thereof.

The present invention also relates to the use of a compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, or a pharmaceutical composition according to the invention for the manufacture of a medicament for the treatment or prevention of, in particular treatment of, a condition in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of allosteric modulators of mGluR2, in particular positive allosteric modulators thereof.

The present invention also relates to a compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, or a pharmaceutical composition according to the invention for use in the treatment, prevention, amelioration, control or reduction of the risk of various neurological and psychiatric disorders associated with glutamate dysfunction in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of positive allosteric modulators of mGluR2.

Also, the present invention relates to the use of a compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, or a pharmaceutical composition according to the invention for the manufacture of a medicament for treating, preventing, ameliorating, controlling or reducing the risk of various neurological and psychiatric disorders associated with glutamate dysfunction in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of positive allosteric modulators of mGluR2.

In particular, the neurological and psychiatric disorders associated with glutamate dysfunction, include one or more of the following conditions or diseases: acute neurological and psychiatric disorders such as, for example, cerebral deficits subsequent to cardiac bypass surgery and grafting, stroke, cerebral ischemia, spinal cord trauma, head trauma, perinatal hypoxia, cardiac arrest, hypoglycemic neuronal damage, dementia (including AIDS-induced dementia), Alzheimer's disease,

Huntington's Chorea, amyotrophic lateral sclerosis, ocular damage, retinopathy, cognitive disorders, idiopathic and drug-induced Parkinson's disease, muscular spasms and disorders associated with muscular spasticity including tremors, epilepsy, convulsions, migraine (including migraine headache), urinary incontinence, substance
5 dependence/abuse, substance withdrawal (including substances such as, for example, opiates, nicotine, tobacco products, alcohol, benzodiazepines, cocaine, sedatives, hypnotics, etc.), psychosis, schizophrenia, anxiety (including generalized anxiety disorder, panic disorder, and obsessive compulsive disorder), mood disorders (including depression, major depressive disorder, treatment resistant depression, mania,
10 bipolar disorders, such as bipolar mania), posttraumatic stress disorder, trigeminal neuralgia, hearing loss, tinnitus, macular degeneration of the eye, emesis, brain edema, pain (including acute and chronic states, severe pain, intractable pain, neuropathic pain, and post-traumatic pain), tardive dyskinesia, sleep disorders (including narcolepsy), attention deficit/hyperactivity disorder, and conduct disorder.

15 In particular, the condition or disease is a central nervous system disorder selected from the group of anxiety disorders, psychotic disorders, personality disorders, substance-related disorders, eating disorders, mood disorders, migraine, epilepsy or convulsive disorders, childhood disorders, cognitive disorders, neurodegeneration, autistic disorders, neurotoxicity and ischemia.

20 In particular, the central nervous system disorder is an anxiety disorder, selected from the group of agoraphobia, generalized anxiety disorder (GAD), mixed anxiety and depression, obsessive-compulsive disorder (OCD), panic disorder, posttraumatic stress disorder (PTSD), social phobia and other phobias. An additional anxiety disorder is panic attack.

25 In particular, the central nervous system disorder is a psychotic disorder selected from the group of schizophrenia, delusional disorder, schizoaffective disorder, schizophreniform disorder and substance-induced psychotic disorder; more in particular, negative symptoms or residual symptoms of schizophrenia. Such disorders manifest psychosis as a prominent symptom. Therefore, the invention also relates to a
30 compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, or a pharmaceutical composition according to the invention for use in the treatment, prevention, amelioration, control or reduction of psychosis.

In particular, the central nervous system disorder is a personality disorder selected from the group of obsessive-compulsive personality disorder, borderline personality disorder and schizoid, schizotypal disorder.

5 In particular, the central nervous system disorder is a substance abuse or substance-related disorder selected from the group of alcohol abuse, alcohol addiction, alcohol dependence, alcohol withdrawal, alcohol withdrawal delirium, alcohol-induced psychotic disorder, amphetamine addiction, amphetamine dependence, amphetamine withdrawal, cocaine addiction, cocaine dependence, cocaine withdrawal, nicotine addiction, nicotine dependence, nicotine withdrawal, opioid dependence and opioid
10 withdrawal. The treatment or prevention of the substance abuse or substance-related disorders referred to herein may involve relapse prevention thereof.

In particular, the central nervous system disorder is an eating disorder selected from the group of anorexia nervosa and bulimia nervosa.

15 In particular, the central nervous system disorder is a mood disorder selected from the group of bipolar disorders (I & II), cyclothymic disorder, depression, dysthymic disorder, major depressive disorder, treatment resistant depression, bipolar depression, and substance-induced mood disorder.

In particular, the central nervous system disorder is migraine.

20 In particular, the central nervous system disorder is epilepsy or a convulsive disorder selected from the group of generalized nonconvulsive epilepsy, generalized convulsive epilepsy, petit mal status epilepticus, grand mal status epilepticus, partial epilepsy with or without impairment of consciousness, infantile spasms, epilepsy partialis continua, and other forms of epilepsy. Additional disorders encompassed under epilepsy or convulsive disorder include any disorder in which a subject
25 (preferably a human adult, child or infant) experiences one or more seizures and / or tremors. Suitable examples include, but are not limited to, epilepsy (including, but not limited to, localization-related epilepsies, generalized epilepsies, epilepsies with both generalized and local seizures, and the like), partial-onset seizures with or without generalization, myoclonic seizures, primary generalized tonic-clonic seizures in
30 particular in patients with idiopathic generalized epilepsy, seizures associated with Lennox-Gastaut syndrome, seizures as a complication of a disease or condition (such as seizures associated with encephalopathy, phenylketonuria, juvenile Gaucher's disease, Lundborg's progressive myoclonic epilepsy, stroke, head trauma, stress, hormonal changes, drug use or withdrawal, alcohol use or withdrawal, sleep deprivation, fever,
35 infection, and the like), status epilepticus (convulsive or non convulsive), essential

tremor, restless limb syndrome, and the like. Preferably, the disorder is selected from epilepsy (regardless of type, underlying cause or origin), essential tremor or restless limb syndrome. More preferably, the disorder is epilepsy (regardless of type, underlying cause or origin) or essential tremor. In particular, the disorder is epilepsy
5 (regardless of type, underlying cause or origin). A more particular example of epilepsy is refractory epilepsy, also referred to as treatment or therapy resistant epilepsy. This term is often used when patients have failed three or more anti-epileptic drugs (AEDs). Refractory epilepsy also includes refractory partial epilepsy and refractory generalized epilepsy (including idiopathic or symptomatic).

10 In particular, the central nervous system disorder is attention-deficit/hyperactivity disorder.

In particular, the central nervous system disorder is an autistic disorder selected from autism and autism spectrum disorders, such as Asperger's syndrome.

In particular, the central nervous system disorder is a cognitive disorder selected
15 from the group of delirium, substance-induced persisting delirium, dementia, dementia due to HIV disease, dementia due to Huntington's disease, dementia due to Parkinson's disease, dementia of the Alzheimer's type, behavioral and psychological symptoms of dementia, substance-induced persisting dementia and mild cognitive impairment.

Particular examples of behavioral and psychological symptoms of dementia (BPSD)
20 include, but are not limited to, aggression, agitation and psychosis.

In particular, the central nervous system disorder is selected from the group of schizophrenia, behavioral and psychological symptoms of dementia, major depressive disorder, treatment resistant depression, bipolar depression, anxiety, depression,
25 generalised anxiety disorder, post-traumatic stress disorder, bipolar mania, epilepsy, attention-deficit/hyperactivity disorder, substance abuse and mixed anxiety and depression.

In particular, the central nervous system disorder is selected from the group of schizophrenia, epilepsy, obsessive compulsive disorder, alcohol addiction, cocaine addiction, nicotine addiction, borderline personality disorder, bipolar disorder,
30 behavioral and psychological symptoms of dementia, Asperger's syndrome, major depressive disorder, treatment resistant depression, anxiety, depression, generalised anxiety disorder, and mixed anxiety and depression.

In particular, the central nervous system disorder is selected from the group of schizophrenia (in particular, negative symptoms or residual symptoms thereof),

generalized anxiety disorder, bipolar disorder (I or II), migraine, behavioral and psychological symptoms of dementia, epilepsy or convulsive disorders, panic disorder, mixed anxiety and depression, and agoraphobia.

In particular, the central nervous system disorder is selected from the group of
5 schizophrenia (in particular, negative symptoms or residual symptoms thereof),
generalized anxiety disorder, bipolar disorder (I or II), migraine, epilepsy, panic
disorder, mixed anxiety and depression, and agoraphobia. Of the disorders mentioned
above, the treatment of psychosis, schizophrenia, behavioral and psychological
symptoms of dementia, major depressive disorder, treatment resistant depression,
10 bipolar depression, anxiety, depression, generalised anxiety disorder, post-traumatic
stress disorder, bipolar mania, substance abuse and mixed anxiety and depression, are
of particular importance.

Of the disorders mentioned above, the treatment of generalized anxiety disorder,
bipolar disorder (I or II), epilepsy, panic disorder, and agoraphobia are of particular
15 importance.

Of the disorders mentioned above, the treatment of anxiety, schizophrenia,
migraine, depression, and epilepsy are of particular importance.

Of the disorders mentioned above, the treatment of anxiety and epilepsy are of
particular importance.

20 At present, the fourth edition of the Diagnostic & Statistical Manual of Mental
Disorders (DSM-IV) of the American Psychiatric Association provides a diagnostic
tool for the identification of the disorders described herein. The person skilled in the art
will recognize that alternative nomenclatures, nosologies, and classification systems for
neurological and psychiatric disorders described herein exist, and that these evolve with
25 medical and scientific progresses.

A skilled person will be familiar with alternative nomenclatures, nosologies,
and classification systems for the diseases or conditions referred to herein. For
example, the “American Psychiatric Association: Diagnostic and Statistical Manual of
Mental Disorders, Fifth Edition. Arlington, VA, American Psychiatric Association,
30 2013” (DSM-5TM) utilizes terms such as anxiety disorders, in particular, agoraphobia,
generalized anxiety disorder, panic disorder, social anxiety disorder (social phobia),
and panic attack; schizophrenia spectrum and other psychotic disorders, in particular,
schizophrenia, delusional disorder, schizoaffective disorder, schizophreniform disorder,
substance/medication-induced and psychotic disorder; personality disorders, in
35 particular, obsessive-compulsive personality disorder, borderline personality disorder,

schizoid personality disorder, and schizotypal personality disorder; substance-related and addictive disorders, in particular, alcohol use disorder, alcohol withdrawal, opioid use disorder, opioid withdrawal, stimulant (amphetamine-type substance, cocaine) use disorder, stimulant (amphetamine-type substance, cocaine) withdrawal, tobacco use disorder, and tobacco withdrawal; depressive disorders, in particular, major depressive disorder, persistent depressive disorder (dysthymia), and substance/medication-induced depressive disorder; bipolar and related disorders, in particular, bipolar I disorder, bipolar II disorder, cyclothymic disorder, substance/medication-induced bipolar and related disorder; obsessive-compulsive disorder and related disorders, in particular, obsessive-compulsive disorder; trauma- and stressor-related disorders, in particular, posttraumatic stress disorder, and acute stress disorder; neurodevelopmental disorder, in particular, autism spectrum disorder, and attention-deficit/hyperactivity disorder; neurocognitive disorders (NCDs) (both major and mild), in particular, delirium, substance intoxication delirium, NCD due to Alzheimer's disease, NCD due to HIV infection, NCD due to Huntington's disease, NCD due to Parkinson's disease, and substance/medication-induced NCD. Such terms may be used by the skilled person as an alternative nomenclature for some of the diseases or conditions referred to herein.

Therefore, the invention also relates to a compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, for use in the treatment of any one of the diseases mentioned hereinbefore.

The invention also relates to a compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, for use in treating any one of the diseases mentioned hereinbefore.

The invention also relates to a compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, for the treatment or prevention, in particular treatment, of any one of the diseases mentioned hereinbefore.

The invention also relates to the use of a compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, for the manufacture of a medicament for the treatment or prevention of any one of the disease conditions mentioned hereinbefore.

The invention also relates to the use of a compound according to the general Formula (I), the stereoisomeric forms and the tautomers thereof and the pharmaceutically acceptable acid or base addition salts and the solvates thereof, for the manufacture of a medicament for the treatment of any one of the disease conditions mentioned hereinbefore.

The compounds of the present invention can be administered to mammals, preferably humans, for the treatment or prevention of any one of the diseases mentioned hereinbefore.

In view of the utility of the compounds of Formula (I), there is provided a method of treating warm-blooded animals, including humans, suffering from any one of the diseases mentioned hereinbefore, and a method of preventing in warm-blooded animals, including humans, any one of the diseases mentioned hereinbefore.

Said methods comprise the administration, i.e. the systemic or topical administration, preferably oral administration, of a therapeutically effective amount of a compound of Formula (I), a stereoisomeric form or a tautomer thereof and a pharmaceutically acceptable addition salt or solvate thereof, to warm-blooded animals, including humans.

Therefore, the invention also relates to a method for the prevention and/or treatment of any one of the diseases mentioned hereinbefore comprising administering a therapeutically effective amount of a compound according to the invention to a patient in need thereof.

One skilled in the art will recognize that a therapeutically effective amount of the PAMs of the present invention is the amount sufficient to modulate the activity of the mGluR2 and that this amount varies *inter alia*, depending on the type of disease, the concentration of the compound in the therapeutic formulation, and the condition of the patient. Generally, an amount of PAM to be administered as a therapeutic agent for treating diseases in which modulation of the mGluR2 is beneficial, such as the disorders described herein, will be determined on a case by case by an attending physician.

Generally, a suitable dose is one that results in a concentration of the PAM at the treatment site in the range of 0.5 nM to 200 μ M, and more usually 5 nM to 50 μ M. To obtain these treatment concentrations, a patient in need of treatment likely will be administered an effective therapeutic daily amount of about 0.01 mg/kg to about 50 mg/kg body weight, preferably from about 0.01 mg/kg to about 25 mg/kg body weight, more preferably from about 0.01 mg/kg to about 10 mg/kg body weight, more

preferably from about 0.01 mg/kg to about 2.5 mg/kg body weight, even more preferably from about 0.05 mg/kg to about 1 mg/kg body weight, more preferably from about 0.1 to about 0.5 mg/kg body weight. The amount of a compound according to the present invention, also referred to here as the active ingredient, which is required to achieve a therapeutically effect will, of course vary on case-by-case basis, vary with the particular compound, the route of administration, the age and condition of the recipient, and the particular disorder or disease being treated. A method of treatment may also include administering the active ingredient on a regimen of between one and four intakes per day. In these methods of treatment the compounds according to the invention are preferably formulated prior to admission. As described herein below, suitable pharmaceutical formulations are prepared by known procedures using well known and readily available ingredients.

