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(19) **HU****MAGYARORSZÁG**
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(73) Jogosult(ak):

Addex Pharma S.A, 1202 Genéve (CH)

(72) Feltaláló(k):

RIGUET, Eric, 1228 Plan-les-Ouates (CH)**CAMPO, Brice, 1228 Plan-les-Ouates (CH)****GIBELIN, Antoine, 1228 Plan-les-Ouates (CH)****MHALLA, Karim, 1228 Plan-les-Ouates (CH)**

(74) Képviselő:

SBGK Szabadalmi Ügyvivői Iroda, Budapest

(54)

GABA-B modulátor új triazindionszármazékok

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(54) **NOVEL TRIAZINEDIONE DERIVATIVES AS GABA-B RECEPTOR MODULATORS**

NEUE TRIAZINDION-DERIVATE ALS GABA-B-REZEPTOR-MODULATOREN

NOUVEAUX DÉRIVÉS DE TRIAZINEDIONE EN TANT QUE MODULATEURS DU RÉCEPTEUR GABA-B

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(73) Proprietor: **Addex Pharma S.A**

1202 Genève (CH)

(72) Inventors:

- **RIGUET, Eric**
1228 Plan-les-Ouates (CH)
- **CAMPO, Brice**
1228 Plan-les-Ouates (CH)
- **GIBELIN, Antoine**
1228 Plan-les-Ouates (CH)
- **MHALLA, Karim**
1228 Plan-les-Ouates (CH)

(74) Representative: **Davies, Jonathan Mark**

Reddie & Grose LLP
16 Theobalds Road
London WC1X 8PL (GB)

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- **DATABASE CA[Online] CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; BADAWEY, EL SAYED A. M. ET AL: "Synthesis of new benzimidazole derivatives as potential antimicrobial agents" XP002479872 retrieved from STN Database accession no. 1991:471532 & ARCHIV DER PHARMAZIE (WEINHEIM, GERMANY) , 324(6), 355-7 CODEN: ARPMAS; ISSN: 0365-6233, 1991,**

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J.F.KENNEDY,BP
1140,MONTLUCON,03103,FRANCE

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KENNEDY,BP
1140,MONTLUCON,03103,FRANCE

Description

SUMMARY OF THE INVENTION

5 [0001] I

[0002] The present invention provides novel compounds which are gamma amino butyric acid receptor-subtype B ("GABA_B"), positive allosteric modulators (enhancers), which are useful for treating or preventing diseases or disorders, including treatment of anxiety, depression, epilepsy, schizophrenia, cognitive disorders, spasticity and skeletal muscle rigidity, spinal cord injury, multiple sclerosis, amyotrophic lateral sclerosis, cerebral palsy, neuropathic pain and craving associated with cocaine and nicotine, panic disorder, posttraumatic stress disorders, urge urinary incontinence, gastro-esophageal reflux disease, transient lower oesophageal sphincter relaxations, functional gastrointestinal disorders and irritable bowel syndrome.

BACKGROUND OF THE INVENTION

15 [0003] The amino acid GABA (γ -aminobutyric acid) is the main inhibitory neurotransmitter in the adult mammalian brain and regulates many physiological and psychological processes. GABA acts through two major classes of receptors: ionotropic GABA_A (including GABA_C) receptors and metabotropic GABA_B receptors (Hill and Bowery, Nature 1981, 290, 149-152; Bormann, J., Trends Pharmacol. Sci. 2000, 21, 16-19). GABA_B receptors are present in most regions of the mammalian brain on presynaptic terminals and postsynaptic neurons, and are involved in the fine-tuning of inhibitory synaptic transmission. Due to their strategic position in neuronal networks to modulate the activity of the various neurotransmitter systems, GABA_B receptors are a target of choice for pharmacological agents intended to treat central and peripheral nervous systems disorders (Bettler et al., Physiol Rev. 2004, 84, 835-867; Cryan and Kaupmann, Trends Pharmacol Sci. 2005, 26, 36-43).

25 [0004] The GABA_B receptor belongs to the Class-III family of G protein-coupled receptors (GPCRs), as are the receptors for glutamate, Ca²⁺, pheromones and putative taste compounds (Pin et al., Pharmacol. Ther. 2003, 98, 325-354). All members of this family share the characteristic of a large extracellular amino-terminal domain that contains a so-called "Venus Flytrap" orthosteric ligand binding site and seven transmembrane (7TM) helical segments plus an intracellular carboxyl-terminal domain that are involved in receptor activation and G-protein coupling (Bockaert and Pin, EMBO J. 1999, 18, 1723; Galvez et al, J. Biol.Chem. 1999, 274, 13362-13369). A distinct feature, however, of the GABA_B receptor is that it operates as a heterodimer of at least two homologous subunits termed GABA_{B1} and GABA_{B2} (Kaupman et al, Nature 1997, 386, 239-246; Gordon et al, J. Biol.Chem. 1999, 274, 7607-7610; Margeta-Mitrovic et al, PNAS 2001, 98, 14649-14654; Bettler et al, Physiol Rev. 2004, 84, 835-867). Orthosteric GABA_B receptor ligands bind only at the N-terminal Venus flytrap region of the GABA_{B1} subunit, which in turn activates the associated GABA_{B2} subunit of the heterodimer. It is this latter subunit that is responsible for coupling and activation of G-protein (Galvez et al, EMBO J. 2001, 20, 2152-2159; Duthey et al, J. Biol. Chem. 2002, 277, 3236-3241; Pin et al, Biochem. Pharmacol. 2004, 68, 1565-1572). The resulting effect is an inhibition of the adenylyl cyclase activity and subsequent cyclic AMP formation and the modulation of activity of inwardly rectifying potassium channels and voltage-sensitive calcium channels.

35 [0005] Several studies using knock-out (KO) mice have demonstrated the role of the heterodimeric GABA_{B1B2} receptor in several CNS disorders. Mice lacking the GABA_{B1} subunit exhibit spontaneous seizures and hyperalgesia (Schuler et al, Neuron 2001, 31, 47-58). These behavioral characteristics are paralleled by a loss of all biochemical and electrophysiological GABA_B responses in these KO mice. In these studies, a clear impairment of passive avoidance performance was also observed indicating impaired memory processes. GABA_{B1} deficient mice were also found to be more anxious than their wild-type counterparts (Mombereau et al, Neuropsychopharmacology 2004, 29, 1050-1062). Analogous results were obtained with GABA_{B2} KO mice, which presented all the same behavioural characteristics than the one observed for the GABA_{B1} KO mice (Gassman et al, J. Neurosci. 2004, 24, 6086-6097). Moreover, it has also been shown that a hypoactivity of the GABA system was linked to spasticity, epilepsy, anxiety, stress, sleep disorders, depression, addiction, and pain (Dalvi and Rodgers, Psychopharmacology 1996, 128, 380-397; for a recent review Ong and Kerr, CNS Drug Dev. 2005, 11, 317-334); while on the contrary, a hyperactivity of the GABAergic system was associated with schizophrenia (Blum and Mann, Int. J. Neuropsychopharmacol. 2002, 5, 159-179).

45 [0006] Baclofen is a potent and selective agonist at the GABA_B receptor and is presently a frequently and only used clinical drug in the treatment of spasticity and rigidity (Montane et al, Neurology 2004, 63, 1357-1363). Moreover, all effective pharmacological agents used to treat panic disorder increase GABA synaptic transmission and anxiolytics and antidepressants that lack GABA activity are not effective in panic disorders. Baclofen was shown to be significantly effective in reducing the number of panic attacks and symptoms of anxiety as assessed with the Hamilton anxiety scale, Zung Scale, and Katz-R nervousness subscale (Breslow et al, Am. J. Psychiatry 1989, 146, 353-356). Drake and co-workers, hypothesized that baclofen would be an effective treatment in the symptomatic management of veterans with chronic posttraumatic stress disorder (PTSD). Their results demonstrated that the therapy, well tolerated, resulted in

significant improvements of the overall symptoms of PTSD and co-morbid depression and anxiety in patients with chronic PTSD due to combat (Drake et al, Ann. Pharmacother. 2003, 37, 1177-1181). More recently, a study looking at the effect of baclofen on the prepulse inhibition (PPI) of the acoustic startle response (ASR), proposed GABA_B receptors as putative new targets in the pharmacological therapy of psychotic disorders (Bortalo et al, Psychopharmacology 2004, 171, 322-330). Despite the demonstration of baclofen as being a potential therapeutic tool for the treatment of disorders such as anxiety and spasticity, its use has been limited due to its poor blood-brain-barrier penetration, very short duration of action, muscle relaxing property, hypothermic and sedative side effects, as well as patients' increasing tolerance (Hefferan et al, Neuroscience Letters 2006, 403, 195-200).

[0007] A new avenue for developing selective compounds acting at GPCRs is to identify molecules that act through allosteric mechanisms, modulating the receptor by binding to a site different from the highly conserved orthosteric binding site. This concept has assumed a greater importance in the pharmacology of family III receptors in general. For example, allosteric modulators have been described for Ca²⁺-sensing receptors (Nemeth et al, USP 6,031,003), for metabotropic glutamate receptors (reviewed in Mutel, Expert Opin. Ther. Patents 12:1-8, 2002; Ritzen, Mathiesen and Thomsen, Basic Clin. Pharmacol. Toxicol 97:202-13, 2005), and most recently for GABA_B receptors (Urwyl et al, Mol. Pharmacol. 2001, 60, 963-971; WO 2005/094828; WO 2006/001750; WO 2006/063732; WO 2006/048146; WO 2006/07486; WO 2006/136442; WO 2007/014843; US 2007/027204; WO 2007/073297; WO 2007/073298; WO 2007/073299; WO 2007/073300). These ligands do not activate the receptor by themselves, but, for example in the case of the GABA_B receptor, increase the potency of GABA in the presence of this endogenous agonist (Pin et al, Mol. Pharmacol. 2001, 60, 881-884; Urwyl et al, Neuropharmacol. 2005, 48, 343-353). Mutational analyses have demonstrated unequivocally that the binding of known GABA_B receptor positive allosteric modulators does not occur at the orthosteric site, but instead at an allosteric site within the seven transmembrane region of the GABA_{B2} subunit at least for CGP7930 (Binet et al, J. Biol. Chem. 2004, 279, 29085-29901).

[0008] As a therapeutic principle, positive allosteric modulators are expected to have several advantages over compounds acting as orthosteric agonists, because they are only effective in the presence of the endogenous ligand and therefore act in line with physiological neurotransmission in its temporal and spatial organization. Orthosteric agonists, on the other hand, activate receptors independently of synaptic activity, possibly leading to unwanted side effects.

[0009] Cryan and co-workers suggested in a recent study, using GS39783 (N,N'-Dicyclopentyl-2-methylsulfanyl-5-nitro-pyrimidine-4,6-diamine), that a positive modulation of GABA_B receptors may serve as a novel therapeutic strategy for the development of anxiolytics with a better side effect profile as compared to baclofen (Cryan et al, J. Pharm. Exp. Therap. 2004, 310, 952-963). They showed that GS39783 is active in models of anxiety such as elevated plus maze (rat), elevated zero maze (mice and rats), and the stress-induced hyperthermia (mice) tests. Moreover, as expected for a positive allosteric modulator that do not have any effect on receptor activity in absence of GABA, but do enhance allosterically the affinity of the GABA_B receptor for the endogenous GABA, no side effect on locomotor activity, rotarod, body temperature and traction test was observed for doses ranging from 0.1 to 200 mg/kg, p.o. In comparison baclofen presented those side effects even at efficacious doses in anxiety models. In conclusion, those data suggest that GS39783, and positive allosteric modulators of GABA_B receptors in general, are useful and innovative anxiolytics without side-effects associated with baclofen. An interesting example of the use of such compounds in preclinical studies have been done for CGP7930 and GS39783 for the treatment of Gastro-Esophageal Reflux Disease (GERD) and cocaine self-administration in rats, where those compounds were found active (Smith et al, Psychopharmacology 2004, 173, 105-111; WO 03/090731).

[0010] Recently the 3,3'-diarylpropyl-1-arylethylamines and 3-aryl propyl-1-arylethylamine have been reported as a new class of GABA_B receptor modulators as they potentiate baclofen-induced responses in the brain (Kerr et al, Aust. J. Chem. 2006, 59, 445-456) and they modulate both pre- and postsynaptic GABA_B receptors in rat brain slices (Ong et al, Eur. J. Pharm. 2005, 507, 35-42).

[0011] Patent Publications DE10255416 and WO9730980 describe triazinediones derivatives having herbicidal properties. These triazinediones are also described in the field of fungicide (EP0438717), growth of protozoa (EP0232932, WO00006172), binding to PPAR- α and - γ (FR02866339), P2X7 inhibitors (US25288288, WO04058270) and IL-5 production inhibitors (WO09902505).

[0012] US 2004/0192692 and WO01/14377 disclose 1,2,4-triazine compounds as ligands for GABA_A receptors, which are useful for treating disorders of the central nervous system.

[0013] It has now surprisingly been found that the compounds of the invention show potent activity and selectivity on GABA_B receptor.

FIGURE

[0014]

Figure A represents the ability of 10 μ M of the Compound **4.02** to increase the [GTP γ ³⁵S] binding in rat brain

membranes induced by 15 μ M of Baclofen, a GABA_B receptor agonist, without having an effect on its own.

DETAILED DESCRIPTION OF THE INVENTION

[0015] According to the present invention, there are provided compounds of the general Formula III, as defined in the attached claims.

[0016] Further preferred compounds of the present invention are compounds selected from the compounds listed in the attached claims.

[0017] The present invention relates to the pharmaceutically acceptable acid addition salts of the compounds and compositions of the compounds with pharmaceutically acceptable carriers or excipients.

[0018] The present invention provides compounds/compositions for use in treating or preventing a condition in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of GABA_B positive allosteric modulators.

[0019] The present invention provides compounds/compositions for use in treating or preventing peripheral and central nervous system disorders selected from the group consisting of: anxiety, depression, epilepsy, schizophrenia, cognitive disorders, spasticity and skeletal muscle rigidity, spinal cord injury, multiple sclerosis, amyotrophic lateral sclerosis, cerebral palsy, neuropathic pain and craving associated with cocaine and nicotine, panic disorder, posttraumatic stress disorders, urge urinary incontinence or gastro-intestinal disorders.

[0020] Pharmaceutical compositions of the invention may provide from about 0.01 to 1000 mg of the active ingredient per unit dose. The compositions may be administered by any suitable route. For example orally in the form of capsules, etc., parenterally in the form of solutions for injection, topically in the form of ointments or lotions, ocularly in the form of eye-drops, rectally in the form of suppositories, intranasally or transcutaneously in the form of delivery system like patches.

[0021] The pharmaceutical formulations of the invention may be prepared by conventional methods in the art; the nature of the pharmaceutical composition employed will depend on the desired route of administration. The total daily dose usually ranges from about 0.05 - 2000 mg.

Definitions

[0022] The following terms and expressions contained herein are defined as follows:

As used herein, the term **"about"** refers to a range of value from $\pm$ 10% of a specified value. For example, the phrase "about 30 mg" include $\pm$ 10% of 30, or from 27 to 33 mg.

[0023] As used herein, a range of values in the form **"x-y"** or **"x to y"** or **"x through y"**, include integers x, y and the integers there between. For example, the phrase "1-6" or "1 to 6" or "1 through 6" are intended to include the integers 1, 2, 3, 4, 5 and 6. Preferred structures include each individual integer in the range, as well as any subcombination of integers. For example, preferred integers for "1-6" can include 1, 2, 3, 4, 5, 6, 1-2, 1-3, 1-4, 1-5, 2-3, 2-4, 2-5 or 2-6, etc.

[0024] As used herein, **"stable compound"** or **"stable structure"** refers to a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture and preferably capable of formulation into an efficacious therapeutic agent.

[0025] As used herein, the specification **"C"** means 1 carbon atom.

[0026] As used herein, in the case where a subscript is the integer **"0"** (zero), the group to which the subscript refers to indicates that the group is absent, i.e. there is a direct bond between the groups.

[0027] **"Halogen"** includes atoms such as fluorine, bromine, chlorine and iodine.

[0028] As used herein, the term **"alkyl"** refers to a straight-chain or branched alkyl group having 1 to 6 carbon atoms, such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, 1-ethylpropyl, 3-methylpentyl, 2,2-dimethylbutyl, 2,3-dimethylbutyl, hexyl, etc. The alkyl moiety of alkyl-containing groups, such as alkoxy, alkoxycarbonyl and alkylaminocarbonyl groups, has the same meaning as alkyl defined above.

[0029] As used herein, the term **"alkenyl"** refers to a straight-chain or branched hydrocarbon group of 2 to 6 carbon atoms having at least one carbon-carbon double bond. A designation "C₂-C₆ alkenyl" refers to an alkenyl radical containing from 2 to 6 carbon atoms. Examples of alkenyl groups include, but are not limited to, ethenyl, propenyl, isopropenyl, allyl, butenyl, pentenyl, 2,4-pentadienyl, etc.

[0030] As used herein, the term **"alkynyl"** refers to a straight-chain or branched hydrocarbon group of 2 to 6 carbon atoms having at least one carbon-carbon triple bond. A designation "C₂-C₆ alkynyl" refers to an alkynyl radical containing from 2 to 6 carbon atoms. Examples include, but are not limited to, ethynyl, propynyl, butynyl, pentynyl, etc.

[0031] The term **"Aryl"** refers to an optionally substituted monocyclic or bicyclic hydrocarbon ring system containing at least one unsaturated aromatic ring. Examples and suitable values of the term "aryl" are phenyl, naphthyl, 1,2,3,4,-tetrahydronaphthyl, indyl, indenyl and the like.

[0032] "**Arylalkyl**" includes (C₆-C₁₀)aryl-(C₁-C₃)alkyl group such as 2-ethyl-phenyl group, 3-ethyl-phenyl group, 4-propyl-phenyl group, 2-propyl-phenyl group, 3-propyl-phenyl group, 2-methyl-naphtyl group, 4-methyl-naphtyl group or the like.