Because such positive allosteric modulators of mGluR2, including compounds of Formula (I), enhance the response of mGluR2 to glutamate, it is an advantage that the present methods utilize endogenous glutamate.

Because positive allosteric modulators of mGluR2, including compounds of Formula (I), enhance the response of mGluR2 to agonists, it is understood that the present invention extends to the treatment of neurological and psychiatric disorders associated with glutamate dysfunction by administering an effective amount of a positive allosteric modulator of mGluR2, including compounds of Formula (I), in combination with an mGluR2 agonist (or mGluR2/3 agonist). Examples of mGluR2/mGluR2/3 agonists include, for example, LY-379268; DCG-IV; LY-354740; LY-404039; LY-544344; LY-2140023; LY-181837; LY-389795; LY-446433; LY-450477; talaglumetad; MGS0028; MGS0039; (-)-2-oxa-4-aminobicyclo[3.1.0]hexane-4,6-dicarboxylate; (+)-4-amino-2-sulfonylbicyclo[3.1.0]hexane-4,6-dicarboxylic acid; (+)-2-amino-4-fluorobicyclo[3.1.0]hexane-2,6-dicarboxylic acid; 1S,2R,5S,6S-2-amino-6-fluoro-4-oxobicyclo[3.1.0]hexane-2,6-dicarboxylic acid; 1S,2R,4S,5S,6S-2-amino-6-fluoro-4-hydroxybicyclo[3.1.0]hexane-2,6-dicarboxylic acid; 1S,2R,3R,5S,6S-2-amino-3-fluorobicyclo[3.1.0]hexane-2,6-dicarboxylic acid; 1S,2R,3S,5S,6S-2-amino-6-fluoro-3-hydroxybicyclo[3.1.0]hexane-2,6-dicarboxylic acid; (+)-4-amino-2-sulfonylbicyclo[3.1.0]hexane-4,6-dicarboxylic acid; (+)-2-amino-4-fluorobicyclo[3.1.0]hexane-2,6-dicarboxylic acid; 1S,2R,5S,6S-2-amino-6-fluoro-4-oxobicyclo[3.1.0]hexane-2,6-dicarboxylic acid; 1S,2R,4S,5S,6S-2-amino-6-fluoro-4-hydroxybicyclo[3.1.0]hexane-2,6-dicarboxylic acid; 1S,2R,3R,5S,6S-2-amino-3-fluorobicyclo[3.1.0]hexane-2,6-dicarboxylic acid; or 1S,2R,3S,5S,6S-2-amino-6-fluoro-3-hydroxybicyclo-

[3.1.0]hexane-2,6-dicarboxylic acid. More preferable mGluR2 agonists include LY-379268; DCG-IV; LY-354740; LY-404039; LY-544344; or LY-2140023.

The compounds of the present invention may be utilized in combination with one or more other drugs in the treatment, prevention, control, amelioration, or reduction of risk of diseases or conditions for which compounds of Formula (I) or the other drugs may have utility, where the combination of the drugs together are safer or more effective than either drug alone.

Pharmaceutical compositions

The present invention also provides compositions for preventing or treating diseases in which modulation of the mGlu2 receptor is beneficial, such as the disorders described herein. While it is possible for the active ingredient to be administered alone, it is preferable to present it as a pharmaceutical composition. Accordingly, the present invention also relates to a pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and, as active ingredient, a therapeutically effective amount of a compound according to the invention, in particular a compound according to Formula (I), a pharmaceutically acceptable salt thereof, a solvate thereof or a stereochemically isomeric form or a tautomer thereof. The carrier or diluent must be “acceptable” in the sense of being compatible with the other ingredients of the composition and not deleterious to the recipients thereof.

The compounds according to the invention, in particular the compounds according to Formula (I), the pharmaceutically acceptable salts thereof, the solvates and the stereochemically isomeric forms and the tautomers thereof, or any subgroup or combination thereof may be formulated into various pharmaceutical forms for administration purposes. As appropriate compositions there may be cited all compositions usually employed for systemically administering drugs.

The pharmaceutical compositions of this invention may be prepared by any methods well known in the art of pharmacy, for example, using methods such as those described in Gennaro et al. Remington's Pharmaceutical Sciences (18th ed., Mack Publishing Company, 1990, see especially Part 8: Pharmaceutical preparations and their Manufacture). To prepare the pharmaceutical compositions of this invention, a therapeutically effective amount of the particular compound, optionally in salt form, as the active ingredient is combined in intimate admixture with a pharmaceutically acceptable carrier or diluent, which carrier or diluent may take a wide variety of forms depending on the form of preparation desired for administration. These pharmaceutical

compositions are desirable in unitary dosage form suitable, in particular, for oral, topical, rectal or percutaneous administration, by parenteral injection or by inhalation. For example, in preparing the compositions in oral dosage form, any of the usual pharmaceutical media may be employed such as, for example, water, glycols, oils, alcohols and the like in the case of oral liquid preparations such as, for example, suspensions, syrups, elixirs, emulsions and solutions; or solid carriers such as, for example, starches, sugars, kaolin, diluents, lubricants, binders, disintegrating agents and the like in the case of powders, pills, capsules and tablets. Because of the ease in administration, oral administration is preferred, and tablets and capsules represent the most advantageous oral dosage unit forms in which case solid pharmaceutical carriers are obviously employed. For parenteral compositions, the carrier will usually comprise sterile water, at least in large part, though other ingredients, for example, surfactants, to aid solubility, may be included. Injectable solutions, for example, may be prepared in which the carrier comprises saline solution, glucose solution or a mixture of saline and glucose solution. Injectable suspensions may also be prepared in which case appropriate liquid carriers, suspending agents and the like may be employed. Also included are solid form preparations that are intended to be converted, shortly before use, to liquid form preparations. In the compositions suitable for percutaneous administration, the carrier optionally comprises a penetration enhancing agent and/or a suitable wetting agent, optionally combined with suitable additives of any nature in minor proportions, which additives do not introduce a significant deleterious effect on the skin. Said additives may facilitate the administration to the skin and/or may be helpful for preparing the desired compositions. These compositions may be administered in various ways, e.g., as a transdermal patch, as a spot-on, as an ointment.

It is especially advantageous to formulate the aforementioned pharmaceutical compositions in unit dosage form for ease of administration and uniformity of dosage. Unit dosage form as used herein refers to physically discrete units suitable as unitary dosages, each unit containing a predetermined quantity of active ingredient calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. Examples of such unit dosage forms are tablets (including scored or coated tablets), capsules, pills, powder packets, wafers, suppositories, injectable solutions or suspensions and the like, teaspoonfuls, tablespoonfuls, and segregated multiples thereof.

Since the compounds according to the invention are orally administrable compounds, pharmaceutical compositions comprising said compounds for oral administration are especially advantageous.

In order to enhance the solubility and/or the stability of the compounds of Formula (I) in pharmaceutical compositions, it can be advantageous to employ α -, β - or γ -cyclodextrins or their derivatives, in particular hydroxyalkyl substituted cyclodextrins, e.g. 2-hydroxypropyl- β -cyclodextrin or sulfobutyl- β -cyclodextrin. Also
5 co-solvents such as alcohols may improve the solubility and/or the stability of the compounds according to the invention in pharmaceutical compositions.

The exact dosage and frequency of administration depends on the particular compound of formula (I) used, the particular condition being treated, the severity of the condition being treated, the age, weight, sex, extent of disorder and general physical
10 condition of the particular patient as well as other medication the individual may be taking, as is well known to those skilled in the art. Furthermore, it is evident that said effective daily amount may be lowered or increased depending on the response of the treated subject and/or depending on the evaluation of the physician prescribing the compounds of the instant invention.

Depending on the mode of administration, the pharmaceutical composition will
15 comprise from 0.05 to 99 % by weight, preferably from 0.1 to 70 % by weight, more preferably from 0.1 to 50 % by weight of the active ingredient, and, from 1 to 99.95 % by weight, preferably from 30 to 99.9 % by weight, more preferably from 50 to 99.9 % by weight of a pharmaceutically acceptable carrier, all percentages being based on the
20 total weight of the composition.

The amount of a compound of Formula (I) that can be combined with a carrier material to produce a single dosage form will vary depending upon the disease treated, the mammalian species, and the particular mode of administration. However, as a
25 general guide, suitable unit doses for the compounds of the present invention can, for example, preferably contain between 0.1 mg to about 1000 mg of the active compound. A preferred unit dose is between 1 mg to about 500 mg. A more preferred unit dose is between 1 mg to about 300 mg. Even more preferred unit dose is between 1 mg to about 100 mg. Such unit doses can be administered more than once a day, for example, 2, 3, 4, 5 or 6 times a day, but preferably 1 or 2 times per day, so that the total dosage
30 for a 70 kg adult is in the range of 0.001 to about 15 mg per kg weight of subject per administration. A preferred dosage is 0.01 to about 1.5 mg per kg weight of subject per administration, and such therapy can extend for a number of weeks or months, and in some cases, years. It will be understood, however, that the specific dose level for any particular patient will depend on a variety of factors including the activity of the
35 specific compound employed; the age, body weight, general health, sex and diet of the individual being treated; the time and route of administration; the rate of excretion;

other drugs that have previously been administered; and the severity of the particular disease undergoing therapy, as is well understood by those of skill in the area.

A typical dosage can be one 1 mg to about 100 mg tablet or 1 mg to about 300 mg taken once a day, or, multiple times per day, or one time-release capsule or tablet
5 taken once a day and containing a proportionally higher content of active ingredient. The time-release effect can be obtained by capsule materials that dissolve at different pH values, by capsules that release slowly by osmotic pressure, or by any other known means of controlled release.

It can be necessary to use dosages outside these ranges in some cases as will be
10 apparent to those skilled in the art. Further, it is noted that the clinician or treating physician will know how and when to start, interrupt, adjust, or terminate therapy in conjunction with individual patient response.

As already mentioned, the invention also relates to a pharmaceutical composition comprising the compounds according to the invention and one or more
15 other drugs for use as a medicament or for use in the treatment, prevention, control, amelioration, or reduction of risk of diseases or conditions for which compounds of Formula (I) or the other drugs may have utility. The use of such a composition for the manufacture of a medicament as well as the use of such a composition for the manufacture of a medicament in the treatment, prevention, control, amelioration or
20 reduction of risk of diseases or conditions for which compounds of Formula (I) or the other drugs may have utility are also contemplated. The present invention also relates to a combination of a compound according to the present invention and an mGluR2 orthosteric agonist (or a mGluR2/3 orthosteric agonist). The present invention also relates to such a combination for use as a medicine. The present invention also relates
25 to a product comprising (a) a compound according to the present invention, a pharmaceutically acceptable salt thereof or a solvate thereof, and (b) a mGluR2 orthosteric agonist (or a mGluR2/3 orthosteric agonist), as a combined preparation for simultaneous, separate or sequential use in the treatment or prevention of a condition in a mammal, including a human, the treatment or prevention of which is affected or
30 facilitated by the neuromodulatory effect of mGluR2 allosteric modulators, in particular positive mGluR2 allosteric modulators. The different drugs of such a combination or product may be combined in a single preparation together with pharmaceutically acceptable carriers or diluents, or they may each be present in a separate preparation together with pharmaceutically acceptable carriers or diluents.

35 The following examples are intended to illustrate but not to limit the scope of the

present invention.

Chemistry

Several methods for preparing the compounds of this invention are illustrated in the following Examples. Unless otherwise noted, all starting materials were obtained from commercial suppliers and used without further purification.

Hereinafter, "aq." means aqueous; "DCE" means 1,2-dichloroethane, "DCM" means dichloromethane; "DIPE" means diisopropylether; "DIPEA" means *N,N*-diisopropylethylamine; "DMF" means *N,N*-dimethylformamide; "ES" means electrospray; "Et₃N" means triethylamine; "Et₂O" means diethyl ether; "EtOAc" means ethyl acetate; "h" means hours; "HPLC" means high performance liquid chromatography; "HRMS" means high-resolution mass spectra/spectrometry; "l" or "L" means liter; "LRMS" means low-resolution mass spectrometry/spectra; "MeOH" means methanol; "min" means minute(s); "mp" means melting point; "Pd(PPh₃)₄" means tetrakis(triphenylphosphine)palladium(0); "RP" means reverse phase; "r.t." means room temperature; "s" means seconds; "sat." means saturated; "SFC" means supercritical fluid chromatography; "sol." means solution; "THF" means tetrahydrofuran.

Microwave assisted reactions were performed in a single-mode reactor: Initiator™ Sixty EXP microwave reactor (Biotage AB), or in a multimode reactor: MicroSYNTH Labstation (Milestone, Inc.).

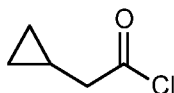
Thin layer chromatography (TLC) was carried out on silica gel 60 F254 plates (Merck) using reagent grade solvents. Open column chromatography was performed on silica gel, particle size 60 Å, mesh = 230-400 (Merck) using standard techniques. Automated flash column chromatography was performed using ready-to-connect cartridges from Merck, on irregular silica gel, particle size 15-40 µm (normal phase disposable flash columns) on a SPOT or LAFLASH system from Armen Instrument.

The absolute stereochemical configuration for some of the compounds was determined using vibrational circular dichroism (VCD). They were measured on a Bruker Equinox 55 equipped with a PMA 37, in a KBr liquid cell using CD₂Cl₂ as solvent (PEM: 1350cm⁻¹, LIA: 1mV, resolution: 4 cm⁻¹). A description on the use of VCD for the determination of absolute configuration can be found in Dyatkin A.B. et. al, Chirality, 14:215-219 (2002).

Whenever the notation “RS” is indicated herein, it denotes that the compound is a racemic mixture, unless otherwise indicated. The stereochemical configuration for some compounds has been designated “*R*” or “*S*” when the mixture was separated; for
5 some compounds, the stereochemical configuration has been designated as “**R*” or “**S*” when the absolute stereochemistry is undetermined although the compound itself has been isolated as a single stereoisomer and is enantiomerically pure. The enantiomeric excess of compounds reported herein was determined by analysis of the racemic mixture by supercritical fluid chromatography (SFC) followed by SFC
10 comparison of the separated enantiomer(s).

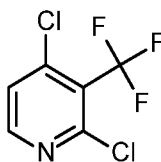
Preparation of intermediates

Description 1 - Intermediate 1

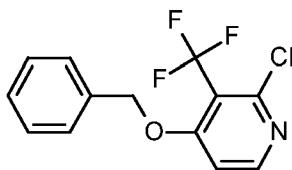


15 Cyclopropylacetic acid ([CAS 5239-82-7], 50 g, 500 mmol) was dissolved in CH₂Cl₂ (300 mL) then SOCl₂ (100 mL) was added. The reaction mixture was stirred at 60 °C for 2 h and then the solvent was evaporated to yield intermediate **1** (53 g, 90%), which was used without further purification.

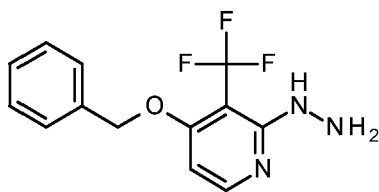
Description 2 - Intermediate 2



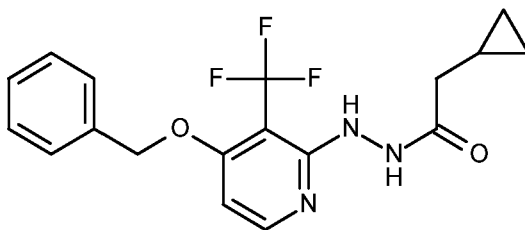
20 To a solution of 2,4-dichloro-3-iodopyridine ([CAS 343781-36-2], 290 g, 1058 mmol) in DMF (1.7 L) was added methyl 2,2-difluoro-2-(fluorosulfonyl)acetate ([CAS 680-15-9], 403 g, 2098 mmol) and CuI (403 g, 2.13 mol), the reaction was then heated at 100 °C for 5 h.
25 The reaction was cooled and filtered. The filtrate was diluted with H₂O and extracted with Et₂O and washed with a NH₃ solution. The organic layer was dried (Na₂SO₄), filtered and concentrated *in vacuo* to yield intermediate **2** (160 g), which was used without further purification.

Description 3 - Intermediate 3

To a solution of NaH (60% in oil, 24 g, 600 mmol) in DMF (2 L) at 0 °C was added benzyl alcohol (35 g, 325 mmol), then the reaction was stirred for 2 min. Intermediate **2** (160 mg, 741 mmol) was added in one portion, and stirred at 0 °C for 1 h. The reaction was diluted by the addition of H₂O and extracted with Et₂O. The organic layer was dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by column chromatography over silica gel (eluent: petroleum ether/EtOAc = 20/1). The pure fractions were collected and the solvent was evaporated to yield intermediate **3** (100 g, 38%).

Description 4 - Intermediate 4

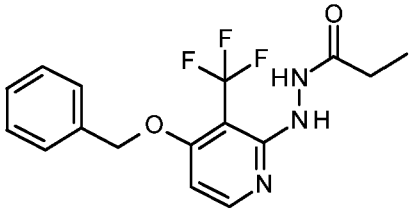
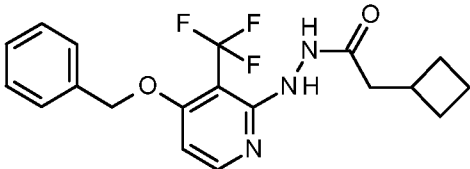
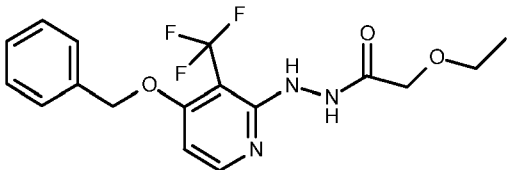
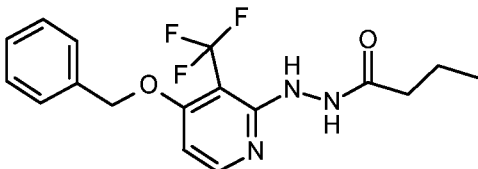
To a solution of intermediate **3** (100 g, 277 mmol) in 1,4-dioxane (1.5 L) was added NH₂NH₂ hydrate (85% solution in water, 300 g, 9.11 mol), the reaction was then heated in sealed tube at 160 °C for 2 h. The mixture was concentrated *in vacuo*, dissolved in DCM washed with NaHCO₃. The organic layer was dried (Na₂SO₄), filtered and concentrated *in vacuo* to yield intermediate **4** (90 g, 90%), which was used without further purification.

Description 5 - Intermediate 5

To a solution of intermediate **4** (90 g, 318 mmol) in CH₂Cl₂ (1.5 L) was added triethylamine (64.3 g, 636 mmol), the mixture was cooled to 0 °C, then a solution of intermediate **1** (53 g, 449 mmol) in CH₂Cl₂ was added. The solution was stirred at RT for 1 h. The reaction mixture was washed with a sat. aq. sol. of NaHCO₃, and extracted

with CH₂Cl₂. The organic layer was dried (Na₂SO₄), filtered and concentrated *in vacuo* to yield intermediate **5** (104.4 g, 90%).

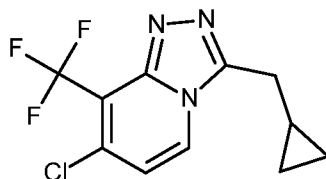
The following intermediates were synthesized following a synthetic sequence analogous to that reported in Description 5 (D5).

Intermediate	Acid chloride	Conditions
 Intermediate 6	propionyl chloride ([CAS 79-03-8])	Addition run at RT.
 Intermediate 7	cyclobutaneacetyl chloride ([CAS 59543-38-3])	Conditions as in D5.
 Intermediate 8	2-ethoxy-acetyl chloride ([CAS 14077-58-8])	Conditions as in D5.
 Intermediate 25	butyryl chloride ([CAS 141-75-3])	Conditions as in D5.

5

Description 6

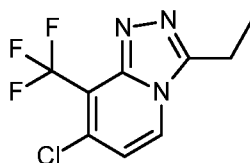
(a) Intermediate 9



To a solution of intermediate **5** (101 g, 277 mmol) in CH₃CN (1.2 L) were added
 10 phosphorus(V) oxychloride (84.7 g, 553 mmol) and *N,N*-diisopropylethylamine (71.3

- g, 553 mmol). The reaction mixture was stirred at 90 °C for 38 h. The reaction was then diluted with DCM and washed with a Na₂CO₃ solution. The organic layer was dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by column chromatography over silica gel (eluent: petroleum ether/EtOAc = 4/1). The pure fractions were collected and the solvent was evaporated to yield intermediate **9** (31.39 g, 41%).

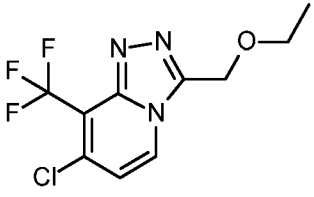
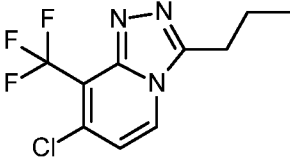
(b) Intermediate 10



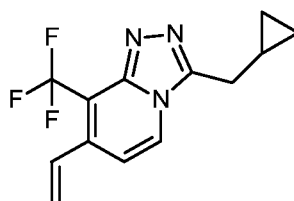
- The reaction was performed in 4 batches then combined for work up and purification.
- To a solution of intermediate **6** (7 g, 20.6 mmol) in DCE (50 mL), was added *N,N*-diisopropylethylamine (3.96 mL, 22.69 mmol) and then phosphorus oxychloride (2.12 mL, 22.69 mmol) and the reaction mixture was heated in a microwave at 150 °C for 5 min. Then DCM was added and the organic layer was washed with a sat. sol. of NaHCO₃, dried (Na₂SO₄) and concentrated *in vacuo* to afford the desired compound, which was purified by column chromatography (gradient elution: DCM 100% to MeOH.NH₃ 2% in DCM) to yield intermediate **10** (2.5 g, 49%).

The following intermediates were synthesized following a synthetic sequence analogous to that reported in Description 6(a) or (b).

Intermediate	Starting material	Conditions
 Intermediate 11	Intermediate 7	Reaction performed as in (a) but in CH ₃ CN. After the reaction was complete, the reaction mixture was poured into ice/water then washed with NaHCO ₃ sat. sol. And extracted with DCM, dried (Na ₂ SO ₄), filtered and concentrated. Purification was performed in Spot (Si cartridge, eluent DCM/EtOAc up to 10-20%).

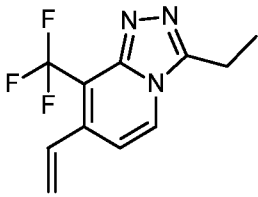
Intermediate	Starting material	Conditions
 Intermediate 12	Intermediate 8	Reaction performed as in (b). Purification by flash column chromatography (silica; EtOAc in DCM 0/100 to 40/60).
 Intermediate 26	Intermediate 25	Reaction performed as in (a). Purification by flash column chromatography (silica; MeOH in CH ₂ Cl ₂ , from 0/100 to 4/96).

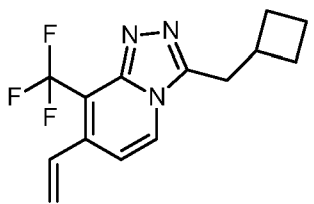
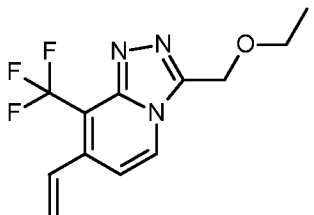
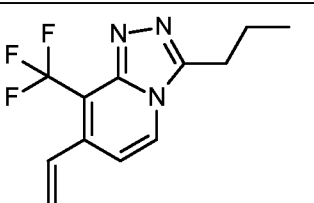
Description 7 - Intermediate 13



(Ph₃P)₄Pd (2.096 g, 1.81 mmol) was added to a stirred solution of intermediate **9** (10 g, 36.28 mmol) and 4,4,5,5-tetramethyl-2-vinyl-1,3,2-dioxaborolane ([CAS 75927-49-0], 7.77 mL, 43.53 mmol) in deoxygenated dioxane (30 mL) and a deoxygenated NaHCO₃ saturated solution (30 mL) under nitrogen. The mixture was stirred at 100 °C for 18 h. The mixture was diluted with EtOAc/water and filtered through a pad of diatomaceous earth. The filtrate was treated with brine and extracted with EtOAc. The organic layer was separated, dried (Na₂SO₄), filtered and the solvents evaporated *in vacuo*. The crude product was purified by flash column chromatography (silica; EtOAc in CH₂Cl₂ 0/100 to 5/95). The desired fractions were collected and concentrated *in vacuo* to yield intermediate **13** (6.08, 63%) as a yellow solid.

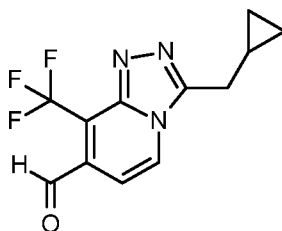
The following intermediates were synthesized following a synthetic sequence analogous to that reported in Description 7.

Intermediate	Starting material	Conditions
 Intermediate 14	Intermediate 10	Reaction performed at 150 °C. Purification by flash column chromatography (silica; 7N solution of ammonia in methanol in DCM 0/100 to 1/9).

Intermediate	Starting material	Conditions
 Intermediate 15	Intermediate 11	Extraction with DCM, purification by flash column chromatography (silica; MeOH in DCM 4/96).
 Intermediate 16	Intermediate 12	Purification by flash column chromatography (silica; EtOAc in DCM 0/100 to 10/90).
 Intermediate 27	Intermediate 26	Reaction mixture performed at 150 °C in microwave. Purification by flash column chromatography (silica; EtOAc in DCM 0/100 to 10/90).

Description 8

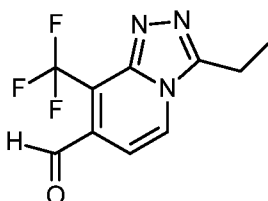
(a) Intermediate 17



- 5 Osmium tetroxide (2.5% in t-BuOH, 10.103 mL, 0.781 mmol) and then, sodium periodate 12.53 g, 58.58 mmol) in water (48.5 mL) were added to a suspension of Intermediate 13 (6.08 g, 20.02 mmol) in dioxane (192 mL). The mixture was stirred at room temperature for 2 h.
- The mixture was treated with water and EtOAc and it was filtered off through a pad of
- 10 diatomaceous earth. The filtrate was extracted with EtOAc. The organic layer was separated, dried (Na₂SO₄), filtered and the solvents evaporated *in vacuo*. The crude

product was washed with Et₂O and it was filtered and dried to yield intermediate **17** (4.25 g, 79%) as a brown solid.

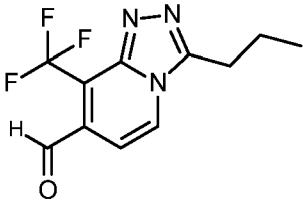
(b) Intermediate 18



- 5 A suspension of sodium periodate (5.04 g, 23.54 mmol) in distilled water (19 mL) was added to a stirred solution of osmium tetroxide (2.5% in t-BuOH, 4.06 mL, 0.31 mmol) and intermediate **14** (2.08 g, 7.85 mmol) in dioxane (75 mL). The mixture was stirred at room temperature for 150 min, and then the mixture was treated with sat NaHCO₃ and brine, and extracted with DCM. The organic layer was separated, dried
- 10 (Na₂SO₄), filtered and concentrated *in vacuo*. The product was triturated with Et₂O and filtered *in vacuo*, and finally put in desiccator at 50 °C for 18 h, to yield intermediate **18** (1.6 g, 80%) as a brown solid.

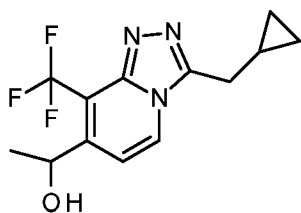
- The following intermediates were synthesized following a synthetic sequence
- 15 analogous to that reported in Description 8.

Intermediate	Starting material	Conditions
 Intermediate 19	Intermediate 15	Procedure as in (a).
 Intermediate 20	Intermediate 16	Procedure as in (a).

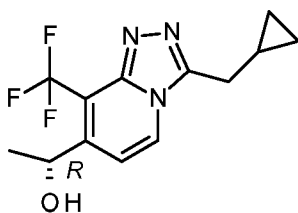
Intermediate	Starting material	Conditions
 Intermediate 28	Intermediate 27	Procedure as in (a), order of addition: osmium tetroxide was added to a stirred solution of intermediate 27 in 1,4-dioxane, then a suspension of sodium periodate in water was added and the reaction mixture was stirred for 2 h at RT. No filtration through a pad of diatomaceous earth.