[0033] "**Alkylaryl**" includes (C₁-C₃)alkyl-(C₆-C₁₀)aryl group such as benzyl group, as phenyl-ethyl group, phenyl-propyl, naphtyl-2-methyl group or the like.

[0034] In this specification, unless stated otherwise, the term "**alkylhalo**" means an alkyl group as defined above, substituted with one or more halogen. The term "C₁-C₆-alkylhalo" may include, but not limited to fluoroethyl, difluoromethyl, trifluoromethyl, fluoroethyl, difluoroethyl, bromoethyl and the like. The term "O-C₁-C₆-alkylhalo" may include, but is not limited to fluoromethoxy, difluoromethoxy, trifluoromethoxy, fluoroethoxy and the like.

[0035] "**Heteroatom**" includes atoms such as nitrogen, oxygen and sulphur.

[0036] In the specification, unless stated otherwise, the term "**Heteroaryl**" refer to an optionally substituted monocyclic or bicyclic unsaturated, aromatic ring system containing at least one heteroatom selected independently from O, N or S to form a ring such as furyl (furan ring), benzofuranyl (benzofuran ring), thienyl (thiophene ring), benzothiophenyl (benzothiophene ring), pyrrolyl (pyrrole ring), imidazolyl (imidazole ring), pyrazolyl (pyrazole ring), thiazolyl (thiazole ring), isothiazolyl (isothiazole ring), triazolyl (triazole ring), tetrazolyl (tetrazole ring), pyridil (pyridine ring), pyrazynyl (pyrazine ring), pyrimidinyl (pyrimidine ring), pyridazinyl (pyridazine ring), indolyl (indole ring), isoindolyl (isoindole ring), benzoimidazolyl (benzimidazole ring), purinyl group (purine ring), quinolyl (quinoline ring), phtalazinyl (phtalazine ring), naphtyridinyl (naphtyridine ring), quinoxalinyl (quinoxaline ring), cinnolyl (cinnoline ring), pteridinyl (pteridine ring), oxazolyl (oxazole ring), isoxazolyl (isoxazole ring), benzoxazolyl (benzoxazole ring), benzothiazolyl (benzothiazole ring), furazanyl (furazan ring), benzotriazolyl (benzotriazole ring), imidazopyridinyl (imidazopyridine ring), pyrazolopyridinyl (pyrazolopyridine ring), 2,3-dihydrobenzofuranyl (2,3-dihydrobenzofuran ring) and the like.

[0037] "**Heteroarylalkyl**" includes heteroaryl-(C₁-C₃)alkyl group, wherein examples of heteroaryl are the same as those illustrated in the above definition, such as 2-methyl-furyl group, 3-methyl-furyl group, 2-methyl-thienyl group, 3-methyl-thienyl group, 1-methyl-imidazolyl group, 2-methyl-imidazolyl group, 2-methyl-thiazolyl group, 2-methyl-pyridyl group, 3-methyl-pyridyl group, 1-methyl-quinolyl group or the like.

[0038] "**Alkylheteroaryl**" includes (C₁-C₃)alkyl-heteroaryl group, wherein examples of heteroaryl are the same as those illustrated in the above definition, such as 2-furylmethyl group, 3-furylmethyl group, 2-thienylmethyl group, 3-thienylmethyl group, 1-imidazolylmethyl group, 2-imidazolylmethyl group, 2-thiazolylmethyl group, 2-pyridylmethyl group, 3-pyridylmethyl group, 1-quinolylmethyl group or the like.

[0039] In this specification, unless stated otherwise, the term "**heterocyclic**" (or "**heterocyclo**") means a 5- or 6-membered ring containing one or more atoms independently selected from N, O, or S, includes monocyclic or bicyclic rings as well which may be saturated or unsaturated.

[0040] "**Heterocycloalkyl**" includes heterocycloalkyl-C₁-C₃-alkyl group, wherein examples of heterocyclo are the same as those illustrated in the above definition such as morpholine, piperazine, methylpiperazine, piperidine, dioxane and the like.

[0041] The term "**cycloalkyl**" means carbocycles containing no heteroatoms, includes mono-, bi-, and tricyclic saturated carbocycles, as well as fused ring systems. Such fused ring systems can include on ring that is partially or fully unsaturated such as a benzene ring to form fused ring systems such as benzo fused carbocycles. Cycloalkyl includes such fused ring systems as spirofused ring systems. Examples of cycloalkyl include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, decahydronaphthalene, adamantane, indanyl, fluorenyl, 1,2,3,4-tetrahydronaphthalene and the like.

[0042] The term "**cycloalkenyl**" refers to a carbocycles includes mono- and bi-carbocycles, as well as fused ring, having at least one carbon-carbon double bond. Examples of cycloalkenyl includes 1,2,3,4,4a,5,6,7-octahydronaphthalene, 2,3,5,6,7,7a-hexahydro-1H-indene, 2,3,4,5,6,7-hexahydro-1H-indene and the like.

[0043] "**Solvate**" refers to a complex of variable stoichiometry formed by a solute (e.g. a compound of formula I) and a solvent. The solvent is a pharmaceutically acceptable solvent as water preferably; such solvent may not interfere with the biological activity of the solute.

[0044] "**Optionally**" means that the subsequently described event(s) may or may not occur, and includes both event(s), which occur, and events that do not occur.

[0045] The term "**substituted**" refers to substitution with the named substituent or substituents, multiple degrees of substitution being allowed unless otherwise stated.

SYNTHESIS

[0046] Compounds may be prepared by methods known in the art of organic synthesis as set forth in part by the following synthesis schemes. In all of the schemes described below, it is well understood that protecting groups for sensitive or reactive groups are employed where necessary in accordance with general principles of chemistry. Protecting groups are manipulated according to standard methods of organic synthesis (T.W. Green and P.G.M. Wuts (1991) Protecting Groups in Organic Synthesis, John Wiley et Sons). These groups are removed at a convenient stage of the

compound synthesis using methods that are readily apparent to those skilled in the art. The compounds may be represented as a mixture of enantiomers, which may be resolved into the individual pure *R*- or *S*-enantiomers. If for instance, a particular enantiomer of the compound is desired, it may be prepared by asymmetric synthesis, or by derivation with a chiral auxiliary, where the resulting diastereomeric mixture is separated and the auxiliary group cleaved to provide the pure desired enantiomers. Alternatively, where the molecule contains a basic functional group such as amino, or an acidic functional group such as carboxyl, this resolution may be conveniently performed by fractional crystallization from various solvents, of the salts of the compounds with optical active acid or by other methods known in the literature, e.g. chiral column chromatography.

Resolution of the final product, an intermediate or a starting material may be performed by any suitable method known in the art as described by E.L. Eliel, S.H. Wilen and L.N. Mander (1984) *Stereochemistry of Organic Compounds*, Wiley-Interscience.

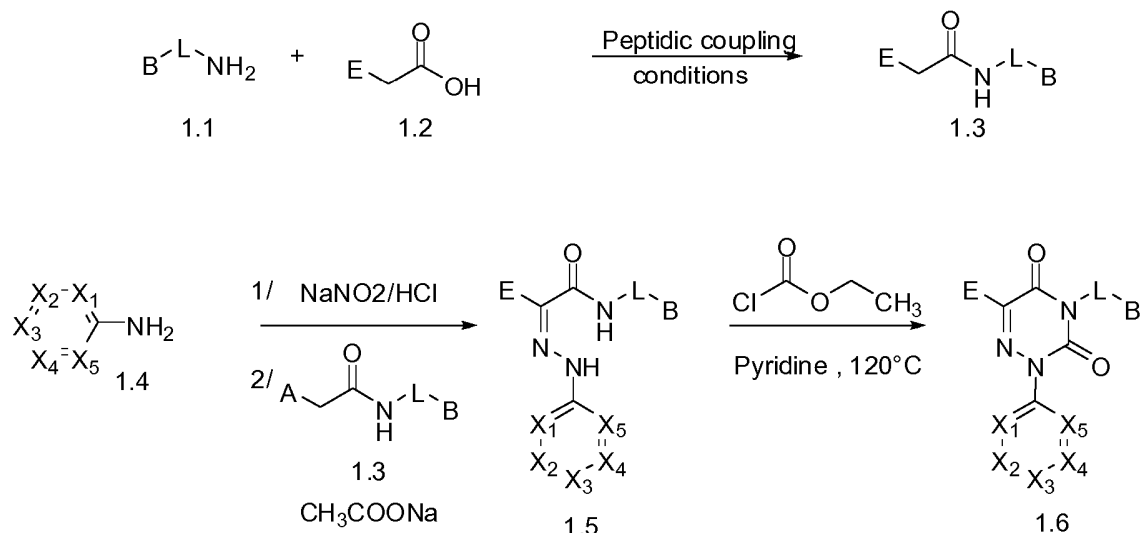
[0047] All the products generated from the following reactions can be isolated and purified employing standard techniques, such as extraction, chromatography, crystallization, distillation, and the like.

[0048] General routes to prepare the Examples of the present invention are shown in the Schemes and Examples that follow.

[0049] All substituents in the synthetic Schemes unless otherwise indicated, are as defined in the attached claims.

[0050] The compounds of Formula III wherein B, L, X₁, X₂, X₃, X₄, X₅ and -E are as defined in the attached claims may be prepared following a cyclisation reaction key step presented in Scheme 1. The amine **1.1** react with the corresponding carboxylic **1.2** acid in a classic peptidic coupling condition (Piskunova, I.P. & al. *Tetrahedron* 1993, 49(21), 4671-4676) where -E can be different electroattractive substituents, to generate **1.3**. Then **1.3** can react with the diazonium salt already obtained by mixing the aniline **1.4** with NaNO₂ to generate **1.5** (Bilek, P.; Slouka, J. *Heterocycl. Commun.* 2004, 10(1), 67-70). The final compound **1.6** is obtained by reaction of the intermediate **1.5** with ethylchloroformate in a presence of pyridine.

Scheme 1



[0051] Alternatively the E substituent can be introduced at a late stage in the synthesis, to provide the compounds of Formula III (Scheme 2). Thus the uracyl derivative is brominated to lead to the intermediates **2.1** (Dudfield, P.J. & al. *J. Chem. Soc. Perkin Trans. 1*, 1999, 20, 2929-2936). The introduction of the L-B group can be achieved through alkylation of **2.2** with Z-L-B (Z= Br, Cl, I, Mesylate, Tosylate, Triflate, OH) in presence of a suitable base such as K₂CO₃, NaH or using a Mitsunobu condition (Z= OH) (Pontillo, J. & al. *Bioorg. Med. Chem. Lett.* 2005, 15(19), 4363-4366) to afford the intermediate **2.3**. Introduction of the suitable substituted aryl/heteroaryl boronic acid is achieved in a presence of copper salt (Lam, P. Y. S. & al. *Tetrahedron Lett.* 2001, 42, 3415-3418) to afford the compound **2.4**. Displacement of the bromoderivative **2.4** by suitable nucleophiles such as amine alcohol in conditions known to the skilled in the art and exemplified in this document lead to compound of formula **2.5** where E=(OR¹², NR¹³R¹⁴), G=(H or a counterion). Additionally reaction of **2.4** with organometallics such as organozinc organoboron, organotin reagent in a presence of palladium catalyst and a co-solvent (M is a metal, zinc or boron or tin or silicon) afford **2.5** whereas E= (alkylaryl, alkyl,

heteroaryl, aryl and the like).

Scheme 2

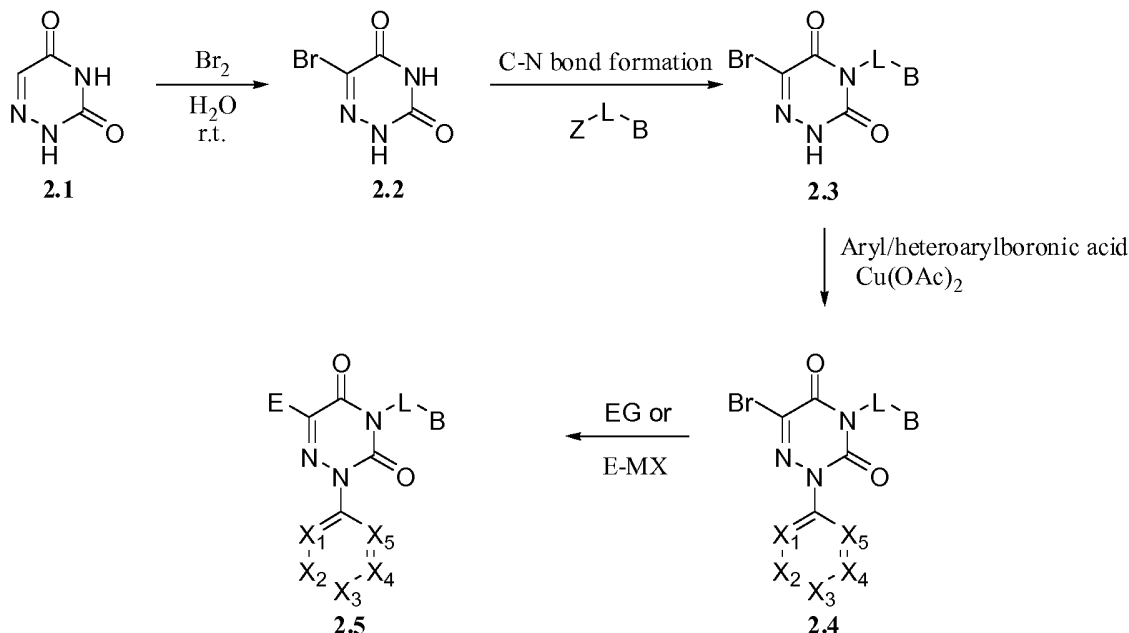
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[0052] Yet another approach to the cpds of Formula III can be represented with the introduction of the L-B group at the final stage of the synthesis (Scheme 3). To achieve such procedure bromouracyl is protected using a suitable protecting group (**P.G.**) such as benzyl or allyl by reaction of **2.2** with benzyl bromide to afford the intermediate **3.1**. The introduction of the suitable substituted aryl/heteroaryl boronic acid is the same as reported in previous scheme to afford **3.2**. The displacement of the bromoderivative was done as reported in Scheme 2 to afford **3.3**. Deprotection of the amide moiety can be achieved with palladium or ruthenium catalyst in presence of nucleophile such as acetate in a manner well known to those skilled in the art. The introduction of the L-B group proceeds in a similar manner described in Scheme 2 to afford the compound of Formula III (ie **3.5**).

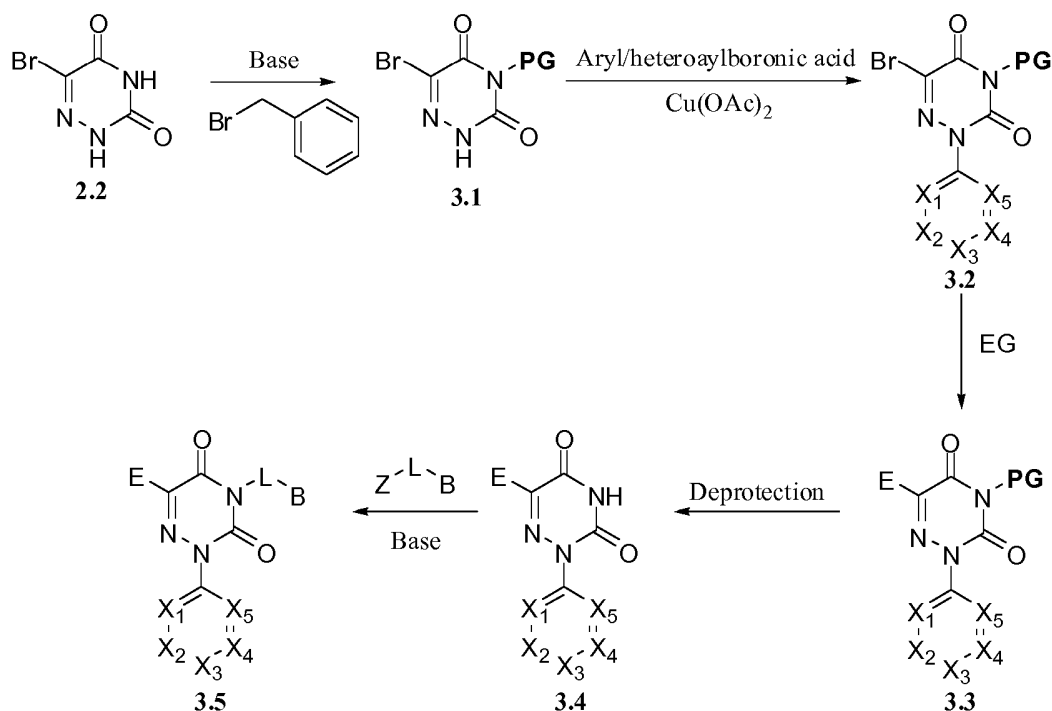
Scheme 3

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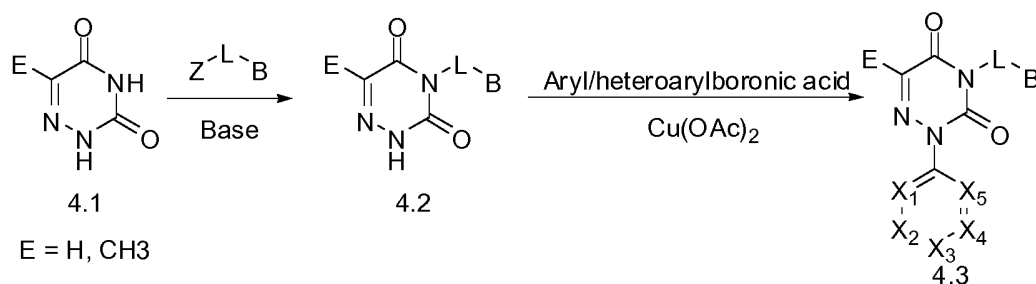
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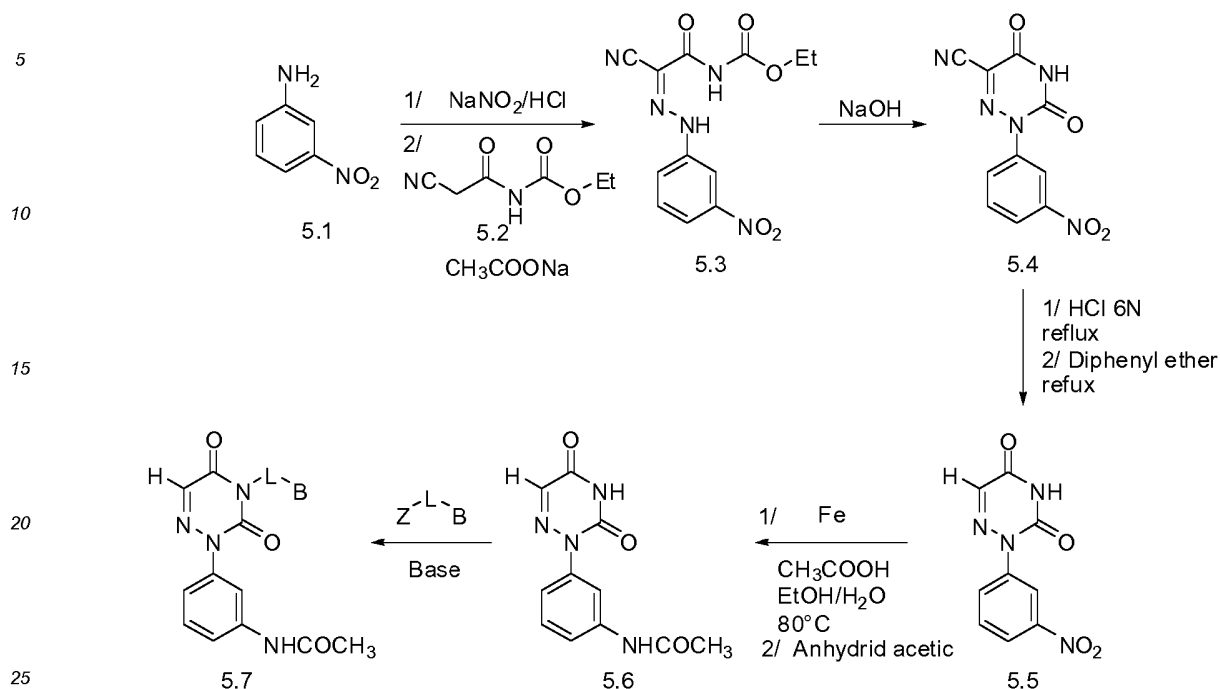
[0053] A similar procedure as those described in Scheme 2 and 3 can be used starting from commercially available uracyl **4.1** (E = H or CH₃). The introduction of the L-B group can be achieved through alkylation of **4.1** with Z-L-B (Z = Br, Cl, I, Mesylate, Tosylate, Triflate, OH) in presence of a suitable base such as K₂CO₃, NaH or using a Mitsunobu condition (Z = OH) (Pontillo, J. & al. Bioorg. Med. Chem. Lett. 2005, 15(19), 4363-4366) to afford the intermediate **4.2**. The introduction of the suitable substituted aryl/heteroaryl boronic acid is the same as reported in a Scheme 2 to afford compound of Formula III (i.e. **4.3**).

Scheme 4



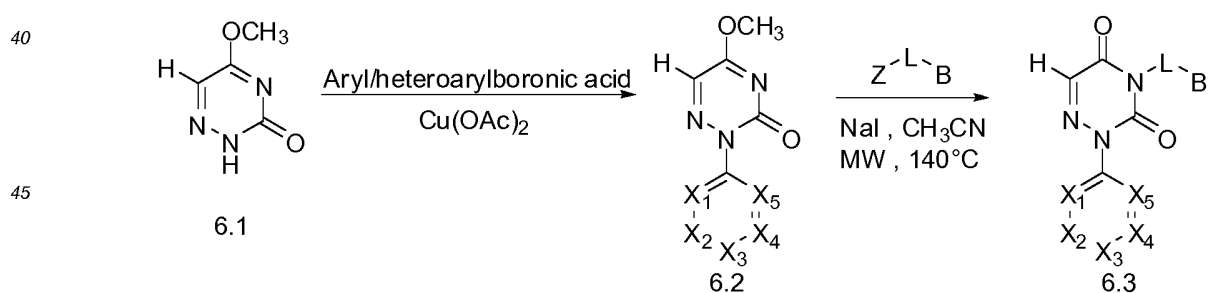
[0054] Yet another approach to the compound of Formula III is described in Scheme 5 in particular for those compounds of Formula III for which the introduction of the aryl or heteroaryl through Cham Lam coupling is difficult. In those cases the azalactam ring already was bearing the aryl substituent. **5.4** is built from the open intermediate **5.3** which is cyclised in basic medium. The intermediate **5.3** was obtained by condensation of a diazonium salt of suitably substituted aniline (in this case R⁶ = NO₂) with ethyl 2-cyanoacetylcarbamate. Further transformation of the E substituent when E = CN in compound of Formula III can be performed (see also Scheme 7 & 8). Thus decyanation was achieved via a two step procedure known in the art such as hydrolysis to the acid and decarboxylation of the acid bearing intermediate (Falco & al. J. Am. Chem. Soc. 1956, 78, 1938-1939). By extension further transformation of the R⁵ group can be also performed such as transformation of the nitro group to an acetamido group. The target compound is then obtained by introduction of the L-B group in a manner similar as that reported in previous to afford the compound **5.7**.

Scheme 5



[0055] Additional possibility of introducing the L-B group at a final stage of the synthesis of compound of Formula III, proceed through clivative alkylation procedure (Scheme 6). Thus **6.1** (described in Collect. Czechoslov. Chem. Commun. 1961, 26, 986-997; Collect. Czechoslov. Chem. Commun. 1975, 40, 2326-2339) is subjected to the Cham Lam coupling in order to introduce the aryl or heteroarylboronic acid in a manner similar to that described in previous schemes to afford **6.2**. The clivative alkylation procedure can be performed with a Z-L-B (Z= Br, Cl, I, Mesylate, Tosylate, Triflate,) in presence of sodium iodide (NaI) under microwave irradiation to afford compounds of Formula III (i.e **6.3**).

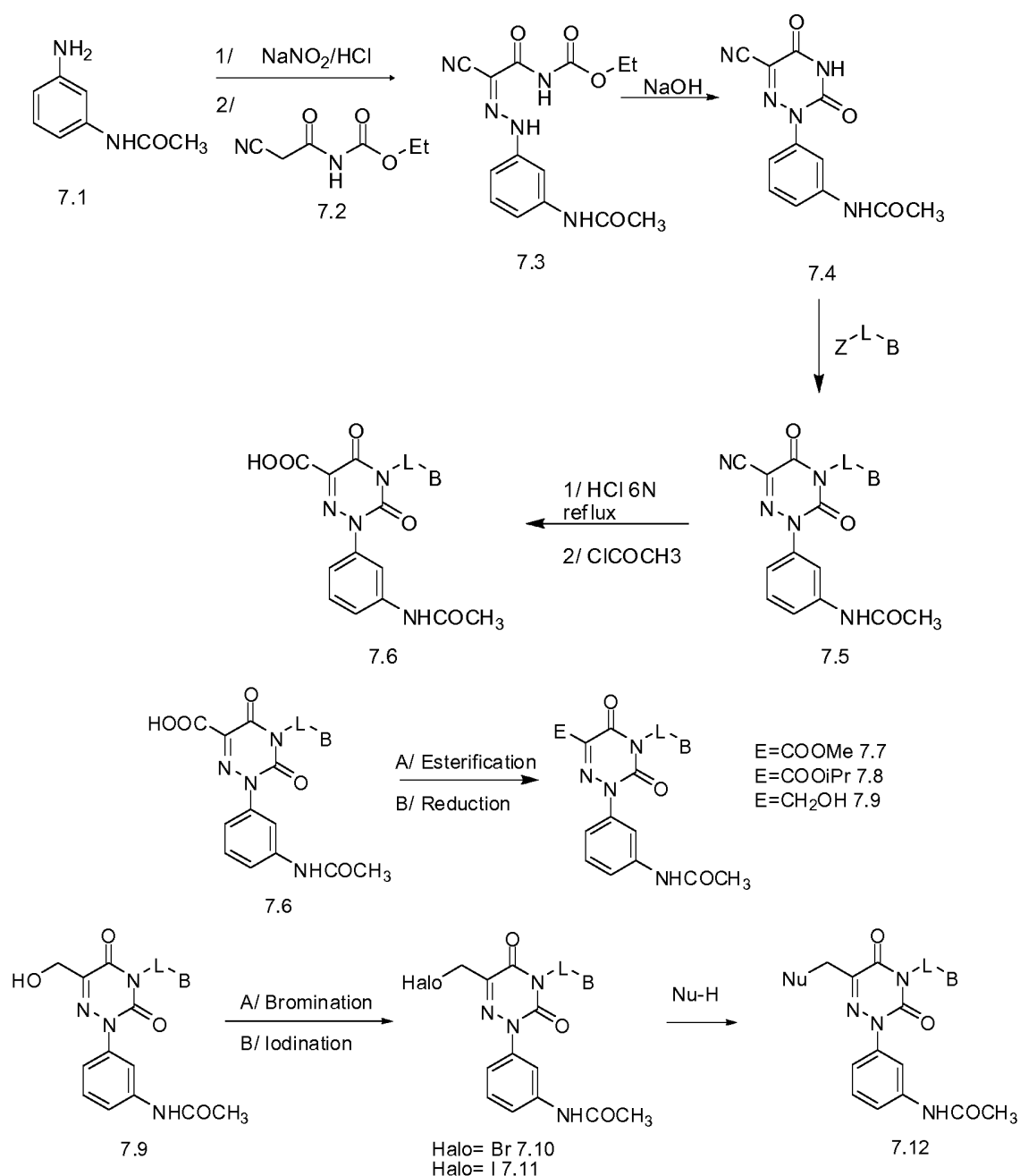
Scheme 6



[0056] The compounds of Formula III wherein B, L, X₁, X₂, X₃, X₄, X₅ are as defined in the attached claims and E= COOH is an important intermediate may be prepared following one variation presented Scheme 7. The commercially available aniline **7.1** is able to react with sodium nitrite in acidic media to generate a diazo intermediate which is able to react with the commercially available compound **7.2** to form the compound **7.3**. In a basic condition, the cyclisation of **7.3** can occur to form the disubstituted azauracil **7.4**. This intermediate can react with different halogenalkylaryl (Z= Br, Cl, I, Mesylate, Tosylate, Triflate) in a presence of base to obtain the compound **7.5**. After refluxing the compound **7.5** in a presence of HCl 6N the nitrile function is transformed into a carboxylic acid function and in a same time the deacetylation of the acetamido is realised. The compound **7.6** can be obtained after acetylation of the aniline. The carboxylic acid function can be transformed in ester function (Colletti, S.L. & al. Tetrahedron Lett. 2000, 41(41), 7825-7830) and can

be reduce in a presence of sodium borohydride to get the alcohol. This alcohol can then react with the phosphorus(III)bro-
mide (Rastelli, G. Bioorg. Med. Chem. Lett. 1997, 7(14), 1897-1902) or with sodium iodide (Baldwin, J.E. Tetrahedron
1991, 47(24), 4089-4100) to afford the bromo or iodo derivative. These halo derivatives could then react with different
nucleophile (**Nu-H**) to obtain the compound **7.12**.

Scheme 7

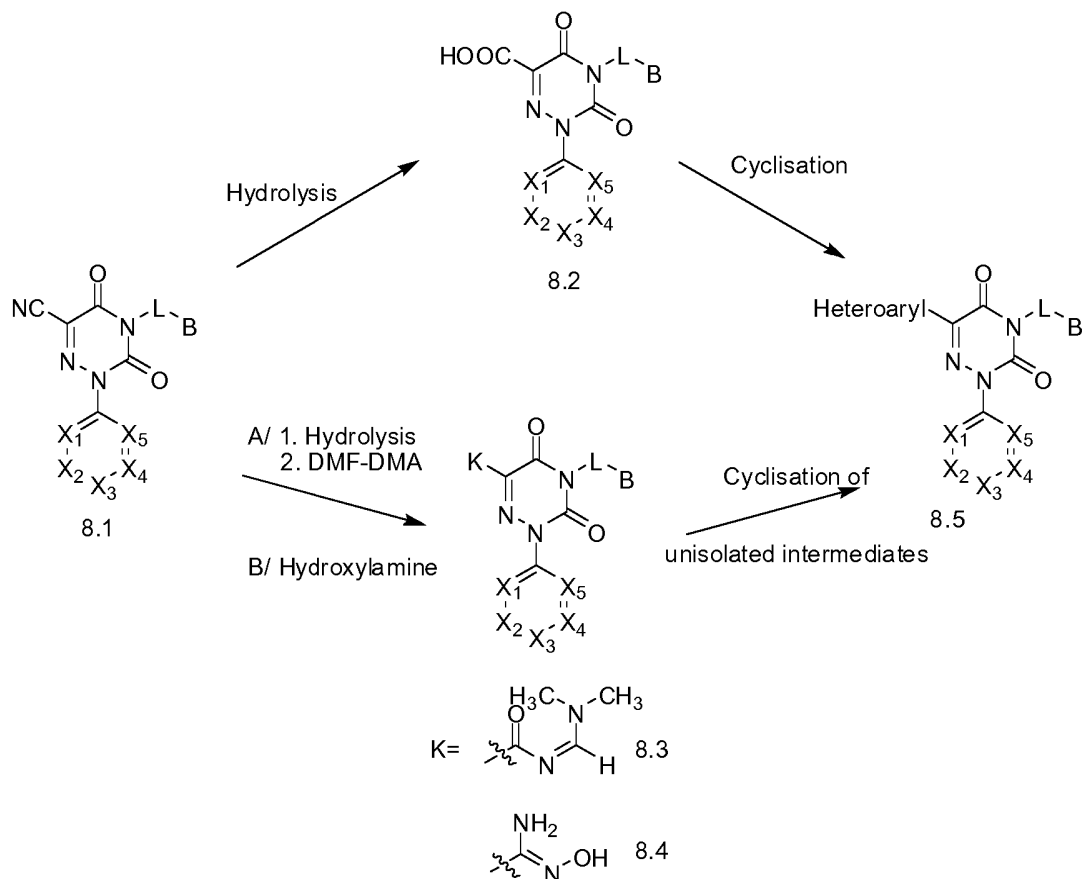


[0057] The compounds of Formula III wherein B, L, X₁, X₂, X₃, X₄, X₅ are as defined in the attached claims and E= CN is an important intermediate for further reactions. Some transformations of the nitrile organic function are presented in Scheme 8. Thus, derivative **8.1** obtained following the Scheme 1 can be hydrolysed into the corresponding carboxylic acid in conditions described in previous schemes. The acid function can itself be the starting for the formation of several

heteoaryl groups in manner known to the skilled in the art and are supported in literature in books (Katrizky A.R. and Rees C.W. (1984) Comprehensive Heterocyclic Chemistry, Pergamon Press), but not limited to. In particular treatment of the acid with hydroxyacetamidine followed by cyclisation of the intermediate acylamidoxine will lead to the the suitable substituted oxadiazole. Alternatively, the nitrile function can be easily transformed into precursor of heterocycles, i.e

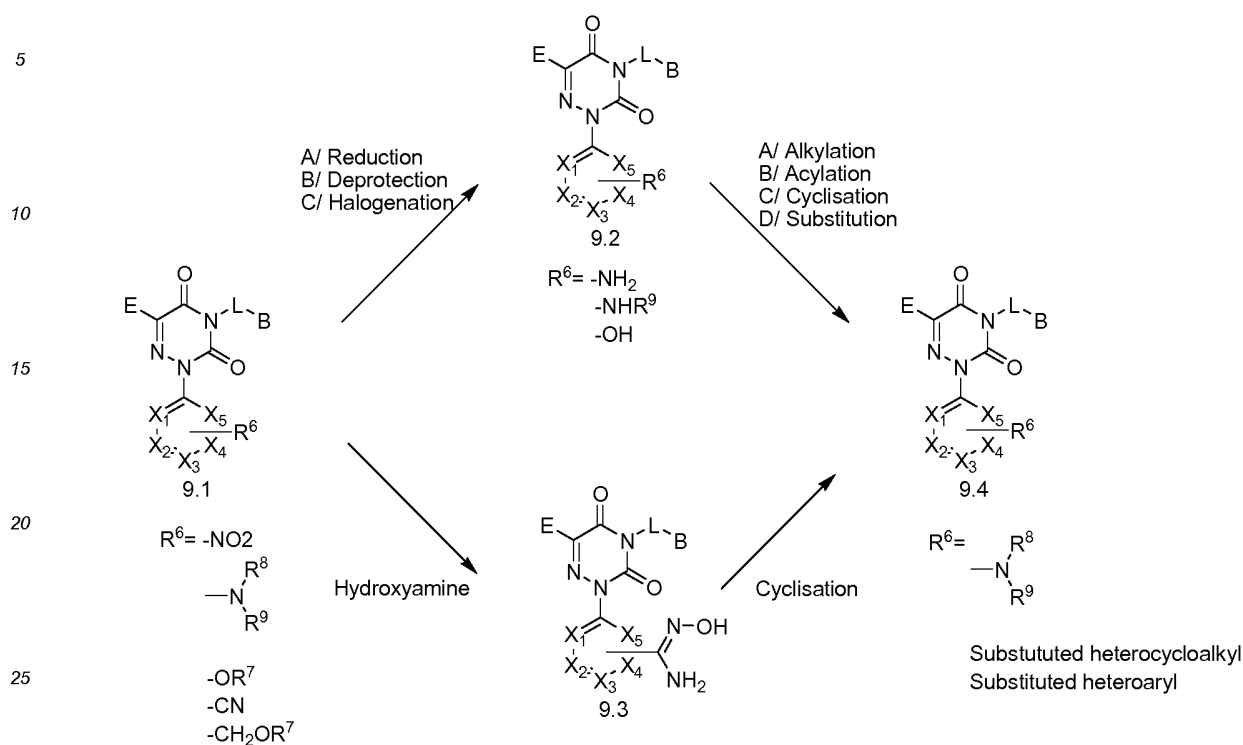
5 N-((dimethylamino)methylene)acetamide (two steps : tranformation to acylamide and reaction with DMF-DMA) or N-hydroxyformimidamide (in direct reaction with suitable hydroxylamine).