Description 9

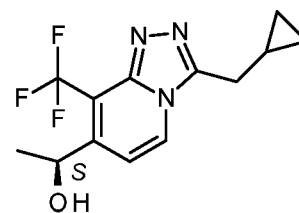
(a) Intermediates **21a**, **21b** and **21c**



Intermediate 21a

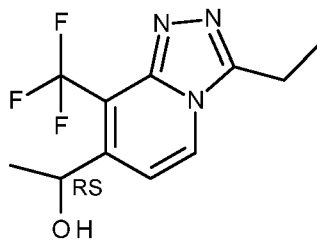


Intermediate 21b



Intermediate 21c

- Methylmagnesium bromide (1.4 M in THF, 12.40 mL, 17.37 mmol) was added dropwise to a stirred suspension of intermediate **17** (4.25 g, 15.79 mmol) in THF (281.07 mL) at -20 °C under N₂ atmosphere. The mixture was stirred at -20 °C for 45 minutes. The crude was treated with a sat. sol. of NH₄Cl and extracted with EtOAc. The organic layer was separated, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by flash column chromatography (silica; MeOH in DCM 0/100 to 4/96). The desired fractions were collected and concentrated *in vacuo* to yield intermediate **21a** (racemic mixture) (2.96 g, 66%). Intermediate **21a** (1.82 g) was purified by chiral SFC: [Stationary phase: CHIRALPAK AD-H (5μm 250 x20 mm), Mobile phase: 80% CO₂, 20% EtOH] yielding **21b** (*R*-enantiomer) (0.453 g, 10%) as a pale grey solid and intermediate **21c** (*S*-enantiomer) (0.439 g, 10%).

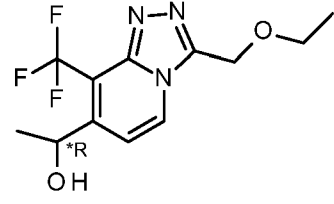
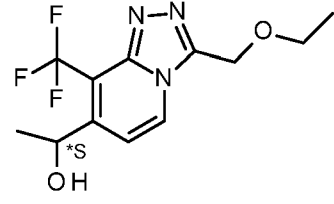
(b) Intermediate 22

Methylmagnesium bromide (1.4 M in THF, 3.97 mL, 5.56 mmol) was added dropwise to a stirred suspension of intermediate **18** (1.23 g, 5.06 mmol) in THF (90 mL) at -20 °C under N₂ atmosphere. The mixture was stirred at -20 °C for 45 minutes. The crude was treated with a sat. sol. of NH₄Cl and extracted with EtOAc. The organic layer was separated, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by flash column chromatography (silica; MeOH in DCM 0/100 to 4/96). The desired fractions were collected and concentrated *in vacuo*. The residue thus obtained was triturated with Et₂O to yield intermediate **22** (620 mg, 35%) as a pale yellow solid.

The following intermediates were synthesized following a synthetic sequence analogous to that reported in Description 9.

Intermediate	Starting material	Conditions
Intermediate 23	Intermediate 19	Procedure (b).
Intermediate 24a	Intermediate 20	Procedure (b).
Intermediate 29	Intermediate 28	Procedure (b).

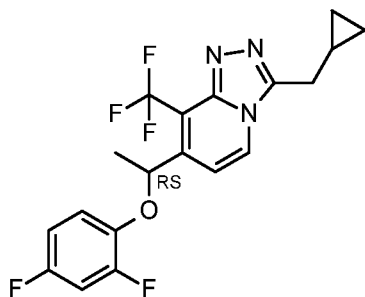
Intermediate **24a** was further separated into Intermediate **24b** and Intermediate **24c**:

 Intermediate 24b	 Intermediate 24c
Chiral SFC conditions: Stationary phase Chiralpak AD-H 5 μm 250*30 mm; Mobile phase: 80% CO₂, 15% EtOH	

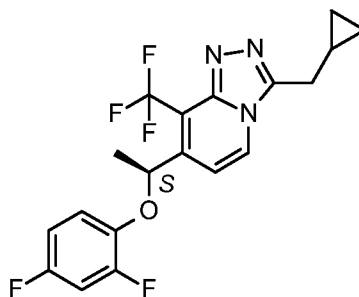
Preparation of the final compounds

Example 1

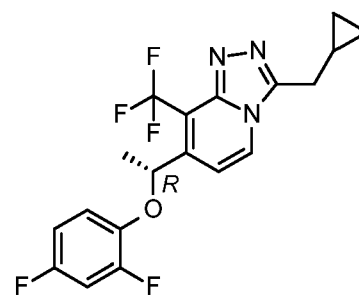
5 (a) Synthesis of compounds 4, 6 and 5



Compound 4

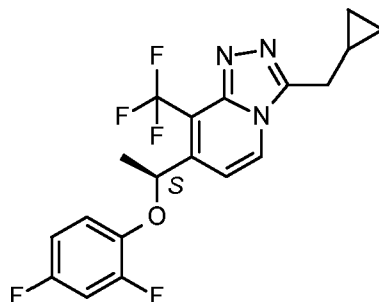


Compound 6



Compound 5

- DIAD (2.07 mL, 10.52 mmol) was added dropwise to a stirred solution of intermediate **21a** (2 g, 7.01 mmol), 2,4-difluorophenol (1.00 mL, 10.52 mmol) and triphenylphosphine (2.76 g, 10.52 mmol) in THF (74.18 mL) at 0 °C and under nitrogen atmosphere. The mixture was stirred at 100 °C for 10 minutes under microwave irradiation. The mixture was diluted with EtOAc and washed with a sat. sol. of NaHCO₃. The organic layer was separated, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by flash column chromatography (silica; MeOH in DCM 0/100 to 97/3). The desired fractions were collected and concentrated *in vacuo*.
- The residue was triturated with DIPE to give compound **4** (1.46 g, 52%) as a white solid, which was purified by chiral SFC [Stationary phase: Chiralpak AD (5 μ m 250*30 mm, Mobile phase: 85% CO₂, 15% iPrOH)], yielding compound **6** (0.659 g, 24%) and compound **5** (0.693 g, 25%).

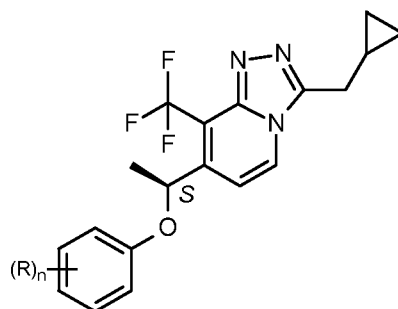
(b) Alternative synthesis of compound 6

DIAD (31.06 μ L, 0.16 mmol) was added dropwise to a stirred solution of intermediate **21b** (30 mg, 0.11 mmol), 2,4-difluorophenol (15.07 μ L, 0.16 mmol) and triphenylphosphine (41.38 mg, 0.16 mmol) in THF (1.11 mL) at 0 °C and under nitrogen atmosphere. The mixture was stirred at 100 °C for 10 minutes under microwave irradiation. The mixture was diluted with EtOAc and washed with a sat. sol. of NaHCO₃. The organic layer was separated, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by flash column chromatography (silica; MeOH in DCM 0/100 to 97/3). The desired fractions were collected and concentrated *in vacuo*. The residue was triturated with DIPE to give compound **6** (40 mg, 96%) as a white solid.

(c) Synthesis of compound 6 hydrochloride salt (.HCl)

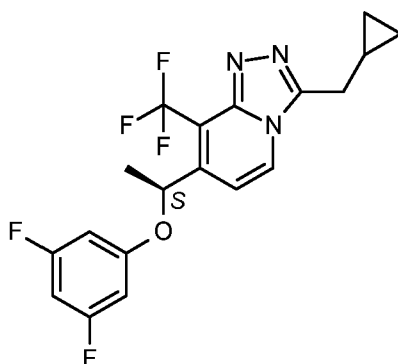
DIAD (207.06 μ L, 1.05 mmol) was added dropwise to a stirred solution of intermediate **21b** (200 mg, 0.70 mmol), 2,4-difluorophenol (100.45 μ L, 1.05 mmol) and triphenylphosphine (275.84 mg, 1.0516 mmol) in THF (4 mL) at 0 °C and under nitrogen atmosphere. The mixture was stirred at 100 °C for 15 minutes under microwave irradiation. The mixture was diluted with EtOAc and washed with a sat. sol. of NaHCO₃. The organic layer was separated, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by RP HPLC (Stationary phase: C18 XBridge 30 x 100 mm 5 μ m, Mobile phase: Gradient from 60% 0.1% NH₄CO₃H/NH₄OH pH 9 solution in Water, 40% CH₃CN to 43% 0.1% NH₄CO₃H/NH₄OH pH 9 solution in Water, 57% CH₃CN), yielding a white solid residue that was dissolved in Et₂O (8 mL) and 1,4-dioxane (0.5 mL). To the solution thus obtained HCl (4M in dioxane, 200 μ L) was added dropwise. The white solid precipitate was filtered, washed with Et₂O, dried (Na₂SO₄) and evaporated under vacuum. The white residue thus obtained was triturated with Et₂O to give compound **6** .HCl (110 mg, 36%) as a white solid.

The following compounds were synthesized following a synthetic sequence analogous to that reported in Example 1(b), starting from intermediate **21b**.



Co. No.	(R) _n	Co. No.	(R) _n
9		12	
10		13	
11		14	

5 Example 2 - Synthesis of compound 7

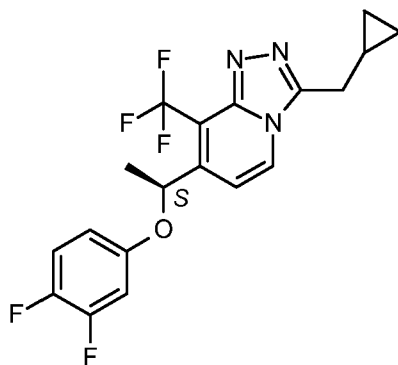


Procedure (a): DIAD (31.06 μ L, 0.158 mmol) was added dropwise to a stirred solution of intermediate **21b** (30 mg, 0.105 mmol), 3,5-difluorophenol (20.52 mg, 0.158 mmol)

and triphenylphosphine (41.38 mg, 0.158 mmol) in THF (1.113 mL) at 0 °C and under nitrogen atmosphere. The mixture was stirred at 100 °C for 10 minutes under microwave irradiation. The mixture was diluted with EtOAc and washed with a sat. sol. of NaHCO₃. The organic layer was separated, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by flash column chromatography (silica; MeOH in DCM 0/100 to 96/4). The desired fractions were collected and concentrated *in vacuo*. The residue was triturated with DIPE to give compound **7** (21 mg, 50%) as a white solid.

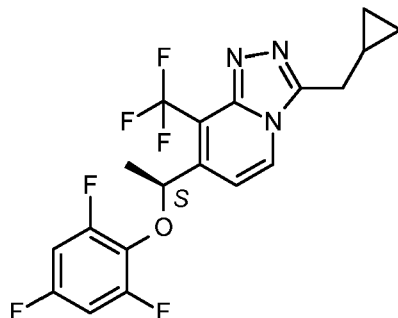
Procedure (b): Alternatively, compound **7** was also synthesized following a synthetic sequence analogous to that reported in Example 1(b), starting from intermediate **21b**.

Example 3 - Synthesis of compound **8**



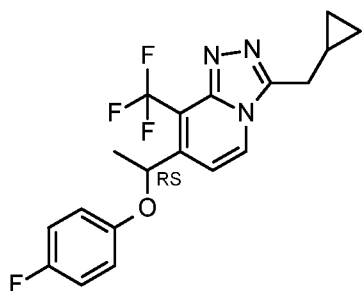
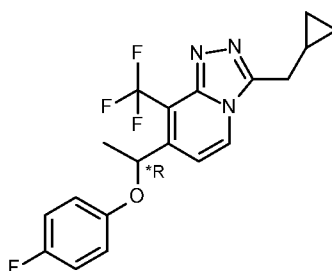
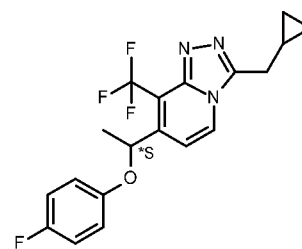
Procedure (a): DIAD (31.06 µL, 0.158 mmol) was added dropwise to a stirred solution of intermediate **21b** (30 mg, 0.105 mmol), 3,4-difluorophenol (20.52 mg, 0.158 mmol) and triphenylphosphine (41.38 mg, 0.158 mmol) in THF (1.11 mL) at 0 °C and under nitrogen atmosphere. The mixture was stirred at 100 °C for 10 minutes under microwave irradiation. The mixture was diluted with EtOAc and washed with a sat. sol. of NaHCO₃. The organic layer was separated, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by flash column chromatography (silica; MeOH in DCM 0/100 to 96/4). The desired fractions were collected and concentrated *in vacuo*. The residue was triturated with DIPE to give compound **8** (10.6 mg, 25%) as a white solid.

Procedure (b): Alternatively, compound **8** was also synthesized following a synthetic sequence analogous to that reported in Example 1(b), starting from intermediate **21b**.

Example 4 - Synthesis of compound 15

Procedure (a): DIAD (155.3 μ L, 0.789 mmol) was added dropwise to a stirred solution of intermediate **21b** (150 mg, 0.526 mmol), 2,4,6-trifluorophenol (116.8 mg, 0.789 mol) and triphenylphosphine (206.88 mg, 0.789 mmol) in THF (5.56 mL) at 0 $^{\circ}$ C and under nitrogen atmosphere. The mixture was stirred at 100 $^{\circ}$ C for 10 minutes under microwave. The mixture was diluted with DCM and washed with a sat. sol. of NaHCO₃. The organic layer was separated, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by flash column chromatography (silica; MeOH/NH₃ 7 N in DCM 0/100 to 90/10). The desired fractions were collected and concentrated *in vacuo*, then purified by RP HPLC (Stationary phase: C18 XBridge 30 x 100 mm 5 μ m, Mobile phase: Gradient from 54% 0.1% NH₄CO₃H/NH₄OH pH 9 solution in Water, 46% CH₃CN to 64% 0.1% NH₄CO₃H/NH₄OH pH 9 solution in Water, 36% CH₃CN) yielding a colourless oil that was crystallized upon standing (2 days). The solid was triturated with heptane to give compound **15** (129.8 mg, 59%) as a white solid.

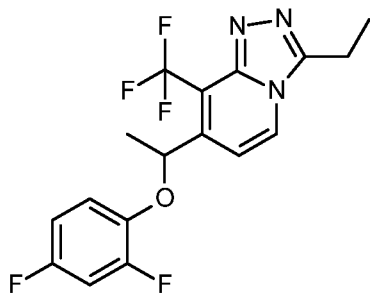
Procedure (b): Alternatively, compound **15** was also synthesized following a synthetic sequence analogous to that reported in Example 1(b), starting from intermediate **21b**.