Scheme 8



[0058] The compounds of Formula III wherein B, L, X₁, X₂, X₃, X₄, X₅, E are as defined in the attached claims and R⁶ is NO₂, NR⁸R⁹, OR⁷ or CN could give compounds of Formula III where R⁶ was transformed into -NR⁸R⁹, substituted-heteroaryl or substitutedheterocycloalkyl are presented in Scheme 9. Thus, derivative 9.1, with R⁶ is -NO₂ can be reduced in primary amine function. The amine function can itself be the starting for the formation of several amide by acylation reaction, secondary or tertiary amine by alkylation reaction and substituted heterocycloalkyl groups in manner known to the skilled in the art and are supported in literature in books (Katrizky A.R. and Rees C.W. (1984) Comprehensive Heterocyclic Chemistry, Pergamon Press), but not limited to. When derivative 9.1, with R⁶ is NR⁸R⁹ can be reduce or deprotected to give the corresponding primary amine, secondary amine which can react as describe before. With R⁶ is -OR⁷, the deprotection reaction can occur, i.e with BBr₃ to give the alcohol. Then when R⁶ is CN the reactions already describe in the Scheme 8 can be realised to finally conduct to heteroaryl.

Scheme 9



[0059] The reagents and starting materials are commercially available and/or, using well-known techniques, can be readily synthesized by one of ordinary skill in the art. Unless otherwise noted, all commercially starting materials were used without further purification.

[0060] Specifically, the following abbreviations may be used in the examples and throughout the specification.

g (grams)	Rt (retention time)
mg (milligrams)	MeOH (methanol)
mL (milliliters)	NaCl (Sodium chloride)
μL (microliters)	Hz (Hertz)
M (molar)	LCMS (Liquid Chromatography Mass Spectrum)
MHz (megahertz)	HPLC (High Pressure Liquid Chromatography)
mmol (millimoles)	NMR (Nuclear Magnetic Resonance)
min (minutes)	^1H (proton)
AcOEt (ethyl acetate)	LiAlH_4 (lithium aluminium hydride)
$\text{Pd}(\text{PPh}_3)_4$ (tetrakis(triphenylphosphine)palladium(0))	MgSO_4 (magnesium sulphate)
$\text{PdCl}_2(\text{PPh}_3)_2$ (Bis(triphenylphosphine) palladium (II) dichloride)	TLC (tin layer chromatography)
CDCl_3 (deuterated chloroform)	HOBT (1-hydroxybenzotriazole)
EDCI.HCl (1-(3-(Dimethylaminopropyl)-3-ethylcarbodiimide, hydrochloride)	R.T. (Room Temperature)

(continued)

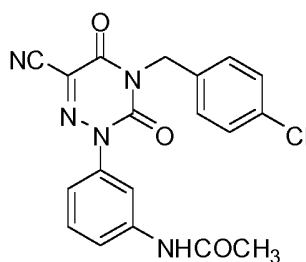
Et ₂ O (diethyl ether)	NaOH (sodium hydroxide)
% (percent)	h (hour)
DCM (dichloromethane)	HCl (hydrochloric acid)
DIEA (diisopropyl ethyl amine)	n-BuLi (n-butyllithium)
Mp (melting point)	THF (tetrahydrofuran)
NaHCO ₃ (sodium hydrogenocarbonate)	NH ₄ Cl (ammonium chloride)
H ₂ SO ₄ (Sulfuric acid)	N ₂ (nitrogen)
DMAP (N,N-dimethylaminopyridine)	DAST (diethylaminosulfur trifluoride)
(BOC) ₂ O (Di-tert-butyl dicarbonate)	mm (millimeters)
SiO ₂ (silicon(IV)oxyde)	V (volt)
°C (Celsius degrees)	MW (Microwave irradiation)
s (singulet)	bs (broad singulet)
d (doublet)	dd (doublet of doublet)
t (triplet)	dt (doublet of triplet)
m (multiplet)	td (triplet of doublet)
q (quintuplet)	dq (doublet of quintuplet)
OTf (triflate)	OMs (mesylate)
OTs (tosylate)	

[0061] All references to brine refer to a saturated aqueous solution of NaCl unless otherwise indicated. All temperatures are expressed in °C. All reactions are not conducted under an inert atmosphere at room temperature unless otherwise noted.

[0062] The microwave oven used is an apparatus from Biotage (Optimizer™) equipped with an internal probe that monitors reaction temperature and pressure, and maintains the desired temperature by computer control.

Reference Example 1: N-(3-(4-(4-chlorobenzyl)-6-cyano-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 1.01 : Compound 1.01 with E = -CN is not a compound of general Formula III as defined in the attached claims).

[0063]



1(A): N-(4-chlorobenzyl)-2-cyanoacetamide.

[0064] According to the Scheme 1 Step 1: A mixture of 4-chlorobenzylamine (10.0g, 71mmol), cyanoacetic acid (6.13g, 72mmol), HOBT (11g, 72mmol), EDCI.HCl (20g, 106mmol) in 100mL of dioxane was stirred for 3h at room temperature. The solvent was removed under vacuum and 300mL of DCM was added. The organic layer was successively washed with 2x100mL of water, 2x50mL of a solution 1N of NaOH, 4x100mL of water until neutral pH. The organic layer was dried over Na₂SO₄, filtered and concentrated to afford N-(4-chlorobenzyl)-2-cyanoacetamide **1(A)** (13.12g, 89%) as a

brown solid.

1(B): (Z)-N-(4-chlorobenzyl)-2-cyano-2-(2-(3-aminophenyl) acetamide)hydrazono)acetamide

[0065] According to the Scheme 1 Step 2: To a suspension of N-(3-aminophenyl)acetamide (1.5g, 10mmol) in concentrated HCl (2.9mL) at 0°C. was added dropwise a solution of sodium nitrite (661mg, 10mmol in 10mL of water); the obtained diazonium salt solution was added under vigorous stirring to a suspension of compound **1(A)** (1.0g, 5mmol) and sodium acetate (3.0g) in ethanol (25mL). The reaction mixture was allowed to rest at 0°C for 2h. The mixture was filtered, washed with cooled ethanol and dried to afford the hydrazone **1(B)** (1.2g, 68%) as yellow crystals.

1(C): N-(3-(4-(4-chlorobenzyl)-6-cyano-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0066] According to the Scheme 1 Step 3: A mixture of the compound **1(B)** (500mg, 1.4mmol) and 3mL of pyridine was cooled at 0°C. Onto this suspension was added dropwise 293mg (2.7mmol) of ethylchloroformiate, with a syringe. The reaction mixture was heated at 120°C for 6h and concentrated under vacuum. The residue was successively dissolved in 3mL of DCM, washed twice with a solution of 5N HCl (1mL), dried over sodium sulfate, filtrated and concentrated to dryness. The crude oil was purified by flash chromatography over Silica gel (AIT Flashmart prepacked column 10g SiO₂) using DCM/MeOH 98/2 as eluant to afford a yellow solid. The solid was washed with diethylether, filtered and dried to afford N-(3-(4-(4-chlorobenzyl)-6-cyano-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **1(C)** (192mg, 36%) as a yellow solid.

mp : 196°C

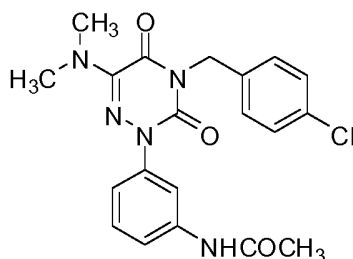
LC₁: Rt= 4.35

MS m/z (ES) [M-H]⁻ = 394

¹H-NMR (DMSO-d₆), δ (ppm): 10.21 (1H, bs), 7.89-7.88 (1H, m), 7.60-7.57 (1H, m), 7.47-7.39 (5H, m), 7.16-7.13 (1H, m), 4.99 (2H, m), 2.05 (3H, s).

Example 2: N-(3-(4-(4-chlorobenzyl)-6-(dimethylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 5.13).

[0067]



2(A): 6-bromo-1,2,4-triazine-3,5(2H,4H)-dione.

[0068] According to Scheme 2 Step 1: To a stirred suspension of 2g of 6-azauracyl (17.7mmol) and water (15mL) was added 2mL of bromine (38.9mmol) at room temperature. The mixture was stirred for 48h at room temperature and filtered. The crude solid was washed with water and dried to afford **2(A)** as a white powder (2.3g, 11% yield).

LC₁ : Rt= 0.69-1.53

MS m/z (ES) [M-H]⁻= 189.9, 191.9

2(B): 4-(4-chlorobenzyl)-6-bromo-1,2,4-triazine-3,5(2H,4H)-dione.

[0069] According to Scheme 2 Step 2: to a solution of 4-chlorobenzyl bromide (1g, 5.2mmol) and **2(A)** (1.1g, 5.2mmol) in DMSO (10mL) was added portionwise NaH(55%) (230mg, 5.2mmol). After 15min at room temperature the reaction mixture was quenched with water and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water (x2), with brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 50gr column, (Eluant: DCM/MeOH 98/2) to obtain 0.856g of 4-(4-chlorobenzyl)-6-bromo-1,2,4-triazine-3,5(2H,4H)-dione **2(B)**.

LC₁ : Rt= 3.93

MS m/z (ES) [M-H]⁻ = 315.8

2(C): N-(3-(4-(4-chlorobenzyl)-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0070] According to Scheme 2 Step 3: To a stirred solution of 4-(4-chlorobenzyl)-6-bromo-1,2,4-triazine-3,5(2H,4H)-dione (0.400g) and 2(B) (411mg, 1.3mmol) in DMF (5mL) were successively added of the 3-acetamidophenylboronic acid (0.34g, 1.9mmol), copper(II) acetate (0.23g, 1.3mmol) and 0.20g of pyridine (2.5mmol). The mixture was stirred overnight at 60°C. The reaction mixture was quenched with a saturated sodium bicarbonate solution and extracted with ethyl acetate (x3). The combined organic layers were washed successively with water, brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by C₁₈ reverse phase column, 65g column: 20% CH₃CN gradient 20 to 60% to afford 0.350g (yield= 62%) of N-(3-(4-(4-chlorobenzyl)-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **2(C)**.

TLC (DCM/MeOH 95/5): rf= 0.35

LC₁: Rt= 4.30

MS m/z (ES) [M+H]⁺ = 450.7

mp= 202°C

RMN 1H (300MHz, DMSO-d₆), δ (ppm): 10.17 (1H, s), 7.83 (1H, s), 7.60-7.54 (1H, s), 7.46-7.37 (5H, m), 5.0 (2H, s), 2.05 (3H, s).

2(D): N-(3-(4-(4-chlorobenzyl)-6-(dimethylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl) acetamide.

[0071] According to Scheme 2 Step 4: To a solution of **2(C)** (100mg, 0.22mmol) in 0.5mL of DMF was added dimethylamine 2M solution in methanol (0.56mL, 1.11mmol). The reaction was heated 5h at 150°C under microwave irradiation. The reaction mixture was quenched with water and extracted with ethyl acetate (x3). The combined organic layers were washed successively with water, with brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 5g column (Eluant: DCM/MeOH 98/2). 10mg, (11% yield) of N-(3-(4-(4-chlorobenzyl)-6-(dimethylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **2(D)** were obtained.

TLC (DCM/MeOH 95/5): rf= 0.44

LC₁: Rt= 4.39

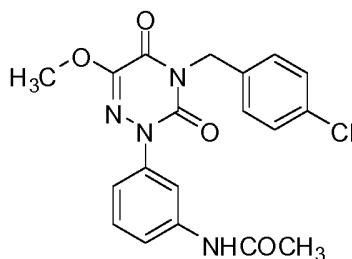
MS m/z (ES) [M+H]⁺ = 414.0

mp= 189-190°C

RMN 1H (300MHz, DMSO-d₆), δ (ppm): 10.07 (1H, s), 7.87 (1H, s), 7.53-7.49 (1H, m), 7.38-7.23 (6H, m), 5 (2H, s), 2.95 (6H, s), 2.04 (3H, s).

Example 3: N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 4.01).

[0072]



[0073] According to Scheme 2 Step 4: To a cooled solution of N-(3-(4-(4-chlorobenzyl)-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **2(C)** (80mg, 0.18mmol) in 0.5mL of methanol (0.5mL) at 0°C, was added a solution of sodium methoxide (2M) in Methanol (85μL 0.18mmol). After 1h at 0°C the reaction mixture was allowed to reach room temperature. After 1h the reaction mixture was quenched with water and extracted with ethyl acetate (x3). The combined organic layers were washed successively with water, with brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 5g column, (Eluant: DCM/AcOEt/MeOH 80/18/2). 25mg (35% yield) of N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide were recovered.

TLC (DCM/MeOH 95/5): rf= 0.36

LC₁: Rt= 3.98

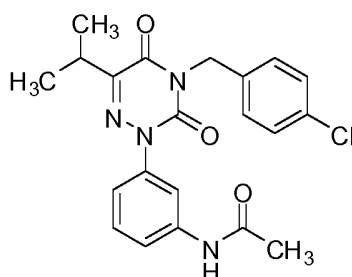
MS m/z (ES) [M+H]⁺= 400.9

mp= 190-191°C

RMN 1H (300MHz, DMSO-d₆), δ (ppm): 10.11 (1H, s), 7.87 (1H, s), 7.56-7.53 (1H, m), 7.43-7.34 (5H, m), 7.25-7.21 (1H, m), 5 (2H, s), 3.82 (3H, s), 2.05 (3H, s).

Example 4: N-(3-(4-(4-chlorobenzyl)-6-isopropyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 5.01).

[0074]



[0075] According to Scheme 2 Step 4: A mixture of N-(3-(4-(4-chlorobenzyl)-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **2(C)** (100mg, 0.22mmol), THF (2mL), dimethoxyethane (69μl, 0.67mmol) and PdCl₂(dppf) (8mg, 0.011mmol) was stirred under nitrogen atmosphere at 50°C. Isopropylzinc(II) chloride solution 0.58M (1.15mL, 0.67mmol) was added dropwise and the mixture was stirred at 50°C during 1hour. The reaction mixture was diluted with 100mL of water and extracted twice with 100mL of CH₂Cl₂. The combined organic layers were dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 15g SiO₂) using AcOEt/cyclohexane 4/6 as eluant to afford a brown solid. The solid was washed with diisopropylether and was filtered to afford N-(3-(4-(4-chlorobenzyl)-6-isopropyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (53mg, 58%) as a white solid.

LC₂: Rt= 2.82

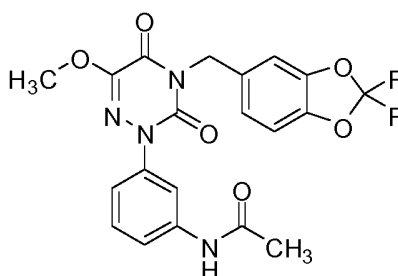
MS m/z (ES) [M+H]⁺= 413.0

mp= 178°C

¹H-NMR (DMSO-d₆), δ (ppm): 10.2 (1H, bs), 7.93 (1H, s), 7.67 (1H, dt), 7.5 (5H, m), 7.3 (1H, dt), 5.11 (2H, s), 3.26 (1H, m), 2.16 (3H, s), 1.27 (6H, d).

Example 5: N-(3-(4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 4.127).

[0076]



5(A): 6-Bromo-4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-1,2,4-triazine-3,5(2H,4H)-dione.

[0077] According to Scheme 2 Step 2: A mixture of 6-bromo-1,2,4-triazine-3,5(2H,4H)-dione **2(A)** (943mg, 4.91mmol), CH₂Cl₂ (40mL), triphenylphosphine (1.76g, 6.69mmol), (2,2-difluorobenzo[d][1,3]dioxol-5-yl)methanol (840mg, 4.46mmol) was stirred at 0°C under nitrogen atmosphere. Di-tertbutyl azodicarboxylate (2.05g, 8.93mmol) was added portionwise and the mixture was stirred for 16hours at room temperature. 200mL of water were added and the aqueous layer was extracted thrice with 200mL of DCM. The organic layers were combined and were successively dried over

MgSO₄, filtered and concentrated under reduce pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepac column SiO₂) using AcOEt/cyclohexane 3/7 as eluant to afford 6-bromo-4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-1,2,4-triazine-3,5(2H,4H)-dione **5(A)** (330mg, 20%) as a white solid.

LC₁: Rt= 2.44
MS m/z (ES) [M-H]⁻ = 362

5(B): N-(3-(6-bromo-4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0078] According to Scheme 2 Step 3: A mixture of 6-bromo-4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-1,2,4-triazine-3,5(2H,4H)-dione (300mg, 0.83mmol), DMF (5mL), 3-acetamidophenylboronic acid (222mg, 1.24mmol), copper acetate (150mg, 0.83mmol) and pyridine (135μL, 1.66mmol) was stirred at 65°C for 15 hours under room atmosphere. A saturated solution of NaHCO₃ (200mL) was added and the aqueous layer was extracted thrice with 200mL of DCM. The organic layers were combined to be dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (AIT Flashmart 25g prepac column C₁₈) using water/acetonitrile 6/4 as eluant to afford N-(3-(6-bromo-4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (250mg, 61%) as a white solid.