Example 5 - Synthesis of compounds 1, 2 and 3**Compound 1****Compound 2****Compound 3**

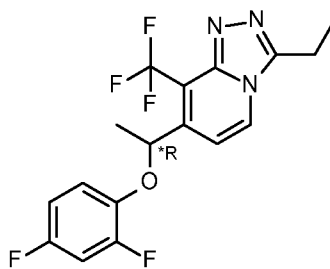
Compounds **1**, **2** and **3** were synthesized following the procedure described in Example 1(a). Thus, reaction of DIAD (500.05 μ L, 2.54 mmol), intermediate **21a** (483 mg, 1.69 mmol), 4-fluorophenol (227.77 mg, 2.03 mmol) and triphenylphosphine (666.14 mg, 2.54 mmol) in THF (17.91 mL) as described in Example 1(a) yielded a residue that was purified by flash column chromatography (silica; EtOAc in DCM 0/100 to 90/10). The

desired fractions were collected and concentrated *in vacuo*. The resulting residue was triturated with DIPE to yield compound **1** (320 mg, 50%) as a white solid, which was purified by chiral SFC [Stationary phase: Chiralpak AD (5 μ m 250*30 mm, Mobile phase: 77% CO₂, 23% MeOH)], yielding compound **2** (131mg, 20%) and compound **3** (129 mg, 20%) as white solids.

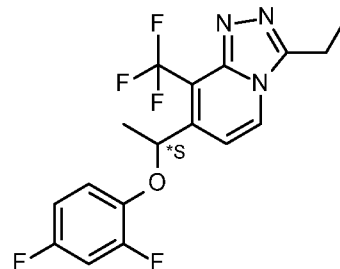
Example 6 - Synthesis of compounds **24**, **26**, and **27**



Compound 24



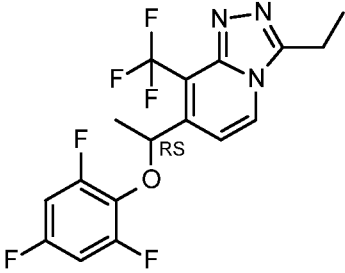
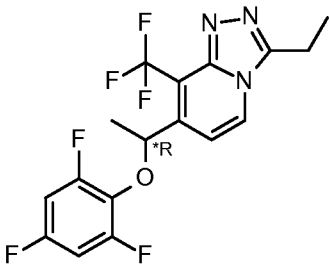
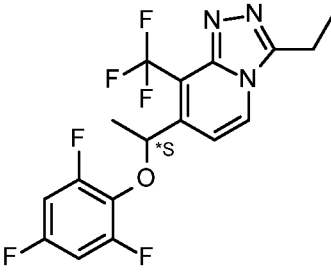
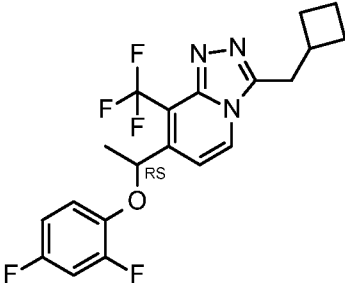
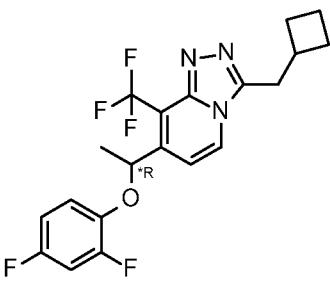
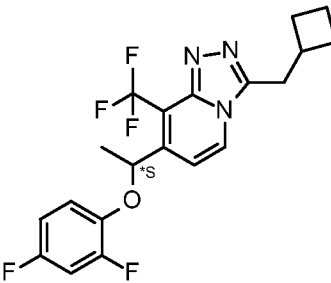
Compound 26



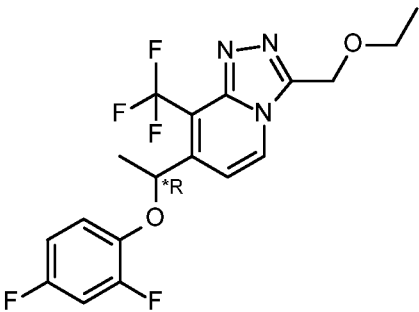
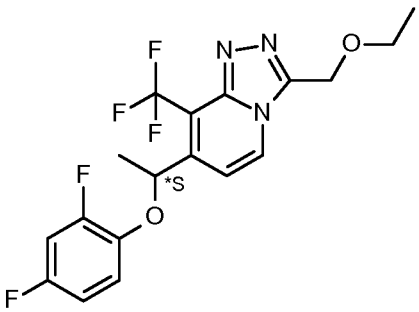
Compound 27

Compounds **24**, **26** and **27** were synthesized following the procedure described in Example 1(a). Thus, reaction of DIAD (364.57 μ L, 1.85 mmol), intermediate **22** (320 mg, 1.23 mmol), 2,4-difluorophenol (176.86 μ L, 1.85 mmol) and triphenylphosphine (485.67 mg, 1.85 mmol) in THF (13.06 mL) as described in Example 1(a) yielded a residue that was purified by flash column chromatography (silica; MeOH in DCM 0/100 to 96/4). The desired fractions were collected and concentrated *in vacuo* to yield a colourless oil that crystallized with DIPE to give compound **24** as a white solid, which was purified by RP HPLC (Stationary phase: C18 XBridge 30 x 100 mm 5 μ m; mobile phase: Gradient from 54% 0.1% NH₄CO₃H/NH₄OH pH 9 solution in Water, 46% CH₃CN to 64% 0.1% NH₄CO₃H/NH₄OH pH 9 solution in Water, 36% CH₃CN) yielding a colourless oil that was crystallized upon trituration with heptane to give 240 mg (52%) of compound **24** as a white solid, which was then purified by chiral SFC (Stationary phase: CHIRALPAK AD-H 5 μ m 250x20mm; mobile phase: 85% CO₂, 15% iPOH (0.3% iPrNH₂)), yielding compound **26** (103 mg, 22%) and compound **27** (107 mg, 23%).

The following compounds were obtained following a synthetic sequence similar to that reported in Example 1(a).

 Compound 25	 Compound 28	 Compound 29
Starting material: Intermediate 22 Chiral SFC conditions: Stationary phase: Chiralpak AD-H 5μm 250 x 20 mm); Mobile phase: 85% CO₂, 15% mixture of EtOH/iPrOH 50/50 v/v (+0.3% iPrNH₂)		
 Compound 16	 Compound 17	 Compound 18
Starting material: Intermediate 23 Chiral SFC conditions: Stationary phase: Chiralpak AD-H (5 μm 250*30 mm); Mobile phase: 80% CO₂, 20% mixture MeOH/iPrOH 50/50 v/v (+0.3% iPrNH₂)		

The following compounds were synthesized following a synthetic sequence analogous to that reported in Example 1(b), starting from the indicated intermediates.

 Compound 20; obtained from intermediate 24c	 Compound 21; obtained from intermediate 24b
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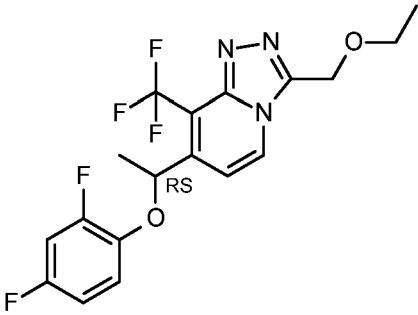
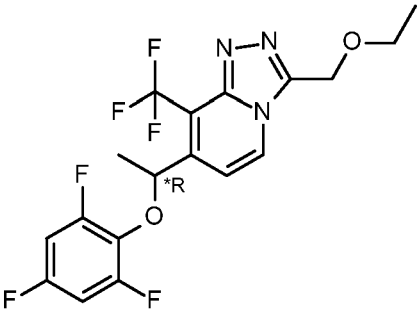
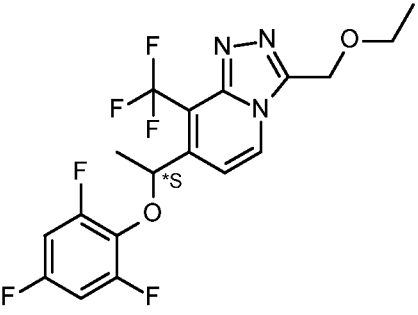
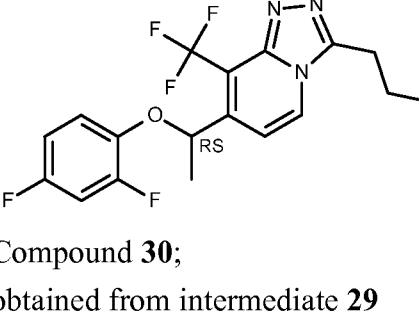
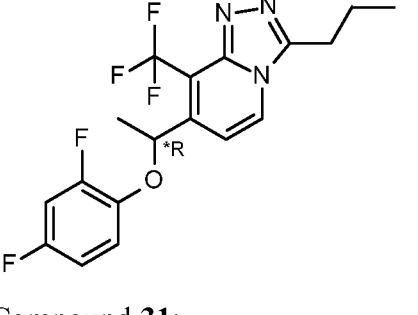
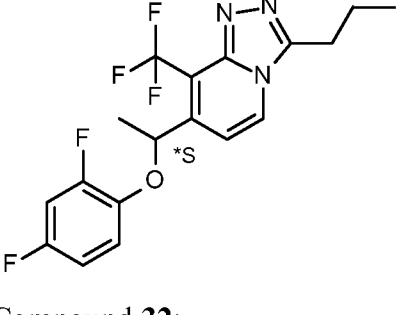
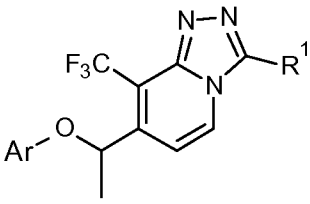
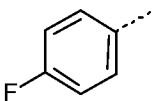
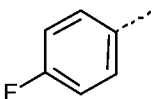
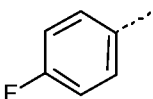
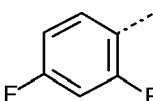
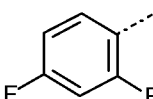
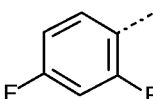
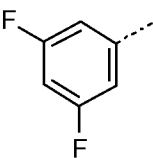
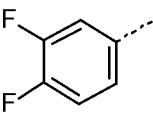
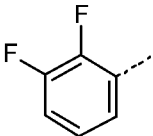
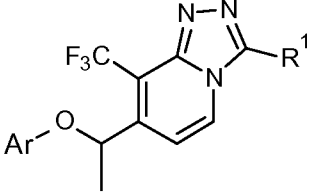
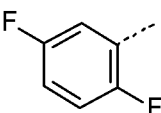
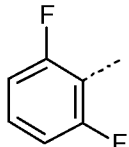
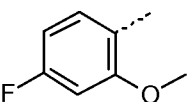
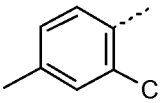
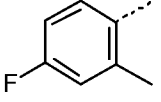
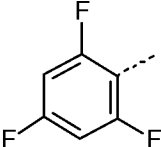
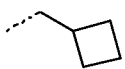
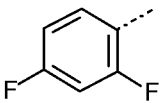
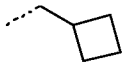
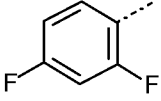
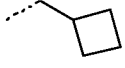
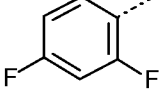
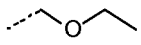
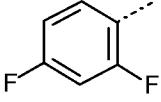
 Compound 19; obtained from intermediate 24a	 Compound 22; obtained from intermediate 24c
 Compound 23; obtained from intermediate 24b	 Compound 30; obtained from intermediate 29
 Compound 31;	 Compound 32;
Starting material: Intermediate 30 Chiral SFC conditions: Stationary phase: Chiralpak AD-H 5μm 250 x 20 mm); Mobile phase: 85% CO₂, 15% iPrOH.	

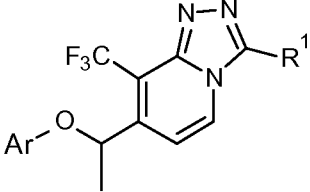
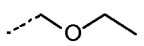
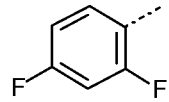
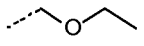
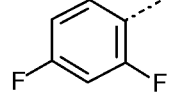
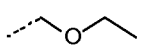
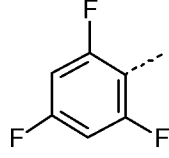
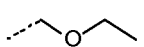
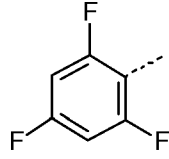
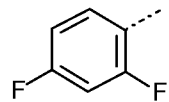
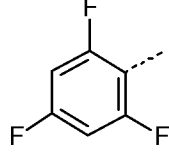
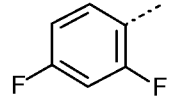
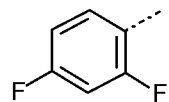
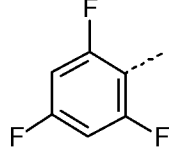
Table 1 below lists additional compounds of Formula (I) which were prepared by analogy to the above examples (Exp. no.).

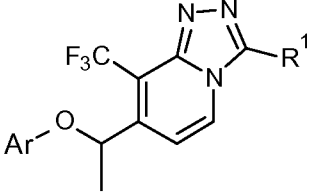
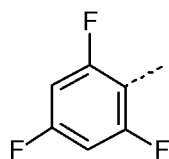
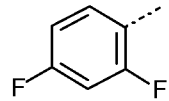
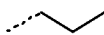
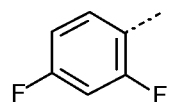
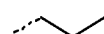
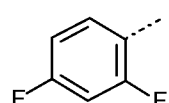
Table 1 : Example compounds according to Formula (I).

indicates that the experimental procedure is described in the examples.