LC₁: Rt= 2.68
MS m/z (ES) [M-H]⁻ = 495

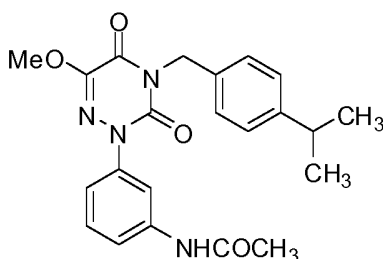
5(C): N-(3-(4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0079] According to Scheme 2 Step 4: A mixture of N-(3-(6-bromo-4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (250mg, 0.51mmol), DMF (6mL) was stirred at 0°C under nitrogen atmosphere. A sodium methoxide solution 2M (303μL, 0.61mmol) was added dropwise and the mixture was stirred at 0°C during 15min. The reaction mixture was diluted with 100mL of a saturated solution of NH₄Cl and the aqueous layer was extracted twice with 100mL of CH₂Cl₂. The organic layers were combined and successively dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepac column 15g SiO₂) using CH₂Cl₂/MeOH 98/2 as eluant to afford a beige solid. The solid was washed with diethylether and filtered to afford N-(3-(4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **5(C)** (141mg, 63%) as a white solid.

mp= 184°C
LC₂: Rt= 2.47
MS m/z (ES) [M+H]⁺ = 447
RMN 1H (300MHz, DMSO-d₆), δ (ppm): 10.12 (1H, bs), 7.89 (1H, t), 7.54 (1H, d), 7.39 (3H, m), 7.24 (1H, d), 5.03 (2H, s), 3.83 (3H, s), 2.06 (3H, s).

Example 6: N-(3-(4-(4-isopropylbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 4.36).

[0080]



6(A): 4-allyl-6-bromo-1,2,4-triazine-3,5(2H,4H)-dione.

[0081] According to Scheme 3 Step 1: To a solution of 6-bromoazauracil (2.00g, 10.4mmol) and allyl bromide (1.26g,

10.4mmol) in DMSO (25mL) was added portionwise NaH (55%) (0.45g, 10.4mmol) at room temperature,. The reaction mixture was stirred during 1h. 100mL of water were added and the aqueous layer was extracted thrice with ethyl acetate. The combined organic layers were successively washed twice with water, saturated brine, dried over MgSO₄, filtered and concentrated under reduce pressure. The crude solid was purified by flash chromatography with silica gel using CH₂Cl₂/MeOH 98/2 as eluant to afford 4-allyl-6-bromo-1,2,4-triazine-3,5(2H,4H)-dione **6(A)** (1.10g, 45%) with a purity of 90% (containing 10% of starting material) as a colorless oil.

LC₂: Rt= 1.40

MS m/z (ES) [M-H]⁻= 230

6(B):N-(3-(4-allyl-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0082] According to Scheme 3 Step 2: A mixture of compound **6(A)** (1.10g, 4.74mmol), 3-acetamidophenylboronic acid (1.20g, 6.64mmol), copper acetate (258mg, 1.42mmol) and pyridine (229g, 2.84mmol) in DMF (30mL) was stirred at 65°C during 15h under room atmosphere. The mixture was diluted with 50mL of a saturated solution of NaHCO₃ and the aqueous layer was extracted twice with 60mL of DCM. The organic layers were combined dried over MgSO₄, filtered and concentrated under reduce pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 10g SiO₂) (Eluant: CH₂Cl₂/MeOH 98/2) to afford N-(3-(4-allyl-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **6(B)** (800mg, 46%) as a colorless oil.

LC₂: Rt= 2.01

MS m/z (ES) [M-H]⁻= 363.

6(C):N-(3-(4-allyl-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0083] According to Scheme 3 Step 3: To a solution of compound **6(B)** (800mg, 2.19mmol) in MeOH (10mL) was added. a solution of sodium methoxyde 2M in methanol (1.1mL, 2.19mol). After 1h of stirring at room temperature, the mixture was quenched with water and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water, brine, dried over MgSO₄, filtered and concentrated under reduced pressure. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 10g SiO₂) using CH₂Cl₂/MeOH 98/2 as eluant to afford N-(3-(4-allyl-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **6(C)** (600mg, 86%) as a white solid.

mp= 72-73°C

LC₂: Rt =1.71

MS m/z (ES) [M+H]⁺= 317 and (ES) [M-H]⁻= 315.

RMN 1H (DMSO-d₆), δ (ppm): 10.12 (1H, s), 7.88-7.87 (1H, m), 7.56-7.36 (1H, m), 7.40-7.35 (1H, t), 7.25-7.22 (1H, m), 5.89-5.80 (1H, m), 5.26-5.14 (2H, m), 4.45-4.43 (2H, m), 3.82 (3H, s), 2.09 (3H, s), 2.05 (3H, s).

6(D):N-(3-(4-allyl-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0084] According to Scheme 3 Step 4: To a solution of compound **6(C)** (600mg, 1.90mmol) in dioxane (5mL) was added dichlorotris(triphenylphosphine)ruthenium (91mg, 0.10mmol) and formic acid (87mg, 1.90mmol). The reaction mixture was heated at 200°C under microwave irradiation for 30min and filtered over celite. The filtrate was concentrated under reduced pressure. The crude product was then dissolved with AcOEt, washed twice with water and once with brine. The organic layer was dried, concentrated until dryness and the crude was then purified by flash chromatography with silica gel (Merck Flashmart prepacked column 10g SiO₂) using CH₂Cl₂/MeOH 98/2 as eluant to afford N-(3-(4-allyl-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **6(D)** (250mg, 50%) as a purple solid.

LC₂: Rt = 1.19

MS m/z (ES) [M+H]⁺= 277 and (ES) [M-H]⁻= 275.

6(E): N-(3-(4-(4-isopropylbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0085] According to Scheme 3 Step 5: A suspension of compound **6(D)** (60mg, 0.22mmol), 4-isopropylbenzyl bromide (41μL, 0.24mmol) and potassium carbonate (90mg, 0.65mmol) was stirred at 60°C for 15h in acetonitrile (2mL). The mixture was diluted with AcOEt and water. The organic layer was washed twice with water and once with brine. The resulting organic layer was dried over MgSO₄ and concentrated under reduced pressure. The crude product was purified by flash chromatography over silica gel (Merck Flashmart prepacked column 10g SiO₂) using CH₂Cl₂/MeOH 98/2 as eluent to afford N-(3-(4-(4-isopropylbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **6(E)** (10mg, 11%) as a yellow solid.

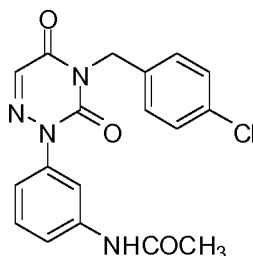
LC₂: Rt = 2.62

MS m/z (ES) $[M+H]^+ = 409.0$ and (ES) $[M-H]^- = 407.0$.

$^1\text{H-NMR}$ (DMSO-d_6), δ (ppm): 10.12 (1H, bs), 7.87 (1H, s), 7.57-7.54 (1H, m), 7.40-7.34 (1H, t), 7.30-7.18 (5H, m), 4.98 (2H, s), 3.81 (3H, s), 2.9-2.81 (1H, q), 2.05 (3H, s), 1.18-1.16 (6H, m).

Example 7: N-(3-(4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 2.01).

[0086]



7(A): 4-(4-Chlorobenzyl)-1,2,4-triazine-3,5(2H,4H)-dione.

[0087] According to Scheme 4 Step 1: To a solution of 6-Azaauracyl (5 g, 44.2mmol) and 4-chlorobenzyl bromide (9.09 g, 44.2mmol) in DMSO (190mL) under nitrogen atmosphere was added portionwise NaH (55%) (1.93g, 44.2mmol) at room temperature. The reaction mixture was stirred during 15min. 300mL of water was added and the aqueous layer was extracted thrice with ethyl acetate (300mL). The combined organic layers were washed twice with water (200mL), dried over MgSO_4 , filtered and concentrated under reduce pressure to afford a yellow solid. The crude solid was purified by flash chromatography with silica gel using $\text{CH}_2\text{Cl}_2/\text{MeOH}$ 98/2 as eluant to afford 4-(4-chlorobenzyl)-1,2,4-triazine-3,5(2H,4H)-dione **7(A)** (6.4g, 61%) as a white solid.

LC_1 : $\text{Rt} = 3.48$

MS m/z (ES) $[M-H]^- = 236$

7(B): N-(3-(4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0088] According to Scheme 4 Step 2: To a stirred solution of 4-(4-chlorobenzyl)-1,2,4-triazine-3,5(2H,4H)-dione (0.250g, 1.1mmol) in DMF (5mL) were added 3-acetamidophenylboronic acid (0.28g, 1.6mmol), copper(II) acetate (0.19g, 1mmol) and pyridine (0.17g, 2.1mmol) and the mixture was stirred overnight at 60°C . The reaction was quenched with a saturated sodium bicarbonate solution and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water then with brine, dried over MgSO_4 , filtered and concentrated. The crude product was purified by C_{18} reverse phase column, 15g column: starting from 20% CH_3CN and gradually increasing the percentage of CH_3CN (10% each 300 ml) the product came out at 60% of CH_3CN . 0.160g (39%) of N-(3-(4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **7(B)** were obtained.

LC_1 : $\text{Rt} = 3.92$

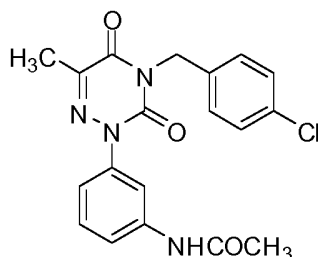
MS m/z (ES) $[M+H]^+ = 371.0$

$^1\text{H-NMR}$ (CDCl_3), δ (ppm): 7.81 (1H, bs), 7.57-7.54 (1H, m), 7.49-7.18 (9H, m), 5.1 (2H, s).

Example 8: N-(3-(4-(4-chlorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 3.01).

[0089]

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8(A): 4-(4-chlorobenzyl)-6-methyl-1,2,4-triazine-3,5(2H,4H)-dione.

[0090] According to Scheme 4 Step 1: To a solution of Azathymine (300mg, 2.36mmol) and 4-chlorobenzyl bromide (485mg, 2.36mmol) in DMSO (10mL) was added portionwise NaH (55%) (103mg, 2.36mmol) at room temperature under nitrogen atmosphere. The mixture was stirred for 15min at room temperature. Ethyl Acetate was then added (150mL) and the organic layer was extracted thrice with water (100mL). The organic layer was dried over MgSO₄, filtered and concentrated under reduced pressure to afford a yellow solid. The crude solid was purified by flash chromatography with silica gel using CH₂Cl₂/MeOH 98/2 as eluant to afford 4-(4-chlorobenzyl)-6-methyl-1,2,4-triazine-3,5(2H,4H)-dione **8(A)** (500mg, 84%) as a white solid.

LC₁: Rt= 3.66

MS m/z (ES) [M-H]⁻= 249

8(B): N-(3-(4-(4-chlorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0091] According to Scheme 4 Step 2: A mixture of compound **8(A)** (100mg, 0.39mmol), 3-acetamidophenylboronic acid (107mg, 0.59mmol), copper acetate (79mg, 0.44mmol) and pyridine (65μL, 0.79mmol) in DMF (3mL) was stirred at 65°C during 15 hours. The reaction mixture was diluted with 100mL of a saturated solution of NaHCO₃ and the aqueous layer was extracted twice with DCM (100mL). The organic layers were combined and successively dried over MgSO₄, filtered and concentrated under reduce pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 20g, C18) using Water/CH₃CN 6/4 as eluant to afford N-(3-(4-(4-chlorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **8(B)** (97mg, 63%) as a white solid.

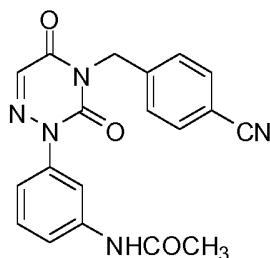
LC₁: Rt= 4.08

MS m/z (ES) [M+H]⁺= 384

¹H-NMR (DMSO-d₆), δ (ppm): 10.12 (1H, bs), 7.8 (1H, t), 7.55 (1H, d), 7.37 (5H, m), 7.15 (1H, dd), 5 (2H, s), 2.19 (3H, s), 2.05 (3H, s).

Example 9: N-(3-(4-(4-cyanobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 2.05).

[0092]



9(A): Ethyl (Z)-2-cyano-2-(2-(3-nitrophenyl)hydrazono)acetylcarbamate.

[0093] According to Scheme 5 Step 1: To a suspension of nitroaniline (5g, 36.2mmol) in concentrated HCl (9mL) and water (50mL) at 0°C was added dropwise over 30minutes a solution of sodium nitrite (5g, 72.4mmol in 200mL water) the crude orange suspension was filtered at 0°C and the obtained diazonium salt was added portionwise over 1hour under vigorous stirring to a suspension of ethyl 2-cyanoacetate (6.22g, 39.8mmol) and sodium acetate (21.0g, 257mmol)

in ethanol (200mL). The reaction mixture was allowed to rest at 0°C for 1h. The mixture was filtered, washed with water then cooled ethanol and dried to afford (Z)-2-cyano-2-(2-(3-nitrophenyl)hydrazono)acetylcarbamate **9(A)** (11.0g, 100%, 75% of purity) as an orange solid. The crude solid was used in the next step without further purification.

Rf= 0.32 CH₂Cl₂/MeOH (95/5)

LC₁: Rt= 3.65

MS m/z (ES) [M+H]⁺= 306

9(B): 2-(3-nitrophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carbonitrile.

[0094] According to Scheme 5 Step 2: The crude compound **9(A)** (11.0g, 75% of purity) was stirred with NaOH 1M (65mL) and 200mL of water. The corresponding heterogeneous mixture was stirred at room temperature. After 1h, two additional set of NaOH 1M (20mL+20mL) was added. After complete conversion, the mixture was diluted with EtOAc and the aqueous layers were successively decanted and acidified. The acidified aqueous layers were extracted with EtOAc and the organic layers were combined and successively washed with brine, dried over MgSO₄ filtered and concentrated to afford the **9(B)** (8g, 47%) as a crude yellow solid. **9B** was used in the next step without further purification.

LC₁: Rt= 3.46

MS m/z (ES) [M-H]⁻= 258

9(C): 2-(3-nitrophenyl)-1,2,4-triazine-3,5(2H,4H)-dione **5.5**.

[0095] According to Scheme 5 Step 3: 2-(3-nitrophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carbonitrile (3.8g) was successively splitted over 8 microwave flasks, diluted with an aqueous solution of HCl (6N, 10mL) and were irradiated under Microwave at 150°C for 1 hour. The clear brown solids were combined filtered and dried to afford the corresponding acid (6.5g, 76% yield).

LC₁: Rt= 2.08

A solution of 2-(3-nitrophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylic acid (3g) in diphenylether (30mL) was refluxed for 1hour and cooled to room temperature. The mixture was quenched with 1M NaOH solution. The basic aqueous layer was then acidified and extracted with ethyl acetate (x3). The organic layers were successively washed with brine, dried over MgSO₄, filtered and concentrated. 1g (40% yield) of 2-(3-nitrophenyl)-1,2,4-triazine-3,5(2H,4H)-dione **9(C)** was recovered and used in next step without any further purification.

LC₁: Rt= 2.95

9(D): N-(3-(3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **5.6**.

[0096] According to Scheme 5 Step 4: To a mixture of compound **9(C)** (2g, 8.54mmol) and 60mL of H₂O/EtOH 50/50 and acetic acid (0.488mL, 8.54mmol) was added iron (2.24 g, 40.1mmol). The reaction mixture was heated at 80°C. A solution of NaOH 1M was added until neutral pH. The mixture was filtered over celite. The aqueous layers were diluted with EtOAc and decanted. The combined organic layers were acidified till pH=2 and extracted with EtOAc. The combined organic layers were successively washed with brine, dried over MgSO₄, filtered and concentrated to afford 1.2g (69% yield) of the crude compound 2-(3-aminophenyl)-1,2,4-triazine-3,5(2H,4H)-dione which is used in next step without any further purification.

LC₁: Rt= 1.74

MS m/z (ES) [M+H]⁺= 205.2

To a stirred solution of 2-(3-aminophenyl)-1,2,4-triazine-3,5(2H,4H)-dione (1.2g, 5.88mmol) in THF (20mL) was added acetic anhydride (1.2 g, 11.8mmol) and the mixture was heated at 50 °C. After 2h the mixture was cooled at room temperature, quenched with water and extracted with ethyl acetate (x3). The organic layers were washed with brine, dried over MgSO₄, filtered and concentrated to afford the 900mg, 62% yield, of N-(3-(3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **9(D)** (900mg, 62%) which is used in next step without any further purification.

LC₁: Rt= 2.23

MS m/z (ES) [M+H]⁺= 247.1

9(E): N-(3-(4-(4-cyanobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0097] According to Scheme 5 Step 5: A suspension of N-(3-(3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (50mg, 0.203mmol), 4-cyanobenzylbromide (40mg, 0.203mmol) and potassium carbonate (31mg, 0.223mmol) in acetonitrile (2mL) was stirred at 60°C for 15h. The reaction mixture was then partitioned between AcOEt and water. The organic layer was washed twice with water and once with brine. The organic layer was dried over MgSO₄ and evaporated under reduced pressure. The crude product was purified by flash chromatography over silica gel (Merck

Flashmart preppacked column 10g SiO₂) using DCM/MeOH 98/2 as eluent to afford N-(3-(4-(4-cyanobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **9(E)** (26mg, 35%) as a white solid.