				
Co. no.	Exp no.	R ¹	Ar	Stereo-chem.
1	E5 [#]			RS
2	E5 [#]			*R
3	E5 [#]			*S
4	E1 [#]			RS
5	E1 [#]			R
6	E1(a) and (b) [#]			S
6.HCl	E1(c)*			
7	E2 [#]			S
8	E3 [#]			S
9	E1(b)			S

				
Co. no.	Exp no.	R ¹	Ar	Stereo-chem.
10	E1(b)			S
11	E1(b)			S
12	E1(b)			S
13	E1(b)			S
14	E1(b)			S
15	E4 [#]			S
16	E1(a)			RS
17	E1(a)			*R
18	E1(a)			*S
19	E1(b)			RS

				
Co. no.	Exp no.	R ¹	Ar	Stereo-chem.
20	E1(b)			*R
21	E1(b)			*S
22	E1(b)			*R
23	E1(b)			*S
24	E6 [#]			RS
25	E1(a)			RS
26	E6 [#]			*R
27	E6 [#]			*S
28	E1(a)			*R

				
Co. no.	Exp no.	R ¹	Ar	Stereo-chem.
29	E1(a)			*S
30	E1(b)			RS
31	E1(b)			*R
32	E1(b)			*S

Analytical part

Optical Rotations

- Optical rotations were measured on a Perkin-Elmer 341 polarimeter with a sodium lamp and reported as follows: $[\alpha]^{\circ}(\lambda, c \text{ g/100ml, solvent, } T^{\circ}\text{C})$.
- 5 $[\alpha]_{\lambda}^T = (100\alpha) / (l \times c)$: where l is the path length in dm and c is the concentration in g/100 ml for a sample at a temperature T ($^{\circ}\text{C}$) and a wavelength λ (in nm). If the wavelength of light used is 589 nm (the sodium D line), then the symbol D might be used instead. The sign of the rotation (+ or -) should always be given. When using this
- 10 equation the concentration and solvent are always provided in parentheses after the rotation. The rotation is reported using degrees and no units of concentration are given (it is assumed to be g/100 ml).

LCMS

- For (LC)MS-characterization of the compounds of the present invention, the following
- 15 methods were used.

General procedure

The High Performance Liquid Chromatography (HPLC) measurement was performed using a LC pump, a diode-array (DAD) or a UV detector and a column as specified in

the respective methods. If necessary, additional detectors were included (see table of methods below).

- Flow from the column was brought to the Mass Spectrometer (MS) which was configured with an atmospheric pressure ion source. It is within the knowledge of the skilled person to set the tune parameters (e.g. scanning range, dwell time...) in order to obtain ions allowing the identification of the compound's nominal monoisotopic molecular weight (MW). Data acquisition was performed with appropriate software. Compounds are described by their experimental retention times (R_t) and ions. If not specified differently in the table of data, the reported molecular ion corresponds to the $[M+H]^+$ (protonated molecule) and/or $[M-H]^-$ (deprotonated molecule). In case the compound was not directly ionizable the type of adduct is specified (i.e. $[M+NH_4]^+$, $[M+HCOO]^-$, etc...). For molecules with multiple isotopic patterns (Br, Cl...), the reported value is the one obtained for the lowest isotope mass. All results were obtained with experimental uncertainties that are commonly associated with the method used.
- Hereinafter, "SQD" means Single Quadrupole Detector, "RT" room temperature, "BEH" bridged ethylsiloxane/silica hybrid, "HSS" High Strength Silica, "DAD" Diode Array Detector.

Table 2. LCMS Method codes (Flow expressed in mL/min; column temperature (T) in °C; Run time in minutes).

Instru- ment	Column	Mobile phase	Gradient	Flow ----- Col T	Run time	LCMS Method
Waters: Acquity® UPLC® - DAD and SQD	Agilent: Eclipse Plus C18 RRHD (1.8µm, 2.1x50mm)	A: 95% CH ₃ COONH ₄ 6.5mM + 5% CH ₃ CN, B: CH ₃ CN	From 95% A to 5% A in 4.6min, held for 0.4min	1 ----- 50	5	1
Waters: Acquity® UPLC® - DAD and SQD	Waters: CSH™ C18 (1.7µm, 2.1x50mm)	A: 95% CH ₃ COONH ₄ 6.5mM + 5% CH ₃ CN, B: CH ₃ CN	From 95% A to 5% A in 4.6min, held for 0.4min	1 ----- 50	5	2

Instru- ment	Column	Mobile phase	Gradient	Flow ----- Col T	Run time	LCMS Method
Waters: Acquity UPLC [®] - DAD and Quattro Micro [™]	Waters: BEH C18 (1.7µm, 2.1x100mm)	A: 95% CH ₃ COONH ₄ 7mM / 5% CH ₃ CN, B: CH ₃ CN	84.2% A for 0.49min, to 10.5% A in 2.18min, held for 1.94min, back to 84.2% A in 0.73min, held for 0.73min.	0.343 ----- 40	6.2	3
Waters: Acquity [®] UPLC [®] - DAD and SQD	Waters: CSH [™] C18 (1.7µm, 2.1x50mm)	A: 95% CH ₃ COONH ₄ 6.5mM + 5% CH ₃ CN, B: CH ₃ CN	From 95% A to 5% A in 7.8min, held for 1.2min	1 ----- 50	9	4

Melting points

Values are peak values, and are obtained with experimental uncertainties that are commonly associated with this analytical method.

5 Mettler FP 81HT / FP90 apparatus

For a number of compounds, melting points were determined in open capillary tubes on a FP 81HT / FP90 apparatus (Mettler-Toledo). Melting points were measured with a temperature gradient of 1, 3, 5 or 10 °C/minute. Maximum temperature was 300 °C. The melting point was read from a digital display.

Table 3 : Physico-chemical data for some compounds, retention time (R_t) in min, $[M+H]^+$ peak (protonated molecule), LCMS method and mp (melting point in °C). (n.d. = not determined).

Co. no.	Mp (°C)	R_t (min)	$[MH^+]$	LCMS method	Optical Rotation
1	156.3	2.32	380	1	
2	176.9	2.93	380	3	-58.5 ° (589 nm, c 0.53 w/v%, DMF, 20 °C)
3	177.3	2.93	380	3	+59.4 ° (589 nm, c 0.52 w/v%, DMF, 20 °C)
4	121.7	2.41	398	1	
5	142	2.99	398.3	3	+95.7 ° (589 nm, c 0.69 w/v%, DMF, 20 °C)
6	142.4	2.99	398.2	3	-95.4 ° (589 nm, c 0.7 w/v%, DMF, 20 °C)
7	170.08	2.37	398	2	-55.7 ° (589 nm, c 0.96 w/v %, DMF, 20 °C)
8	n.d.	2.32	398	2	n.d.
9	n.d.	2.32	398	2	n.d.
10	n.d.	2.25	398	2	n.d.
11	n.d.	2.28	398	2	n.d.
12	n.d.	2.16	410	2	n.d.
13	144.1	2.68	410	2	n.d.
14	161.7	2.51	394	2	n.d.
15	80.3	2.37	416	2	-167.0 ° (589 nm, c 0.55 w/v%, DMF, 20 °C)

Co. no.	Mp (°C)	R _t (min)	[MH ⁺]	LCMS method	Optical Rotation
16	n.d.	2.50	412	2	n.d.
17	n.d.	3.12	412	3	n.d.
18	n.d.	3.12	412	3	n.d.
19	n.d.	2.39	402	2	n.d.
20	n.d.	2.3	402	2	n.d.
21	n.d.	3.36	402		n.d.
22	n.d.	2.35	420	2	n.d.
23	n.d.	2.35	420	2	n.d.
24	135.7	2.05	372	2	n.d.
25	138.3	2.13	390	2	n.d.
26	n.d.	2.80	372	3	-83.9 ° (589 nm, c 0.52 w/v %, DMF, 25 °C)
27	n.d.	2.80	372	3	+92.1 ° (589 nm, c 0.55 w/v %, DMF, 25 °C)
28	n.d.	2.85	390	3	-129.2 ° (589 nm, c 0.5 w/v %, DMF, 25 °C)
29	n.d.	2.85	390	3	+137.3 ° (589 nm, c 0.51 w/v %, DMF, 25 °C)
30	130.6	2.29	386	2	n.d.
31	127.85	2.29	386	2	-67.5 ° (589 nm, c 0.83 w/v %, DMF, 20 °C)
32	127.69	2.29	386	2	+89.5 ° (589 nm, c 0.83 w/v %, DMF, 20 °C)

SFC-MS***General procedure***

The SFC measurement was performed using Analytical system from Berger instrument comprising a FCM-1200 dual pump fluid control module for delivering carbon dioxide (CO₂) and modifier, a CTC Analytics automatic liquid sampler, a TCM-20000 thermal control module for column heating from room temperature to 80°C. An Agilent 1100 UV photodiode array detector equipped with a high-pressure flow cell standing up to 400 bars was used. Flow from the column was split to a MS spectrometer. The MS detector was configured with an atmospheric pressure ionization source. The following ionization parameters for the Waters ZQ mass spectrophotometer are: corona: 9µa, source temp: 140°C, cone: 30 V, probe temp 450°C, extractor 3 V, desolvation gas 400L/hr, cone gas 70 L/hr. Nitrogen was used as the nebulizer gas. Data acquisition was performed with a Waters-Micromass MassLynx-Openlynx data system.

Method 1: In addition to the general procedure: The analytical chiral separation in SFC-MS was carried out on a CHIRALPAK AD DAICEL column (10 µm, 4.6 x 250 mm) at 35 °C with a flow rate of 3.0 ml/min. The mobile phase is 85% CO₂, 15% iPrOH (+ 0.3% iPrNH₂) hold 7 min in isocratic mode.

Method 2: In addition to the general procedure: The analytical chiral separation in SFC-MS was carried out on a CHIRALPAK AD DAICEL column (10 µm, 4.6 x 250 mm) at 35 °C with a flow rate of 3.0 ml/min. The mobile phase is 75% CO₂, 15% iPrOH (+ 0.3% iPrNH₂) hold 7 min in isocratic mode.

Method 3: In addition to the general procedure: The analytical chiral separation in SFC-MS was carried out on a CHIRALPAK AD DAICEL column (10 µm, 4.6 x 250 mm) at 35 °C with a flow rate of 3.0 ml/min. The mobile phase is 80% CO₂, 10% Methanol + 10% iPrOH (+ 0.3% iPrNH₂) hold 7 min in isocratic mode.

Table 4: Analytical SFC data – R_t means retention time (in minutes), [M+H]⁺ means the protonated mass of the compound, method refers to the method used for SFC/MS analysis of enantiomerically pure compounds. The measurement was compared against the mixture.

Co. Nr.	R _t	[M+H] ⁺	UV Area %	Method	Isomer Elution Order*
6	4.28	398	100	1	A
5	5.98	398	100	1	B
2	2.13	380	100	2	A

Co. Nr.	R _t	[M+H] ⁺	UV Area %	Method	Isomer Elution Order*
3	2.97	380	100	2	B
17	2.46	412	100	3	A
18	3.12	412	100	3	B
31	2.93	386	100	1	A
32	3.81	386	100	1	B

*A means the first isomer that elutes. B means the second isomer that elutes.

Nuclear Magnetic Resonance (NMR)

For a number of compounds, ¹H NMR spectra were recorded either on a Bruker DPX-400 or on a Bruker AV-500 spectrometer with standard pulse sequences, operating at 400 MHz and 500 MHz respectively. Chemical shifts (δ) are reported in parts per million (ppm) downfield from tetramethylsilane (TMS), which was used as internal standard.

Co. No. 6: ¹H NMR (400 MHz, CDCl₃) δ ppm 0.30 - 0.38 (m, 2 H), 0.59 - 0.68 (m, 2 H), 1.14 - 1.22 (m, 1 H), 1.72 (d, *J*=6.5 Hz, 3 H), 3.02 - 3.14 (m, 2 H), 5.84 (q, *J*=6.3 Hz, 1 H), 6.67 - 6.73 (m, 1 H), 6.80 - 6.89 (m, 2 H), 7.30 (d, *J*=7.4 Hz, 1 H), 8.11 (d, *J*=7.4 Hz, 1 H)

Co. No. 7: ¹H NMR (400 MHz, CDCl₃) δ ppm 0.30 - 0.39 (m, 2 H), 0.59 - 0.68 (m, 2 H), 1.11 - 1.23 (m, 1 H), 1.70 (d, *J*=6.5 Hz, 3 H), 3.01 - 3.14 (m, 2 H), 5.83 (q, *J*=6.2 Hz, 1 H), 6.35 - 6.45 (m, 3 H), 7.13 (d, *J*=7.2 Hz, 1 H), 8.08 (d, *J*=7.4 Hz, 1 H)

Co. No. 8: ¹H NMR (400 MHz, CDCl₃) δ ppm 0.30 - 0.38 (m, 2 H), 0.58 - 0.68 (m, 2 H), 1.11 - 1.22 (m, 1 H), 1.69 (d, *J*=6.2 Hz, 3 H), 3.01 - 3.13 (m, 2 H), 5.79 (q, *J*=6.2 Hz, 1 H), 6.53 (dtd, *J*=9.2, 3.1, 3.1, 1.7 Hz, 1 H), 6.72 (ddd, *J*=11.6, 6.5, 3.1 Hz, 1 H), 6.95 - 7.04 (m, 1 H), 7.15 (d, *J*=7.4 Hz, 1 H), 8.07 (d, *J*=7.4 Hz, 1 H)

Co. No. 15: ¹H NMR (500 MHz, CDCl₃) δ ppm 0.30 - 0.41 (m, 2 H), 0.59 - 0.71 (m, 2 H), 1.16 - 1.25 (m, 1 H), 1.70 (d, *J*=6.4 Hz, 3 H), 3.05 - 3.16 (m, 2 H), 5.80 (q, *J*=6.4 Hz, 1 H), 6.62 - 6.70 (m, 2 H), 7.45 (d, *J*=7.5 Hz, 1 H), 8.16 (d, *J*=7.2 Hz, 1 H)

25

Co. No. 13: ¹H NMR (500 MHz, CDCl₃) δ ppm 0.27 - 0.39 (m, 2 H), 0.58 - 0.67 (m, 2 H), 1.12 - 1.21 (m, 1 H), 1.73 (d, *J*=6.4 Hz, 3 H), 2.22 (s, 3 H), 3.06 (qd, *J*=15.4, 6.6

Hz, 2 H), 5.92 (q, $J=6.4$ Hz, 1 H), 6.71 (d, $J=8.4$ Hz, 1 H), 6.89 (dd, $J=8.4$, 1.4 Hz, 1 H), 7.18 (d, $J=1.7$ Hz, 1 H), 7.32 (d, $J=7.2$ Hz, 1 H), 8.07 (d, $J=7.2$ Hz, 1 H)

5 **Co. No. 14:** ^1H NMR (500 MHz, CDCl_3) δ ppm 0.28 - 0.39 (m, 2 H), 0.57 - 0.69 (m, 2 H), 1.12 - 1.21 (m, 1 H), 1.70 (d, $J=6.6$ Hz, 3 H), 2.31 (s, 3 H), 3.01 - 3.12 (m, 2 H), 5.79 (q, $J=6.6$ Hz, 1 H), 6.55 (dd, $J=9.0$, 4.3 Hz, 1 H), 6.69 (td, $J=8.5$, 3.0 Hz, 1 H), 6.87 (dd, $J=9.0$, 2.9 Hz, 1 H), 7.17 (d, $J=7.5$ Hz, 1 H), 8.06 (d, $J=7.2$ Hz, 1 H)

10 **Co. No. 20:** ^1H NMR (500 MHz, CDCl_3) δ ppm 1.22 (t, $J=7.1$ Hz, 3 H), 1.72 (d, $J=6.4$ Hz, 3 H), 3.58 (q, $J=7.1$ Hz, 2 H), 5.03 - 5.10 (m, 2 H), 5.84 (q, $J=6.5$ Hz, 1 H), 6.67 - 6.74 (m, 1 H), 6.81 - 6.88 (m, 2 H), 7.34 (d, $J=7.2$ Hz, 1 H), 8.40 (d, $J=7.5$ Hz, 1 H)

15 **Co. No. 22:** ^1H NMR (500 MHz, CDCl_3) δ ppm 1.23 (t, $J=6.9$ Hz, 3 H), 1.70 (d, $J=6.4$ Hz, 3 H), 3.58 (q, $J=7.0$ Hz, 2 H), 5.05 - 5.12 (m, 2 H), 5.81 (q, $J=6.6$ Hz, 1 H), 6.62 - 6.70 (m, 2 H), 7.48 (d, $J=7.5$ Hz, 1 H), 8.45 (d, $J=7.2$ Hz, 1 H)

20 **Co. No. 31:** ^1H NMR (400 MHz, CDCl_3) δ ppm 1.07 (t, $J=7.40$ Hz, 3 H) 1.72 (d, $J=6.24$ Hz, 3 H) 1.92 (sxt, $J=7.63$ Hz, 2 H) 2.98 - 3.14 (m, 2 H) 5.84 (q, $J=6.47$ Hz, 1 H) 6.65 - 6.74 (m, 1 H) 6.78 - 6.89 (m, 2 H) 7.29 (d, $J=7.40$ Hz, 1 H) 8.02 (d, $J=7.40$ Hz, 1 H).

Pharmacological examples

A) In vitro pharmacology

25 The compounds provided in the present invention are positive allosteric modulators of mGluR2. These compounds appear to potentiate glutamate responses by binding to an allosteric site other than the glutamate binding site. The response of mGluR2 to a concentration of glutamate is increased when compounds of Formula (I) are present. Compounds of Formula (I) are expected to have their effect substantially at mGluR2 by virtue of their ability to enhance the function of the receptor. The effects of positive
30 allosteric modulators tested at mGluR2 using the [^{35}S]GTP γ S binding assay method described below and which is suitable for the identification of such compounds, and more particularly the compounds according to Formula (I), are shown in Table 5.

[^{35}S]GTP γ S binding assay

35 The [^{35}S]GTP γ S binding assay is a functional membrane-based assay used to study G-protein coupled receptor (GPCR) function whereby incorporation of a

non-hydrolysable form of GTP, [³⁵S]GTPγS (guanosine 5'-triphosphate, labelled with gamma-emitting ³⁵S), is measured. The G-protein α subunit catalyzes the exchange of guanosine 5'-diphosphate (GDP) by guanosine triphosphate (GTP) and on activation of the GPCR by an agonist, [³⁵S]GTPγS, becomes incorporated and cannot be cleaved to continue the exchange cycle (Harper (1998) Current Protocols in Pharmacology 2.6.1-10, John Wiley & Sons, Inc.). The amount of radioactive [³⁵S]GTPγS incorporation is a direct measure of the activity of the G-protein and hence the activity of the agonist can be determined. mGlu2 receptors are shown to be preferentially coupled to Gαi-protein, a preferential coupling for this method, and hence it is widely used to study receptor activation of mGlu2 receptors both in recombinant cell lines and in tissues. Here we describe the use of the [³⁵S]GTPγS binding assay using membranes from cells transfected with the human mGlu2 receptor and adapted from Schaffhauser *et al.* (Molecular Pharmacology, **2003**, 4:798-810) for the detection of the positive allosteric modulation (PAM) properties of the compounds of this invention.

15 **Membrane preparation**

CHO-cells were cultured to pre-confluence and stimulated with 5 mM butyrate for 24 h. Cells were then collected by scraping in PBS and cell suspension was centrifuged (10 min at 4000 RPM in benchtop centrifuge). Supernatant was discarded and pellet gently resuspended in 50 mM Tris-HCl, pH 7.4 by mixing with a vortex and pipetting up and down. The suspension was centrifuged at 16,000 RPM (Sorvall RC-5C plus rotor SS-34) for 10 minutes and the supernatant discarded. The pellet was homogenized in 5 mM Tris-HCl, pH 7.4 using an ultra-turrax homogenizer and centrifuged again (18,000 RPM, 20 min, 4 °C). The final pellet was resuspended in 50 mM Tris-HCl, pH 7.4 and stored at -80 °C in appropriate aliquots before use. Protein concentration was determined by the Bradford method (Bio-Rad, USA) with bovine serum albumin as standard.

[³⁵S]GTPγS binding assay

Measurement of mGluR2 positive allosteric modulatory activity of test compounds was performed as follows. Test compounds and glutamate were diluted in assay buffer containing 10 mM HEPES acid, 10 mM HEPES salt, pH 7.4, 100 mM NaCl, 3 mM MgCl₂ and 10 μM GDP. Human mGlu2 receptor-containing membranes were thawed on ice and diluted in assay buffer supplemented with 14 μg/ml saponin. Membranes were pre-incubated with compound alone or together with a predefined (~EC₂₀) concentration of glutamate (PAM assay) for 30 min at 30 °C. After addition of [³⁵S]GTPγS (f.c. 0.1 nM), assay mixtures were shaken briefly and further incubated to allow [³⁵S]GTPγS incorporation on activation (30 minutes, 30 °C). Final assay

mixtures contained 7 μ g of membrane protein in 10 mM HEPES acid, 10 mM HEPES salt, pH 7.4, 100 mM NaCl, 3 mM MgCl₂, 10 μ M GDP and 2 μ g/ml saponin. Total reaction volume was 200 μ l. Reactions were terminated by rapid filtration through Unifilter-96 GF/B plates (Perkin Elmer, Massachusetts, USA) using a 96-well filtermate universal harvester. Filters were washed 6 times with ice-cold 10 mM NaH₂PO₄/10 mM Na₂HPO₄, pH 7.4. Filters were then air-dried, and 40 μ l of liquid scintillation cocktail (Microscint-O) was added to each well. Membrane-bound radioactivity was counted in a Microplate Scintillation and Luminescence Counter from Perkin Elmer.

10 Data analysis

The concentration-response curves of representative compounds of the present invention -obtained in the presence of EC₂₀ of mGluR2 agonist glutamate to determine positive allosteric modulation (PAM)- were generated using the Lexis software interface (developed at J&J). Data were calculated as % of the control glutamate response, defined as the maximal response that is generated upon addition of glutamate alone. Sigmoid concentration-response curves plotting these percentages versus the log concentration of the test compound were analyzed using non-linear regression analysis. The concentration producing half-maximal effect is then calculated as EC₅₀. The pEC₅₀ values below were calculated as the -log EC₅₀, when the EC₅₀ is expressed in M. E_{max} is defined as relative maximal effect (i.e. maximal % effect relative to the control glutamate response).

Table 5 below shows the pharmacological data obtained for compounds of Formula (I).

Table 5. Pharmacological data for compounds according to the invention.

Co. No.	GTP γ S - hmGluR2 PAM pEC ₅₀	GTP γ S - hmGluR2 PAM E _{max}
1	6.59	296
2	6.84	228
3	5.79	187
6	7.39	256
5	6.06	141
4	7.04	329
7	7.31	292
8	7.04	244
9	7.3	260

Co. No.	GTPγS - hmGluR2 PAM pEC₅₀	GTPγS - hmGluR2 PAM E_{max}
10	7.47	218
11	8.25	239
12	6.99	178
16	7.54	284
13	7.75	280
14	7.53	281
15	8.16	293
19	6.71	297
25	6.9	233
24	6.42	193
17	7.73	317
18	6.24	213
22	7.61	325
23	5.94	167
21	6.32	102
20	7.07	332
26	6.78	214
27	n.c.	51
30	6.9	227
28	7.19	234
29	5.85	77
31	7.05	251
32	5.71	116

n.c. means that the pEC₅₀ could not be calculated

pEC₅₀ values were not calculated in cases where the concentration-response curve did not reach a plateau level.

- 5 All compounds were tested in presence of mGluR2 agonist glutamate at a predetermined EC₂₀ concentration, to determine positive allosteric modulation. pEC₅₀ values were calculated from a concentration-response experiment of at least 8 concentrations.

B) In vivo pharmacologyMotor Activity (Video tracking)Apparatus and General Procedure

On the day of experiments, the mice were brought into the procedural room. They
5 were housed individually and allowed to acclimate for at least a half hour prior to
testing. Although the studies were conducted during the light cycle (from 8:00 to 16:00
h), the procedure room was only sparsely lit (3 to 30 LUX) to provide better contrast
for the video tracking. Local lighting was used for the injection procedures. During
each trial, an individual mouse was placed in an open field arena (grey PVC cylinder
10 with a height of 40 cm and a diameter of 22.5 cm). Each arena was placed on an
infrared LED (8 x 8 LEDs)-lit box (white PVC squared box; 40 x 40 cm²; height
12.5 cm). Each mouse was placed in the center of the arena and allowed to explore
freely for 30 min. After each trial, the arena was cleaned with a wet and subsequently
with a dry cleaning cloth. An infrared sensitive tube camera and a white light source
15 (in arena: 4-7 LUX) were mounted to the ceiling above the observation chamber to
record and input activity to a computer. Animal behavior was recorded and analyzed
using the Noldus Ethovision XT Video Tracking System (Version 3.1; Noldus,
Wageningen, The Netherlands). The total distance traveled (cm) was calculated. Data
were then exported to data management systems for further analysis and reporting.

20

1) Phencyclidine (PCP)-induced Hyperlocomotion in Mice

Test compound or solvent was administered at a pre-defined time before measurement
(standard: 30 min) to male NMRI mice that were challenged with phencyclidine (PCP;
5 mg/kg, s.c.) 30 min before measurement. Activity was measured for a period of 30
25 min. Criterion for drug-induced inhibition of hyperlocomotion: total distance < 5500
counts (3.9% false positives in controls; n = 154).

2) Conditioned avoidance response (CAR) test in Rats**Apparatus**

30 The apparatus consisted of an inner box surrounded by an outer box. The inner box
was composed of four walls of transparent, synthetic material (length x width x height:
30 x 30 x 30 cm), an open top, and a grid floor made of 15 pairs of iron bars (2 mm
diameter; 6 mm inter-bar distance). Odd and even bars were connected with a source
of alternative current (1.0 mA; Coulbourn Instruments Solid State Shocker/Distributor),
35 which could be interrupted by a switch. The outer box was composed of the same
material (length x width x height: 40 x 40 x 36 cm), also with an open top, with a

distance of 5 cm between the inner and outer box on all sides. To decrease the amount of environmental stimuli, three walls of the outer box were made non-transparent. The front wall was left transparent to allow the necessary inspection of the animal during the test. The upper edge of the outer and inner box served as a target for the rats on which to jump with fore- and hind-paws, respectively.

Avoidance Conditioning and Selection of Animals

From their arrival in the laboratory on the experimental day, male Wiga Wistar rats (230 ± 30 g) were housed in individual cages provided with bedding material. The rats received 5 training sessions at 15-min time intervals over a 1-h period during which, the rats were conditioned to avoid an electric shock: the rat was placed on the non-electrified grid floor and the grid was electrified 10 s later for not more than 30 s, if the rat did not jump out of the box. Only rats that showed correct avoidance responses in all the last 3 training sessions were included for further experiments, and received the test compound or solvent immediately after the last training session.

Experimental Sessions

The rats were tested 3 times, i.e. at 60, 90 and 120 min after the injection of test compound or solvent. Latency to avoidance was recorded. The median avoidance response obtained over the three experimental sessions for each rat were used for further calculations. A median avoidance latency > 8 s was selected as an all-or-none criterion for drug-induced inhibition of avoidance (occurring in only 1.5% of solvent-pretreated control rats; $n = 66$).

The results of tests 1) and 2) are shown in table 6 below.

Table 6. Pharmacological data for compounds according to the invention in tests 1) and 2) described above. PCP means Drug-induced inhibition of PCP-induced hyperlocomotion; CAR means Conditioned avoidance response; ED₅₀ means median effective dose; PO means oral route.

Co. No.	Route	ED ₅₀ (mg/kg)	
		PCP	CAR
6	PO	28.3	8.1*

* compound 6 was in suspension in 20% hydroxypropyl- β -cyclodextrin containing 1% polysorbate 80.

3) Anticonvulsant Studies

Preparation of Test Article and Controls

Test compounds were administered using an optimal fluid volume to body fluid ratio. Test compounds were administered to mice in a volume of 0.01 ml/g of body weight (White, H.S., et al., General principles: Experimental selection, quantification, and evaluation of antiepileptic drugs, in Antiepileptic Drugs, Fourth Edition, R.H. Levy, R.H. Mattson, and B.S. Meldrum, Editors. 1995, Raven Press, Ltd.: New York. p. 99-110). The test compound number 6 was administered orally (p.o.). For each of the tests performed on the test compound, a 40% Hydroxypropyl- β -cyclodextrin (Hp- β -CD) stock solution was first prepared and utilized for formulating the test compound number 6 at the desired concentrations for testing via the oral (p.o.) route. Final compound concentrations were administered as suspensions in 20% Hp- β -CD. A 20% Hp- β -CD solution was used for the vehicle groups.

15 *Critical Reagents*

a) Vehicle Solutions

40% Hydroxypropyl- β -cyclodextrin (Hp- β -CD) stock solution

b) Miscellaneous Solutions

20 Tetracaine (0.5% solution w/v) was added dropwise from a plastic dropper bottle onto the eyes of all animals that would subsequently receive electrical stimulation via corneal electrodes.

Animals and animal husbandry

Adult male CF No 1 albino mice (26-35 g) were obtained from Charles River, Portage, Michigan. The animals were maintained on an adequate diet (Prolab RMH 3000) and allowed free access to food and water, except during the short time they were removed from their cage for testing. Animals newly received in the laboratory were allowed sufficient time to correct for possible food and water restriction incurred during transit before being employed in testing. All mice were housed in plastic cages in specially constructed rooms with controlled humidity, exchange of air and controlled lighting (12 hours on - 12 hours off). The animals were housed, fed, and handled in a manner consistent with the recommendations in the National Council Publication, "Guide for the Care and Use of Laboratory animals."