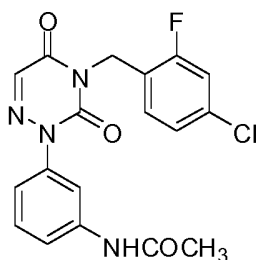
LC₁: Rt = 3.47

MS m/z (ES) [M+H]⁺ = 362 and (ES) [M-H]⁻ = 360.

¹H-NMR (DMSO-d₆), δ (ppm): 10.14 (1H, bs), 7.85-7.80 (4H, m), 7.60-7.55 (3H, m), 7.41-7.36 (1H, t), 7.18-7.16 (1H, m), 5.07 (2H, s), 2.05 (3H, s).

Example 10: N-(3-(4-(4-chloro-2-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 2.20).

[0098]



10(A): N-(3-(5-methoxy-3-oxo-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0099] According to Scheme 6 Step 1: To a stirred solution of 5-methoxy-1,2,4-triazin-3(2H)-one (1g, 7.9mmol) in DMF (5mL) were successively added the 3-acetamidophenylboronic acid (1.4g, 7.9mmol), copper(II) acetate (1.4g, 7.9mmol) and pyridine (1.2g, 7.9mmol). The mixture was stirred overnight at 60°C. The reaction mixture was quenched with a saturated sodium bicarbonate solution and extracted with ethyl acetate (x3). The combined organic layers were washed successively with water, brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 70g column, CH₂Cl₂/MeOH 98/2 to afford (0.493g, 24%) of N-(3-(5-methoxy-3-oxo-1,2,4-triazin-2(3H)-yl)phenyl)acetamide

LC₁: Rt = 2.46

MS m/z (ES) [M+H]⁺ = 261.2

10(B): N-(3-(4-(4-chloro-2-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0100] According to Scheme 6 Step 2: To a stirred solution of N-(3-(5-methoxy-3-oxo-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (40mg, 0.15mmol) in acetonitrile (3mL) were added 1-(bromomethyl)-4-chloro-2-fluorobenzene (52mg, 0.23mmol) and sodium iodide (46mg, 0.31mmol). The mixture was heated irradiated at 160°C under microwave irradiation. The reaction mixture was quenched with water and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water then with brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 5g column, CH₂Cl₂/MeOH 98/2 to afford the N-(3-(4-(4-chloro-2-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **10(B)** as a yellow solid (13mg, 22%).

LC₁: Rt = 3.98

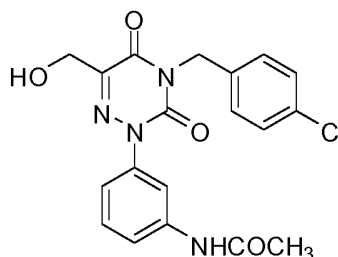
MS m/z (ES) [M+H]⁺ = 389.2

mp = 184°C

¹H-NMR (DMSO-d₆), δ (ppm): 10.14 (1H, s), 7.85-7.81 (2H, m), 7.57-7.13 (6H, m), 5 (2H, s), 2.06-2.03 (3H, m).

Example 11: N-(3-(4-(4-chlorobenzyl)-6-(hydroxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 5.14).

[0101]



11(A): (Z)-ethyl 2-(2-(3-acetamidophenyl)hydrazono)-2-cyanoacetylcarbamate.

[0102] According to Scheme 7 Step 1: To a suspension of 3-aminophenylacetamide (1.50g, 9.9mmol), in concentrated HCl (2.5mL) and water (11mL) at 0°C was added dropwise over 30min a solution of sodium nitrite (1.38g, 19.9mmol) in water (20mL). The crude orange suspension was stirred at 0°C and the obtained diazonium salt was added portionwise over 1hour under vigorous stirring to a suspension made of compound N-cyanoacetylurethane (1.71g, 10.9mmol) and sodium acetate (5.82g, 70.9mmol) in ethanol (37mL). The reaction mixture was stirred at 0°C for 2h. The brown resulting mixture was successively filtered, washed with water, cooled ethanol, Et₂O and dried to afford the hydrazone Ethyl (Z)-2-cyano-2-(2-(3-acetamidophenyl)hydrazono)acetylcarbamate **11(A)** (3.18g, 99%).

[0103] **11(B):** N-(3-(6-cyano-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide. According to Scheme 7 Step 2: A mixture of compound **11(A)** (3.18g, 9.9mmol) in water (10mL) and sodium hydroxide (0.40g, 9.9mmol) was stirred during 30min at room temperature. The mixture was filtered. The precipitate was dried under vacuum to afford N-(3-(6-cyano-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **11(B)** (1.9g, 69%) as an orange solid. LC₁: Rt= 2.75

MS m/z (ES) [M-H]⁻ = 270

11(C): N-(3-(4-(4-chlorobenzyl)-6-cyano-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0104] According to Scheme 7 Step 3: A suspension of **11(B)** (500mg, 1.8mmol), 4-chlorobenzyl bromide (455mg, 2.2mmol) and potassium carbonate (510mg, 3.7mmol) in CH₃CN (10mL) was stirred 12h under reflux. The mixture was cooled at room temperature and 100mL of water were added. The aqueous layer was extracted thrice with DCM (100mL). The organic layers were mixed, dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (AIT Flashmart prepacked column 50g SiO₂) using CH₂Cl₂/MeOH (98/2) as eluant to afford N-(3-(4-(4-chlorobenzyl)-6-cyano-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **11(C)** (460mg, 63%) as a beige solid.

11(D): 2-(3-acetamidophenyl)-4-(4-chlorobenzyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylic acid.

[0105] According to Scheme 7 Step 4: A mixture of compound **11(C)** (2.00g, 5.1mmol) and hydrochloric acid 33% (30mL) was stirred at reflux during 3h. The solvent was removed concentrated under reduce pressure and then the crude residue was treated with a saturated solution of Na₂CO₃ (50mL). The aqueous layer was acidified (pH=4) with acetic acid (pH=4) and the reaction mixture was filtered to afford 4-(4-chlorobenzyl)-2-(3-aminophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylic acid (1.65g, 88%) as a beige solid.

To a solution of the crude chlorobenzyl)-2-(3-aminophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylic acid (1.65g) in THF (50mL) was added the acetyl chloride (751μL, 10.4mmol) dropwise. The mixture was stirred for 1hour. The reaction mixture was quenched with water (2mL) and the resulting mixture was concentrated under reduced pressure to afford 4-(4-chlorobenzyl)-2-(3-acetamidophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylic acid **11(D)** (1.70g, 98%) as a beige solid.

11(E): N-(3-(4-(4-chlorobenzyl)-6-(hydroxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0106] According to Scheme 7 Step 5: To a solution of **11(D)** (380mg, 0.92mmol), and diisopropylethylamine (189μL, 1.1mmol) in THF (10mL) at 0°C under nitrogen atmosphere was added ethyl chloroformate (101μL, 1.06mmol) dropwise. The reaction mixture was stirred for 1h then treated dropwise with a solution of sodium borohydride (90mg, 2.38mmol) in water (2mL). The mixture was stirred overnight at room temperature and quenched slowly with a saturated solution of NaHCO₃ (100mL). The aqueous layer was extracted thrice with CH₂Cl₂ (100mL). The organic layers were combined, dried over MgSO₄, filtered and concentrated under reduced pressure to afford a yellow solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 20g SiO₂) using CH₂Cl₂/MeOH 97/3 as

eluant to afford N-(3-(4-(4-chlorobenzyl)-6-(hydroxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **11(E)** (32mg, 9%) as a white solid.

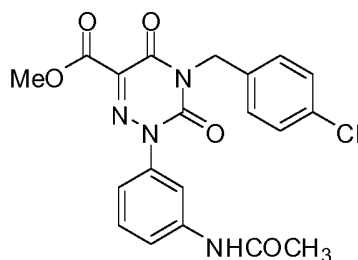
LC₂: Rt= 2.07

MS m/z (ES) [M+H]⁺= 400.8

RMN 1H (300MHz, DMSO-d₆), δ (ppm): 10.14 (1H, bs), 7.82 (1H, t), 7.56 (1H, dd), 7.4 (5H, m), 7.17 (1H, dd), 5 (2H, s), 5.31 (1H, t, 6Hz), 4.4 (2H, d, 6Hz), 2.05 (3H, s).

Example 12: Methyl 4-(4-chlorobenzyl)-2-(3-acetamidophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylate (Final Compound 5.17).

[0107]



[0108] According to Scheme 7 Step 5: To a solution of 2-(3-acetamidophenyl)-4-(4-chlorobenzyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylic acid **11(D)** (80mg, 0.19mmol) in MeOH (3mL) was added dropwise at 0°C a solution of trimethylsilyldiazomethane 2M (1.3mL, 2.70mmol). The mixture was stirred at room temperature for 18h. The mixture was diluted with 100mL of a saturated solution of NaHCO₃ and the aqueous layer was extracted twice with 100mL of DCM. The organic layers were combined, dried over MgSO₄, filtered and concentrated under reduced pressure to afford a yellow solid. The crude solid was purified by flash chromatography with silica gel (AIT Flashmart prepacked column 7g SiO₂) using CH₂Cl₂/MeOH 98/2 as eluant to afford a yellow solid. The solid was further triturated with diisopropylether and filtered to afford methyl 4-(4-chlorobenzyl)-2-(3-acetamidophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylate (35mg, 42%) as a beige solid.

LC₁: Rt= 3.95

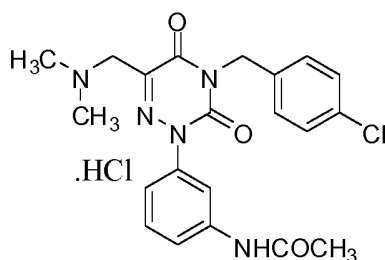
MS m/z (ES) [M+H]⁺= 428.0

mp= 193°C

¹H-NMR (DMSO-d₆), δ (ppm): 7.84 (1H, bs), 7.45 (5H, m), 7.3 (3H, m), 5.15 (2H, s), 3.97 (3H, s), 2.2 (3H, s).

Example 13: N-(3-(4-(4-chlorobenzyl)-6-((dimethylamino)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide hydrochloride (Final Compound 5.15).

[0109]



13(A): N-(3-(6-(bromomethyl)-4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0110] According to Scheme 7 Step 6: To a solution of N-(3-(4-(4-chlorobenzyl)-6-(hydroxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (500mg, 1.25mmol) in acetonitrile (20mL) under nitrogen atmosphere was added phosphorus tribromide (176μL, 1.87mmol) dropwise and the mixture was refluxed for 90min. The reaction mixture was diluted with 200mL a saturated solution of NaHCO₃ (200mL) and the aqueous layer was extracted twice with DCM (200mL). The organic layers were combined to be dried over MgSO₄, filtered and concentrated under reduced

pressure to afford N-(3-(4-(4-chlorobenzyl)-6-(bromomethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **13(A)** (540mg, 93%) as a yellow solid.

13(B): N-(3-(4-(4-chlorobenzyl)-6-((dimethylamino)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide hydrochloride.

[0111] According to Scheme 7 Step 7: A solution of **13(A)** (60mg, 0.13mmol), (1mL), and dimethylamine 2M (in methanol, 23 μ L, 0.65mmol) in DMF (1ml) was irradiated at 80°C under microwave irradiation for 5min. The mixture was diluted with 50mL of a saturated solution of NaHCO₃, and the aqueous layer was extracted twice with 50mL of DCM. The organic layers were combined and successively dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 5g SiO₂) using CH₂Cl₂/MeOH 95/5 as eluant to afford a white solid. The solid was dissolved in DCM (3mL) and hydrogen chloride 1N (2mL) was added. The solution was concentrated under reduced pressure to afford the HCl salt N-(3-(4-(4-chlorobenzyl)-6-((dimethylamino)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide hydrochloride **13(B)** (17mg, 28%) as a beige solid.

LC₂ : Rt= 1.74

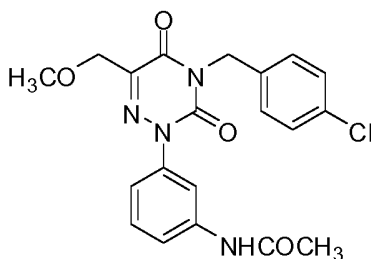
MS m/z (ES) [M+H]⁺= 427.9

mp= 143°C

¹H-NMR (DMSO-d₆), δ (ppm): 10.27 (1H, bs), 10.09 (1H, bs), 7.98 (1H, t), 7.47 (6H, m), 7.21 (1H, dq), 5.02 (2H, s), 4.3 (2H, s), 2.87 (6H, s), 2.06 (3H, s).

Example 14: N-(3-(4-(4-chlorobenzyl)-6-(methoxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (Final Compound 5.16).

[0112]



14(A): N-(3-(4-(4-chlorobenzyl)-6-(iodomethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0113] According to Scheme 7 Step 6: A mixture of N-(3-(4-(4-chlorobenzyl)-6-(bromomethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (200mg, 0.43mmol) and sodium iodide (97mg, 0.65mmol) in acetone (10mL) was stirred overnight at room temperature. The mixture was diluted with water (100mL) and the aqueous layer was extracted twice with 100mL of DCM. The organic layer were combined and successively dried over MgSO₄, filtered and concentrated under reduced pressure to afford N-(3-(4-(4-chlorobenzyl)-6-(iodomethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **14(A)** (220mg) as a white solid. The tittle product was used in the next reaction without any further purification.

LC₂: Rt= 2.69

MS m/z (ES) [M+H]⁺= 510.7

14(B): N-(3-(4-(4-chlorobenzyl)-6-(methoxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

[0114] According to Scheme 7 Step 7: To a cooled solution of **14(A)** (220mg, 0.43mmol) in DMF (2mL) under nitrogen atmosphere was added dropwise at 0°C sodium methoxide 2M (in methanol, 215 μ L, 0.43mmol). The mixture was stirred at 0°C for 1h. The reaction mixture was diluted with water (100mL) and the aqueous layer was extracted twice with DCM (100mL). The organic layers were combined and successively dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 25g C₁₈) using water/CH₃CN 6/4 as eluant to afford N-(3-(4-(4-chlorobenzyl)-6-(methoxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide **14(B)** (16mg, 9%) as a yellow solid.

LC₂ : Rt= 2.36

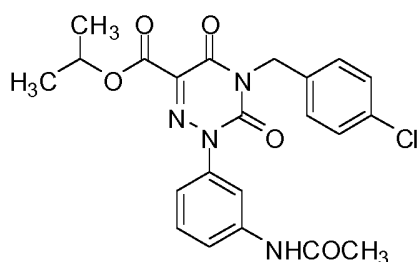
MS m/z (ES) [M+H]⁺= 414.8

mp= 97°C

¹H-NMR (DMSO-d₆), δ (ppm): 10.15 (1H, bs), 7.82 (1H, t), 7.57 (1H, dd), 7.4 (5H, m), 7.17 (1H, dd), 5 (2H, s), 4.34 (2H, s), 2.05 (3H, s).

Example 15: Isopropyl 2-(3-acetamidophenyl)-4-(4-chlorobenzyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylate (Final Compound 5.18).

[0115]



[0116] According to Scheme 7 Step 5: A mixture of 4-(4-chlorobenzyl)-2-(3-acetamidophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylic acid **7.6** (100mg, 0.24mmol), isopropylalcohol (3mL) and sulfuric acid (2drops) was heated at 80°C for 20 hours. The reaction mixture was diluted with a saturated solution of NaHCO₃ (50mL) and the aqueous layer was extracted twice with DCM (50mL). The organic layers were combined and selectively dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 7g SiO₂) using CH₂Cl₂/MeOH 98/2 as eluant to afford isopropyl 4-(4-chlorobenzyl)-2-(3-acetamidophenyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylate (33mg, 30%) as a beige solid.

LC₂ : Rt= 2.65

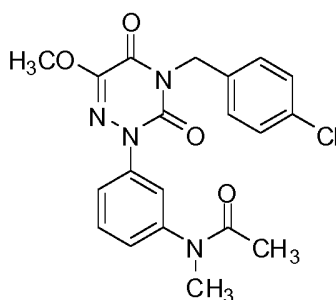
MS m/z (ES) [M+H]⁺= 457.0

mp= 183°C

¹H-NMR (DMSO-d₆), δ (ppm): 10.19 (1H, bs), 7.84 (1H, s), 7.59 (1H, d), 7.41 (5H, m), 7.16 (1H, dd), 5.13 (1H, m), 4.97 (2H, s), 2.05 (3H, s), 1.27 (6H, d).

Example 16: N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide (Final Compound 4.133).

[0117]



[0118] According to Scheme 9 Step 1: To a solution of N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide (70mg, 0.17mmol) in THF (3mL) was added portionwise at 0°C sodium hydride (55%) 11mg, 0.26mmol). The suspension was stirred 10 minutes at room temperature. Then, methyl iodide (87μL, 1.40mmol) was added dropwise and the reaction mixture was stirred at room temperature under nitrogen atmosphere overnight. The mixture was diluted with 100mL of a saturated solution of ammonium chloride and the aqueous layer was extracted twice with 100mL of DCM. The organic layers were combined to be successively dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 10g SiO₂) using CH₂Cl₂/MeOH 99/1 as eluant to afford

N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide (54mg, 75%) as a white solid.

LC₁ : Rt= 4.09

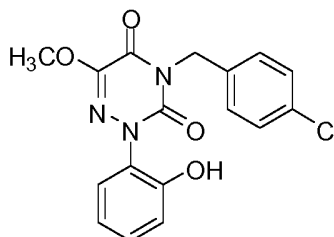
MS m/z (ES) [M+H]⁺= 414.0

mp= 157°C

¹H-NMR (DMSO-d₆), δ (ppm): 7.5 (8H, m), 5.01 (2H, s), 3.85 (3H, s), 3.18 (3H, s), 2.5 (3H, s).

Example 17: 4-(4-chlorobenzyl)-2-(2-hydroxyphenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione (Final Compound 4.143).

[0119]



17(A): 4-(4-chlorobenzyl)-6-bromo-2-(2-hydroxyphenyl)-1,2,4-triazine-3,5(2H,4H)-dione.