Experimental Design

35 *General Methods*

Minimal Motor Impairment (MMI): Acute MMI was assessed by a combination of direct observations of the animal for overt symptoms of the animal's neurological or

muscular function. In mice, the rotarod procedure was used to disclose minimal muscular or neurological impairment. When a mouse is placed on a rod that rotates at a speed of 6 rpm, the animal can maintain its equilibrium for long periods of time. The animal was considered toxic if it fell off this rotating rod three times during a 1 min period.

TPE Determination: Groups of four animals each were administered test compounds and each group was tested at one of five time points: 0.25, 0.5, 1, 2, or 4h post-treatment (White et al. 1995). TPE was determined using the 6Hz (32mA) assay. The time (0.25, 0.5, 1, 2, or 4h post-treatment) at which maximal protection was observed was considered the Time of Peak Effect (TPE).

At the TPE determined for this study, or determined previously, compounds were tested in the 6Hz assay (32 and/or 44 mA), across several doses and comprising doses that elicited little or no protection to full protection. An ED₅₀ and 95% confidence interval (CI) were calculated using Probit analysis on a computer program provided in the laboratory (Finney "Probit Analysis" 34d ED 1971, London: Cambridge University Press).

Serum Collection for pK/pD Analysis: In various tests, animals were sacrificed following testing, and trunk blood and/or brain tissue (whole brains) was collected for quantification of drug levels. Immediately after testing, animals were decapitated and trunk blood was collected into a BD Vacutainer® tube containing K2EDTA and chilled on ice until centrifugation. Following centrifugation (13000 – 18000 rpm, 5-7 min), the plasma was removed and transferred to a labeled microcentrifuge tube and stored at -80 °C. For brain tissue collection, brains were removed immediately following decapitation and flash frozen. The frozen sample was placed in a labeled centrifuge tube and stored at -80 °C.

6 Hz Psychomotor Seizure Test in Mice

The 6Hz seizure test is used as a model of pharmacoresistant limbic seizures. The 6Hz seizure displays a resistance to phenytoin, carbamazepine, lamotrigine, and topiramate (Barton et al. "Pharmacological characterization of the 6 Hz psychomotor seizure model of partial epilepsy" Epilepsy Research 2001, Vol. 47, pp. 217-222).

Method for 6 Hz Psychomotor Seizure Test

Focal seizures were induced in mice via corneal stimulation (6Hz, 0.2 msec rectangular pulse, 3 sec duration; Barton et al. 2001). Mice were tested at either 32mA or 44mA. Prior to stimulation, drops of 0.5% tetracaine were applied to each eye. The seizures

that arise from corneal stimulation in this assay are characterized by a minimal clonic phase followed by stereotyped automatic behaviors including stun, forelimb clonus, twitching of the vibrissae, and Straub-tail. Animals not displaying these behaviors were considered protected.

5

Table 7: Time-to-Peak Effect Determination for Co. No. 6 (p.o.) in the 6Hz (32mA) Assay.

Dose (mg/kg, p.o.)	Time (h)	# protected / # tested	# rotarod motor impairment / # tested
10	0.25	1/4	0/4
	0.5	3/4	0/4
	1	0/4	0/4
	2	1/4	0/4
	4	0/4	0/4
20	0.25	4/4	0/4
	0.5	3/4	0/4
	1	4/4	0/4
	2	0/4	0/4
	4	1/4	0/4

TPE determined to be 0.5h.

10 **Table 8: Dose-Response Study for Co. No. 6 in the 6Hz Assay (32mA and 44mA; 0.5h TPE).**

Test	Dose (mg/kg, p.o.)	# protected / # tested	# rotarod motor impairment / # tested
6Hz 32mA	20	7/8	0/8
	10	6/8	0/8
	5	2/8	0/8
	2.5	1/8	0/8
ED₅₀ (95% CI): 7.2 mg/kg (4.2 to 11.8)			
6Hz 44mA	40	8/8	0/8

	20	6/8	0/8
	15	4/8	0/8
	10	0/8	0/8
ED₅₀ (95% CI): 16.1 mg/kg (13.0 to 20.1)			

Prophetic composition examples

“Active ingredient” as used throughout these examples relates to a final compound of Formula (I), the pharmaceutically acceptable salts thereof, the solvates and the stereochemically isomeric forms and the tautomers thereof.

5 Typical examples of recipes for the formulation of the invention are as follows:

1. Tablets

	Active ingredient	5 to 50 mg
	Di-calcium phosphate	20 mg
	Lactose	30 mg
10	Talcum	10 mg
	Magnesium stearate	5 mg
	Potato starch	ad 200 mg

In this Example, active ingredient can be replaced with the same amount of any of the compounds according to the present invention, in particular by the same amount of any
 15 of the exemplified compounds.

2. Suspension

An aqueous suspension is prepared for oral administration so that each 1 milliliter contains 1 to 5 mg of one of the active compounds, 50 mg of sodium carboxymethyl cellulose, 1 mg of sodium benzoate, 500 mg of sorbitol and water ad 1 ml.

20 3. Injectable

A parenteral composition is prepared by stirring 1.5 % by weight of active ingredient of the invention in 10% by volume propylene glycol in water.

4. Ointment

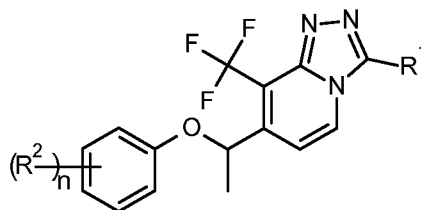
	Active ingredient	5 to 1000 mg
25	Stearyl alcohol	3 g
	Lanoline	5 g
	White petroleum	15 g
	Water	ad 100 g

In this Example, active ingredient can be replaced with the same amount of any of the compounds according to the present invention, in particular by the same amount of any of the exemplified compounds.

- Reasonable variations are not to be regarded as a departure from the scope of the
- 5 invention. It will be obvious that the thus described invention may be varied in many ways by those skilled in the art.

CLAIMS

1. A compound of Formula (I)



or a stereochemically isomeric form thereof, wherein

5 R^1 is selected from the group consisting of C_{1-6} alkyl, $(C_{3-8}$ cycloalkyl) C_{1-3} alkyl, and $(C_{1-3}$ alkyloxy) C_{1-3} alkyl;

each R^2 is independently selected from F, Cl, C_{1-3} alkyl, C_{1-3} alkyloxy, mono- or polyhalo C_{1-3} alkyl, and mono- or polyhalo C_{1-3} alkyloxy;

n is an integer selected from 1, 2, and 3;

10 or a pharmaceutically acceptable salt or solvate thereof.

2. The compound according to claim 1, or a stereoisomeric form thereof, wherein

R^1 is selected from the group consisting of CH_3CH_2 , $CH_3CH_2CH_2$,

(cyclopropyl)methyl, (cyclobutyl)methyl, ethyloxymethyl, and methyloxymethyl; and

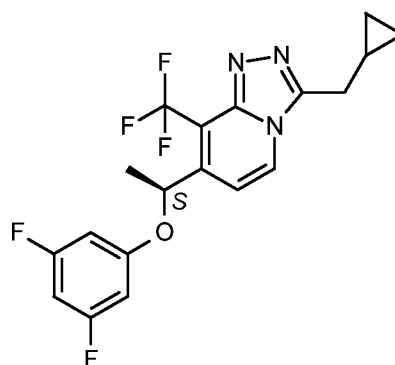
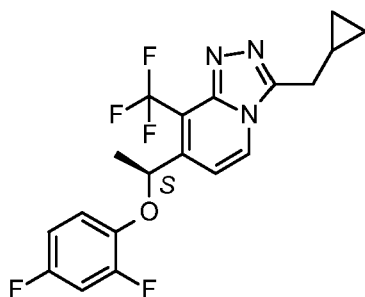
15 the rest of variables are as defined in claim 1.

3. The compound according to claim 1 or 2, wherein

each R^2 is independently selected from F, Cl, CH_3 , CH_3O and CF_3 ; and the rest of variables are as defined in claim 1 or 2.

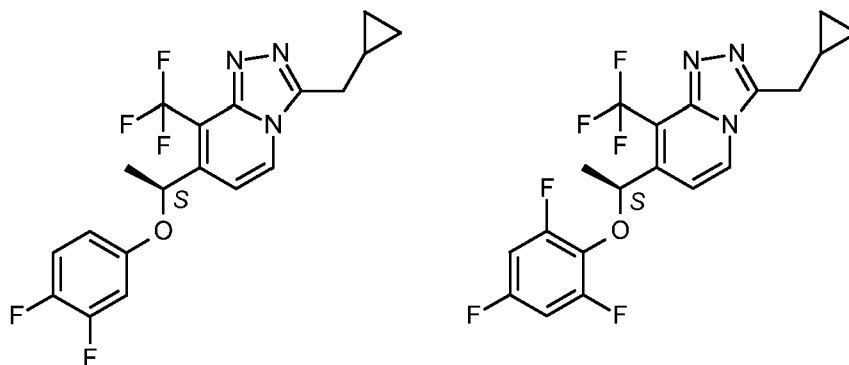
20

4. The compound according to any one of claims 1 to 3, wherein the compound is



; or a hydrochloride salt thereof;

;



; or

5. A pharmaceutical composition comprising a therapeutically effective amount of a compound according to any one of claims 1 to 4 and a pharmaceutically acceptable carrier or excipient.

5

6. A compound according to any one of claims 1 to 4 or a pharmaceutical composition according to claim 5, for use as a medicament.

7. A compound according to any one of claims 1 to 4 or a pharmaceutical composition according to claim 5 for use in the treatment or prevention of a central nervous system disorder selected from the group of anxiety disorders, psychotic disorders selected from the group consisting of schizophrenia, schizoaffective disorder and schizophreniform disorder, personality disorders, substance-related disorders, eating disorders, mood disorders, migraine, epilepsy or convulsive disorders, childhood disorders, cognitive disorders, neurodegeneration, autistic disorders, neurotoxicity and ischemia.

8. The compound or the pharmaceutical composition for use according to claim 7, wherein the psychotic disorders are selected from the group of schizophrenia, schizoaffective disorder and schizophreniform disorder;

the anxiety disorders are selected from the group of agoraphobia, generalized anxiety disorder (GAD), mixed anxiety and depression, obsessive-compulsive disorder (OCD), panic disorder, panic attack, posttraumatic stress disorder (PTSD), social phobia and other phobias;

the personality disorders are selected from the group of obsessive-compulsive personality disorder, borderline personality disorder and schizoid, schizotypal disorder;

the substance abuse or substance-related disorders are selected from the group of alcohol abuse, alcohol addiction, alcohol dependence, alcohol withdrawal, alcohol withdrawal delirium, alcohol-induced psychotic disorder, amphetamine addiction, amphetamine dependence, amphetamine withdrawal, cocaine addiction, cocaine dependence, cocaine withdrawal, nicotine addiction, nicotine dependence, nicotine withdrawal, opioid dependence and opioid withdrawal;

the eating disorders are selected from the group of anorexia nervosa and bulimia nervosa;

the mood disorders are selected from the group of bipolar disorders (I & II), cyclothymic disorder, depression, dysthymic disorder, major depressive disorder, treatment resistant depression, bipolar depression, and substance-induced mood disorder;

the epilepsy or convulsive disorders are selected from the group of generalized nonconvulsive epilepsy, generalized convulsive epilepsy, petit mal status epilepticus, grand mal status epilepticus, partial epilepsy with or without impairment of consciousness, infantile spasms, epilepsy partialis continua, and other forms of epilepsy;

the cognitive disorders are selected from the group of delirium, substance-induced persisting delirium, dementia, dementia due to HIV disease, dementia due to Huntington's disease, dementia due to Parkinson's disease, dementia of the Alzheimer's type, behavioral and psychological symptoms of dementia, substance-induced persisting dementia and mild cognitive impairment;

the autistic disorders are selected from autism and autism spectrum disorders, such as Asperger's syndrome.

25

9. A compound according to any one of claims 1 to 4 or a pharmaceutical composition according to claim 5 for use in the treatment or prevention of a central nervous system disorder selected from

anxiety disorders selected from the group of agoraphobia, generalized anxiety disorder, panic disorder, social anxiety disorder (social phobia), and panic attack;

schizophrenia spectrum and other psychotic disorders selected from the group of schizophrenia, schizoaffective disorder and schizophreniform disorder;

- personality disorders selected from the group of obsessive-compulsive personality disorder, borderline personality disorder, schizoid personality disorder, and schizotypal personality disorder;
- 5 substance-related and addictive disorders selected from the group of alcohol use disorder, alcohol withdrawal, opioid use disorder, opioid withdrawal, stimulant (amphetamine-type substance, cocaine) use disorder, stimulant (amphetamine-type substance, cocaine) withdrawal, tobacco use disorder, and tobacco withdrawal;
- 10 depressive disorders selected from the group of major depressive disorder, persistent depressive disorder (dysthymia), and substance/medication-induced depressive disorder;
- bipolar and related disorders selected from the group of bipolar I disorder, bipolar II disorder, cyclothymic disorder, substance/medication-induced bipolar and related disorder;
- obsessive-compulsive disorder;
- 15 trauma- and stressor-related disorders selected from the group of posttraumatic stress disorder, and acute stress disorder;
- neurodevelopmental disorders selected from the group of autism spectrum disorder, and attention-deficit/hyperactivity disorder;
- 20 neurocognitive disorders (NCDs) (both major and mild) selected from the group of delirium, substance intoxication delirium, NCD due to Alzheimer's disease, NCD due to HIV infection, NCD due to Huntington's disease, NCD due to Parkinson's disease, and substance/medication-induced NCD; and
- 25 epilepsy or convulsive disorders selected from the group of generalized nonconvulsive epilepsy, generalized convulsive epilepsy, petit mal status epilepticus, grand mal status epilepticus, partial epilepsy with or without impairment of consciousness, infantile spasms, epilepsy partialis continua, and other forms of epilepsy.
- 10.** The compound or the pharmaceutical composition for use according to claim 8 or 9, wherein the central nervous system disorder is selected from the group of
- 30 schizophrenia (in particular, negative symptoms or residual symptoms thereof), generalized anxiety disorder, bipolar disorder (I or II), migraine, behavioral and psychological symptoms of dementia, epilepsy or convulsive disorders, panic disorder, mixed anxiety and depression, and agoraphobia.

11. A product comprising
- (a) a compound as defined in any one of claims 1 to 4; and
 - (b) an mGluR2 orthosteric agonist,
- 5 as a combined preparation for simultaneous, separate or sequential use in the treatment or prevention of a central nervous system disorder selected from the group of schizophrenia (in particular, negative symptoms or residual symptoms thereof), generalized anxiety disorder, bipolar disorder (I or II), migraine, behavioral and psychological symptoms of dementia, epilepsy or convulsive disorders, panic disorder,
- 10 mixed anxiety and depression, and agoraphobia.