[0120] According to Scheme 9 Step 1: To a solution of 4-(4-chlorobenzyl)-6-bromo-2-(2-methoxyphenyl)-1,2,4-triazine-3,5(2H,4H)-dione (150 mg, 0.35 mmol) in DCM (2 ml) cooled at -78°C was added dropwise BBr₃ (0.710 mL, 0.710 mmol). The reaction mixture was stirred 1h at -78°C and was quenched with MeOH. The mixture was allowed to reach room temperature. Water was added and the aqueous layer was extracted with DCM (x3). The combined organic layers were successively washed with water, brine, dried over MgSO₄, filtered and concentrated. 131 mg (yield=90%) of 4-(4-chlorobenzyl)-6-bromo-2-(2-hydroxyphenyl)-1,2,4-triazine-3,5(2H,4H)-dione were recovered as a beige solid which was used in the next step without any purification.

17(B): 4-(4-chlorobenzyl)-2-(2-hydroxyphenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione.

[0121] According to Scheme 9 Step 2: To a solution of 4-(4-chlorobenzyl)-6-bromo-2-(2-hydroxyphenyl)-1,2,4-triazine-3,5(2H,4H)-dione (0.130g, 0.32 mmol) in DMF (3 ml) was added, dropwise, at 0°C a solution 2M of sodium methoxide (0.32 mmol, 0.160 ml). The mixture was stirred 1h at 0°C. 1 more equivalent of a solution 2M of sodium methoxide was necessary to complete the conversion. Then the reaction mixture was quenched with water. The aqueous layers was neutralised till pH=7 and extracted with ethyl acetate (x3). The organic layer were combined and successively washed with water then, brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 10 g prepacked column, DCM/MeOH 98/2 to afford (30mg, Yield= 26%) 30 mg (Yield= 26%) of 4-(4-chlorobenzyl)-2-(2-hydroxyphenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione **4.143** were recovered as a white solid.

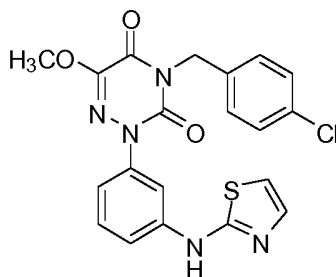
LC₂ : Rt= 2.45

mp= 164-165°C

¹H-NMR (DMSO-d₆), δ (ppm): 7.4 (4H, s), 7.30-7.23 (2H, m), 6.95-6.82 (2H, m), 5 (2H, s), 3.74 (3H, s).

Example 18: 4-(4-chlorobenzyl)-6-methoxy-2-(3-(thiazol-2-ylamino)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione (Final Compound 4.144).

[0122]



18(A): 2-(3-aminophenyl)-4-(4-chlorobenzyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione.

[0123] According to Scheme 9 Step 1: A mixture of N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)yl)phenyl)acetamide (500mg, 1.24mmol) in a solution 6N of HCl (10mL) and THF (10mL) was stirred at reflux for 3hours. The THF was evaporated and the aqueous layer was basified with saturated solution of NaHCO₃; until neutral pH. The aqueous layer was then extracted with ethyl acetate (x3). The organic layers were combined to be successively washed with water then with brine, dried over MgSO₄, filtered and concentrated. The crude product was azeotroped with acetonitrile to afford 2-(3-aminophenyl)-4-(4-chlorobenzyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione as a beige solid (423mg, 94.5%).

18(B): 4-(4-chlorobenzyl)-6-methoxy-2-(3-(thiazol-2-ylamino)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione.

[0124] According to Scheme 9 Step 2: To a Solution of 2-(3-Aminophenyl)-4-(4-chlorobenzyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione (125mg, 0.34mmol) in ethanol (10ml) was added bromothiazole (124μL, 1.4mmol) and 5 mL of concentrated hydrochloric acid. The reaction mixture was heated to 100°C for 48 hours. The reaction mixture was quenched and neutralised with sat NaHCO₃ and extracted in ethyl acetate (x3). The organic layers were combined and successively washed with water, brine, dried over MgSO₄, filtered and concentrated. The crude oily solid was purified by flash chromatography, 10 g prepacked column, cyclohexane/ ethyl acetate 70/30 to afford 4-(4-chlorobenzyl)-6-methoxy-2-(3-(thiazol-2-ylamino)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione **4.144** (10mg (Yield= 6.5%) as a white solid.

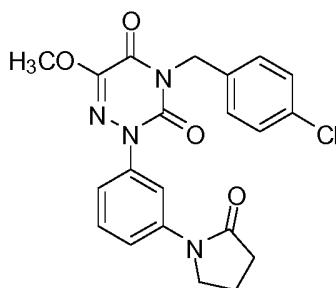
LC₂ : Rt= 2.82

mp= 176-178°C

¹H-NMR (CDCl₃), δ (ppm): 10.4 (1H, s), 8.0 (1H,s), 7.6-6.9 (9H, m), 5.0 (2H,s), 3.8 (3H,s).

Example 19: 4-(4-chlorobenzyl)-6-methoxy-2-(3-(2-oxopyrrolidin-1-yl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione (Final Compound 4.145).

[0125]



[0126] According to Scheme 9 Step 2: To a solution of 2-(3-aminophenyl)-4-(4-chlorobenzyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione (100mg, 0.27mmol) (1mL) and NEt₃ (78μL, 0.55mmol) in DMF was added bromobutyryl chloride (36μL, 0.30mmol). The reaction mixture was heated at 80°C for 4 hours then quenched with water and extracted with ethyl acetate. The organic layers were combined and successively washed with water, brine, dried over MgSO₄, filtered and concentrated. The crude product was purified on a prepacked silica column (10g) 8:2 cyclohexane/Ethyl acetate 80/20 to afford 4-(4-chlorobenzyl)-6-methoxy-2-(3-(2-oxopyrrolidin-1-yl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione **4.145** as a white solid (6mg, 5%).

LC₂ : Rt= 2.64

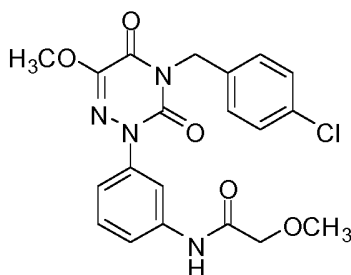
MS m/z (ES) [M+H]⁺ = 427.0

mp = 91-93°C

¹H-NMR (CDCl₃), δ (ppm): 7.9 (1H, s), 7.6-7.3 (7H, m), 5.2 (2H, s), 3.9 (3H, s), 1.5-0.9 (6H, m).

Example 20: N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-2-methoxyacetamide (Final Compound 4.141).

[0127]



[0128] According to Scheme 9 Step 2: A mixture of 2-(3-aminophenyl)-4-(4-chlorobenzyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione (100mg, 0.28mmol), DMF (4mL), diisopropylethylamine (95μl, 0.56mmol) and methoxyacetyl chloride (30μl, 0.33mmol) was stirred under nitrogen at room temperature during 1hour. The mixture was diluted with 50mL of a saturated solution of sodium carbonate and the aqueous layer was extracted twice with 50mL of DCM. The organic layers were combined to be dried over MgSO₄, filtered and concentrated under reduced pressure to afford a brown solid. The crude solid was purified by flash chromatography with silica gel (Merck Flashmart prepacked column 15g SiO₂) using AcOEt/cyclohexane 4/6 as eluant to afford a beige solid. The solid was washed with diethylether to afford N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-2-methoxyacetamide **4.141** (84mg, 70%) as a white solid.

LC₂ : Rt = 2.48

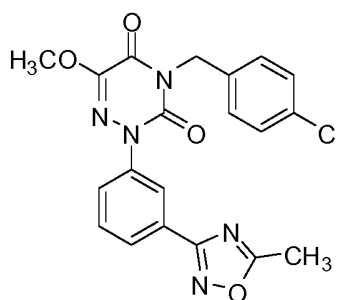
MS m/z (ES) [M+H]⁺ = 431.0

mp = 143°C

¹H-NMR (CDCl₃), δ (ppm): 10.1 (1H, bs), 8.01 (1H, t), 7.65 (1H, d), 7.4 (5H, m), 7.33 (1H, dd), 5.01 (2H, s), 4.02 (2H, s), 3.81 (3H, s), 3.38 (3H, s)

Example 21: 4-(4-chlorobenzyl)-6-methoxy-2-(3-(5-methyl-1,2,4-oxadiazol-3-yl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione (Final Compound 4.146).

[0129]



21(A): 3-(4-(4-chlorobenzyl)-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile.

[0130] According to Scheme 2 Step 3: To a solution of 6-bromo-4-(4-chlorobenzyl)-1,2,4-triazine-3,5(2H,4H)-dione (0.600 g, 1.9 mmol) in DCM (8 ml) were added 3-cyanophenylboronic acid (0.42g, 2.8 mmol), copper(II) acetate (0.069g, 0.38 mmol), pyridine (0.3g, 3.8 mmol) and pyridine N-oxide (0.2g, 2.1 mmol) and the mixture was stirred overnight at room temperature. The reaction mixture was quenched with a saturated sodium bicarbonate solution and extracted with ethyl acetate (x3). The combined organic layers were washed with water then with brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 30 g prepacked column, cyclohexane/ethyl acetate

85/15 to afford 3-(4-(4-chlorobenzyl)-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile (300mg, 38%) which was used in the next step without any purification

21(B): 3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile.

[0131] According to Scheme 2 Step 4: To a stirred solution of 3-(4-(4-chlorobenzyl)-6-bromo-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile (0.300g, 0.72 mmol) in DMF (5 ml) was added, dropwise, at 0°C, 2M solution of sodium methoxide (0.72 mmol, 0.359 ml). The mixture was stirred 1 hour at 0°C then quenched with water and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water then with brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 10 g prepacked column, DCM 100%. 150 mg (Yield=57%) of 3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile were recovered and used for the next step.

21(C): (Z)-3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-N'-hydroxybenzamidine.

[0132] According to Scheme 9 Step 1: To a stirred solution of 3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile (150mg, 0.41 mmol) in ethanol (5ml) was added hydroxylamine (0.100ml of a 50% solution in water, 1.2 mmol) and the mixture was heated at 80°C overnight. The reaction mixture was quenched with a saturated ammonium chloride solution and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water, with brine, dried over MgSO₄, filtered and concentrated to afford (Z)-3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-N'-hydroxybenzamidine (150 mg, 99%) which was used in the next step without any further purification.

21(D): 4-(4-chlorobenzyl)-6-methoxy-2-(3-(5-methyl-1,2,4-oxadiazol-3-yl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione.

[0133] According to Scheme 9 Step 2: A mixture of (Z)-3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-N'-hydroxybenzamidine (150mg, 0.4mmol), triethyl orthoacetate (2mL) and of sulfuric acid (1 drop) was stirred at 100°C 4 hours. The reaction mixture was quenched with water and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water, brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 10 g prepacked column, cyclohexane/ethyl acetate 80/20 to afford 4-(4-chlorobenzyl)-6-methoxy-2-(3-(5-methyl-1,2,4-oxadiazol-3-yl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione **4.146** (40mg, Yield= 25%) as a white solid.

LC₂ : Rt= 2.88

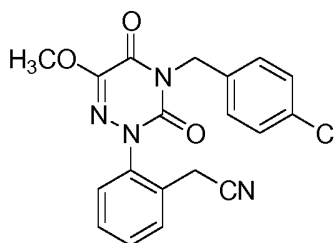
MS m/z (ES) [M+H]⁺= 426.0

mp= 163-165°C

¹H-NMR (CDCl₃), δ (ppm): 8.27 (1H, bs), 7.98 (1H, d), 7.82-7.78 (1H, m), 7.68 (1H, t), 7.44-7.36 (4H, m), 5.03 (2H, s), 3.85 (3H, s), 2.68 (3H, s)

Example 22: 2-(2-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetonitrile (Final Compound 4.167).

[0134]



22(A): 6-bromo-4-(4-chlorobenzyl)-2-(2-(hydroxymethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione.

[0135] According to Scheme 2 Step 3: To a solution of 6-bromo-4-(4-chlorobenzyl)-1,2,4-triazine-3,5(2H,4H)-dione (0.400g, 1.3mmol) in DMF (8mL) were added 2-(hydroxymethyl)phenylboronic acid (0.29g, 1.9mmol), copper(II) acetate (0.23g, 1.3 mmol) and pyridine (0.20g, 2.5mmol) and the mixture was left stirring overnight at 60°C. The reaction mixture was quenched with a saturated sodium bicarbonate solution. The precipitate formed was filtered, washed first with water

and then with diethyl ether. 330 mg (Yield=62%) of 6-bromo-4-(4-chlorobenzyl)-2-(2-(hydroxymethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione were recovered. The product was pure enough to be used without any purification.

22(B): 4-(4-chlorobenzyl)-2-(2-(hydroxymethyl)phenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione.

[0136] According to Scheme 2 Step 4: To a stirred solution of 6-bromo-4-(4-chlorobenzyl)-2-(2-(hydroxymethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione (0.330g, 0.79mmol) in DMF (10 ml) was added, dropwise, at 0°C, 2M solution of sodium methoxide (0.79mmol, 0.40ml). The mixture was stirred 1 hour at 0°C then quenched with water and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water then with brine, dried over MgSO₄, filtered and concentrated. 270mg (Yield=92.5%) of 4-(4-chlorobenzyl)-2-(2-(hydroxymethyl)phenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione were recovered. The product was pure enough to be used in the next step without any purification.

22(C): 2-(2-(bromomethyl)phenyl)-4-(4-chlorobenzyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione.

[0137] According to Scheme 9 Step 1: To a stirred solution of 4-(4-chlorobenzyl)-2-(2-(hydroxymethyl)phenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione (70mg, 0.19mmol) in CH₃CN (10mL) was added phosphorus tribromide (76 mg, 0.28mmol) and the mixture was heated at 60°C during two hours. The reaction mixture was diluted with 100mL of water and the aqueous layer was extracted with DCM (x2). The combined organic layers were successively washed with water, with brine, dried over MgSO₄, filtered and concentrated to afford 70mg (Yield=86%) of 2-(2-(bromomethyl)phenyl)-4-(4-chlorobenzyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione. The product was pure enough to be used in the next step without any purification.

22(D): 2-(2-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetonitrile.

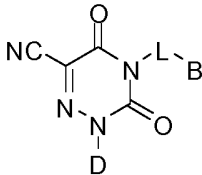
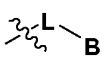
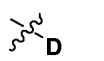
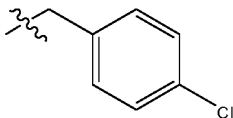
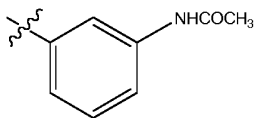
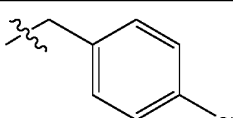
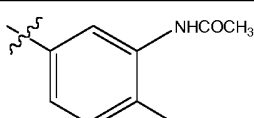
[0138] According to Scheme 9 Step 2: A mixture of 4-(4-chlorobenzyl)-2-(2-(bromomethyl)phenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione (70mg, 0.16mmol) in DMF (2mL) and potassium cyanide (31mg, 0.13 mmol) was stirred at 20°C 12 hours. The reaction mixture was quenched with water and extracted with ethyl acetate (x3). The combined organic layers were successively washed with water, brine, dried over MgSO₄, filtered and concentrated. The crude product was purified by flash chromatography, 10 g prepacked column, DCM/MeOH 99/1 to afford 2-(2-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetonitrile **4.167** (50mg, Yield= 81%) as a white oil. LC₂ : Rt= 2.63

MS m/z (ES) [M+H]⁺= 383.0

¹H-NMR (DMSO-d₆), δ (ppm): 7.58-7.36 (4H, m), 7.44-7.36 (4H, m), 5 (2H, s), 4.02 (2H, s), 3.8 (3H, s)

[0139] The compounds in the following Tables have been synthesized according to the previous examples, as denoted in the column as "Exp. Nr". The compounds denoted with the asterisk have been exemplified in the Examples. Compounds 1.01 to 1.07 inclusive with E = -CN are not compounds of general Formula III as defined in the attached claims.

Table 1:

			
Compound Nr	Exp. Nr.		
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1.02	1.01		

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(continued)

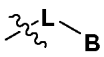
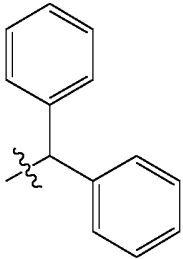
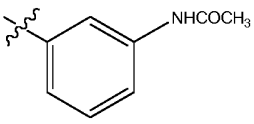
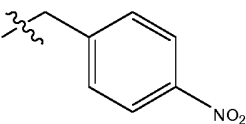
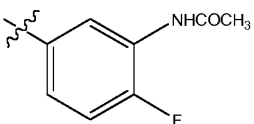
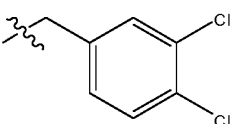
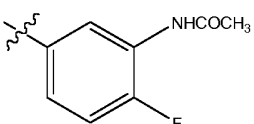
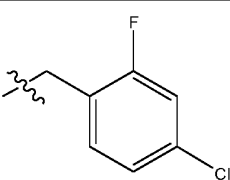
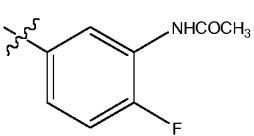
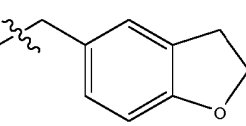
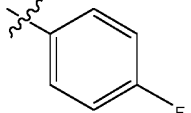
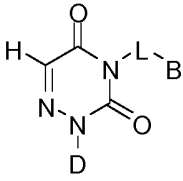
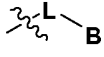
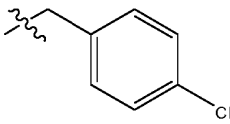
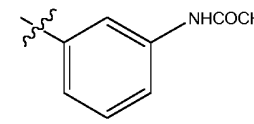
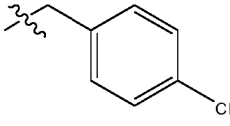
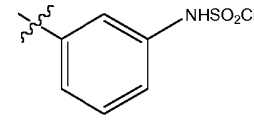
Compound Nr	Exp. Nr.		
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1.05	1.01		
1.06	1.01		
1.07	1.01		

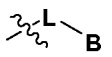
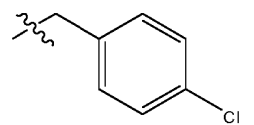
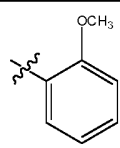
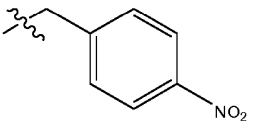
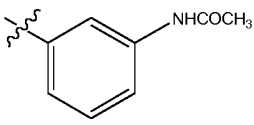
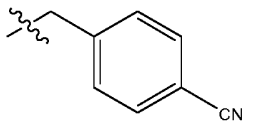
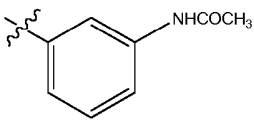
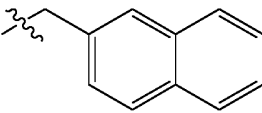
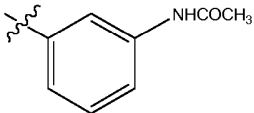
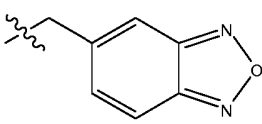
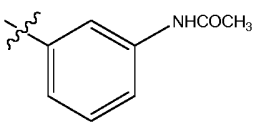
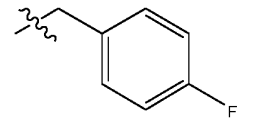
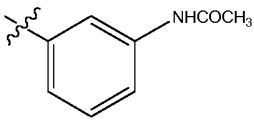
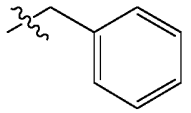
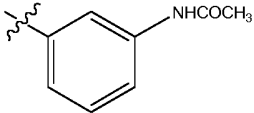
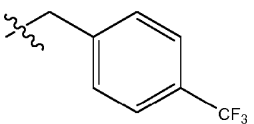
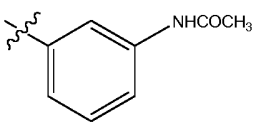
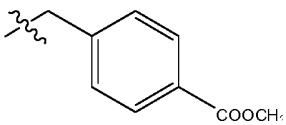
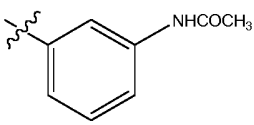
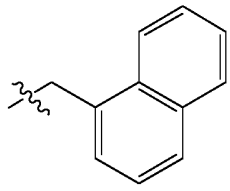
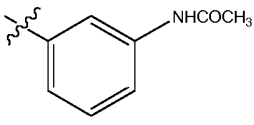
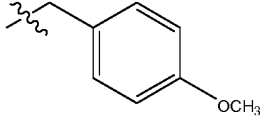
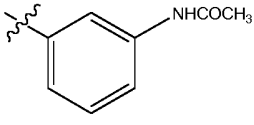
Table 2:

			
Compound Nr	Exp. Nr.		
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2.02	2.01		

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(continued)

5
10
15
20
25
30
35
40
45
50
55

Compound Nr	Exp. Nr.		
2.03	2.01		
2.04	2.01		
2.05*	2.05		
2.06	2.05		
2.07	2.05		
2.08	2.05		
2.09	2.05		
2.10	2.05		
2.11	2.05		
2.12	2.05		
2.13	2.05		

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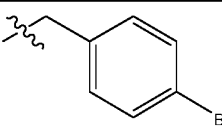
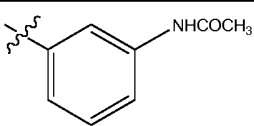
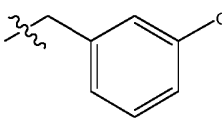
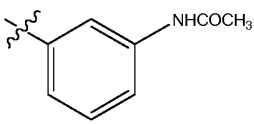
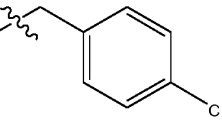
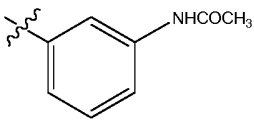
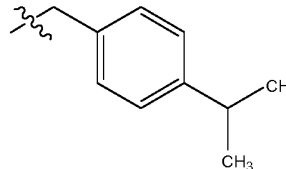
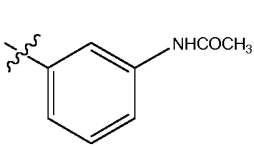
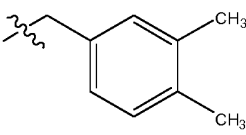
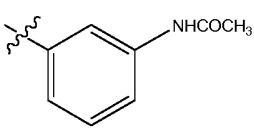
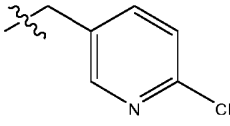
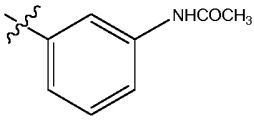
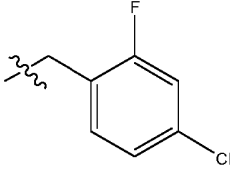
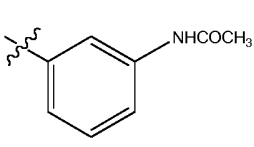
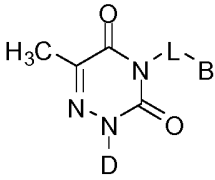
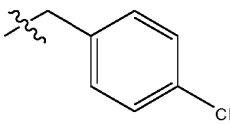
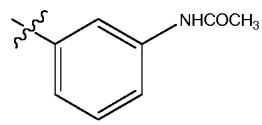
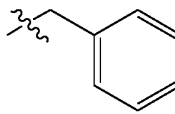
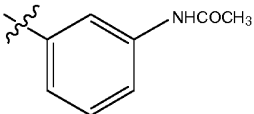
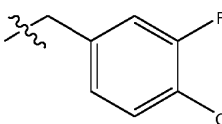
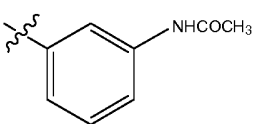
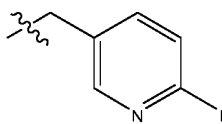
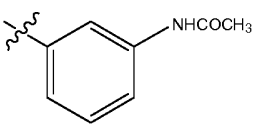
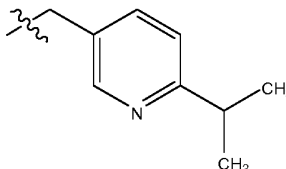
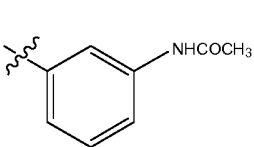
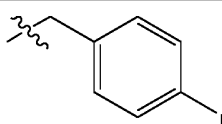
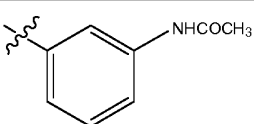
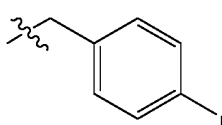
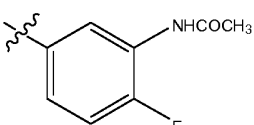
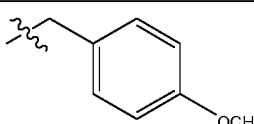
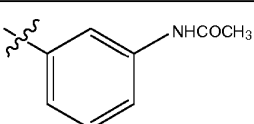
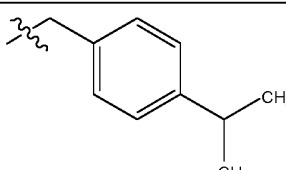
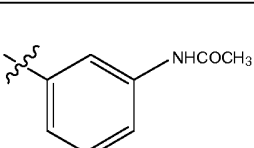
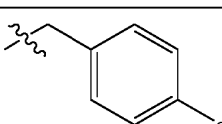
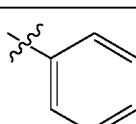
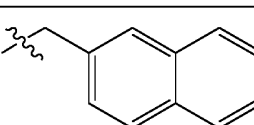
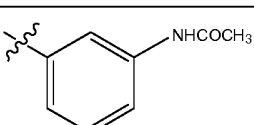
Compound Nr	Exp. Nr.		
2.14	2.05		
2.15	2.05		
2.16	2.05		
2.17	2.05		
2.18	2.05		
2.19	2.05		
2.20*	2.20		

Table 3:

			
Compound Nr	Exp. Nr.		
3.01*	3.01		

(continued)

Compound Nr	Exp. Nr.		
3.02	3.01		
3.03	3.01		
3.04	3.01		
3.05	3.01		
3.06	3.01		
3.07	3.01		
3.08	3.01		
3.09	3.01		
3.10	3.01		
3.11	3.01		

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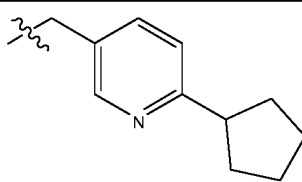
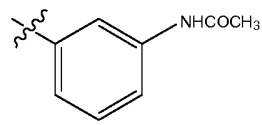
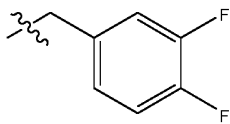
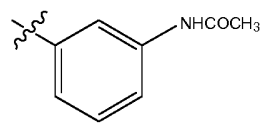
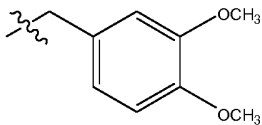
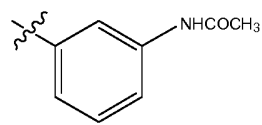
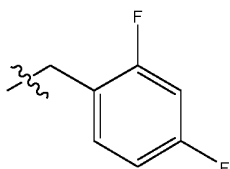
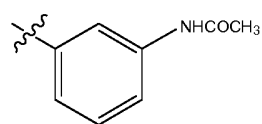
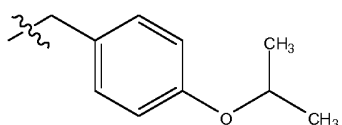
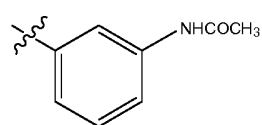
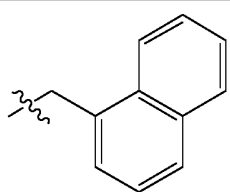
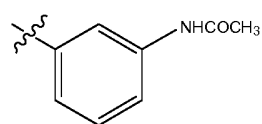
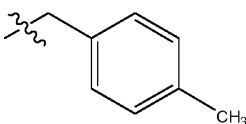
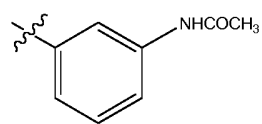
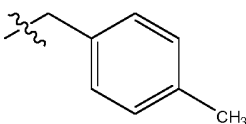
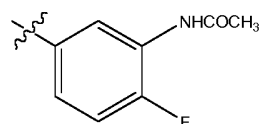
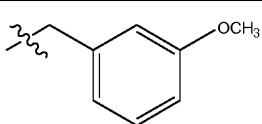
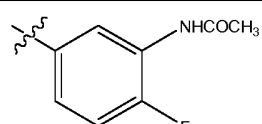
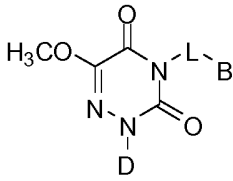
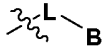
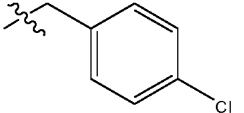
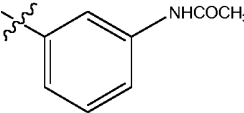
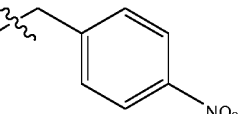
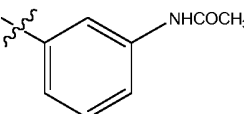
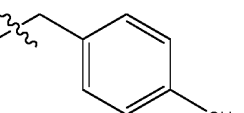
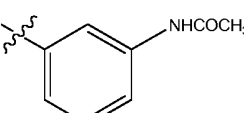
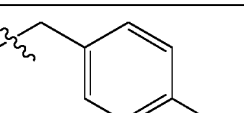
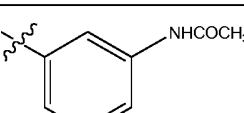
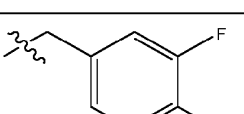
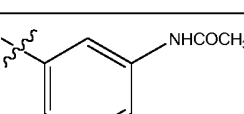
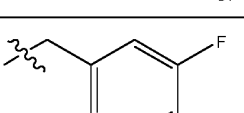
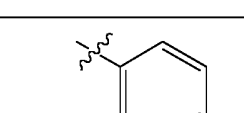
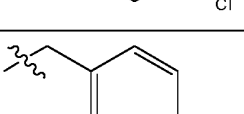
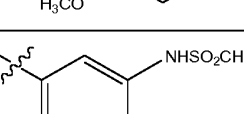
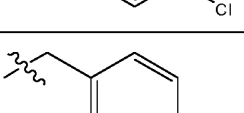
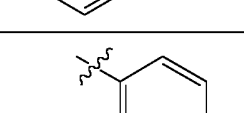
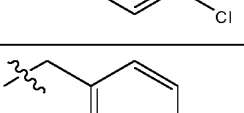
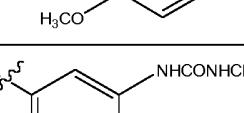
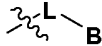
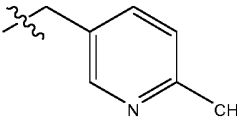
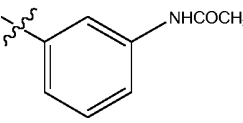
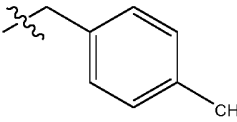
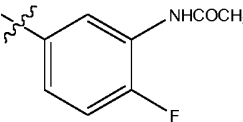
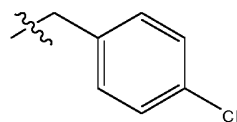
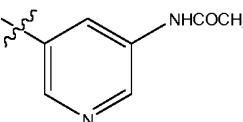
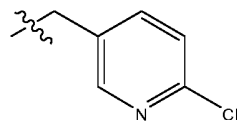
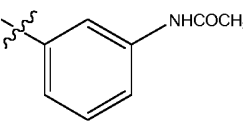
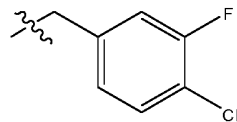
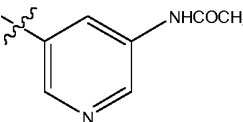
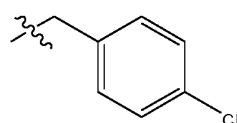
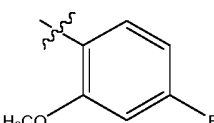
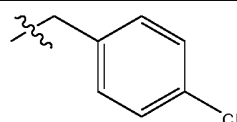
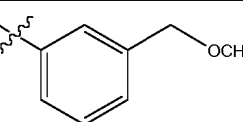
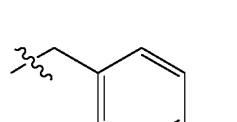
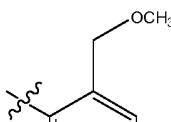
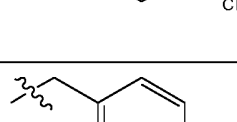
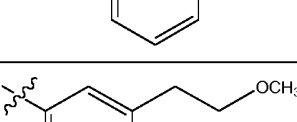
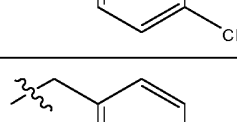
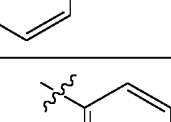
Compound Nr	Exp. Nr.		
3.12	3.01		
3.13	3.01		
3.14	3.01		
3.15	3.01		
3.16	3.01		
3.17	3.01		
3.18	3.01		
3.19	3.01		
3.20	3.01		

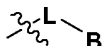
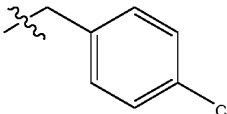
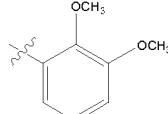
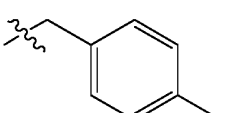
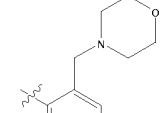
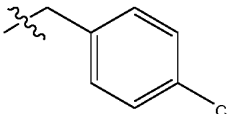
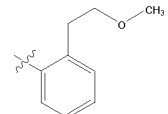
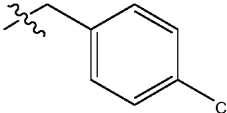
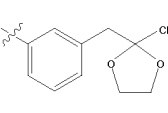
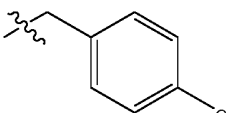
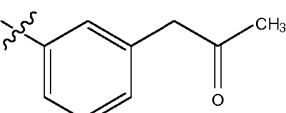
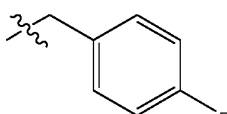
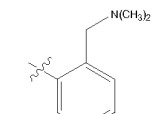
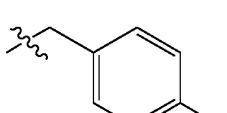
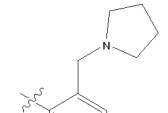
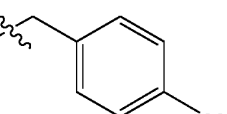
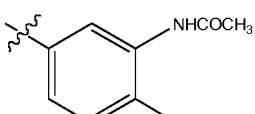
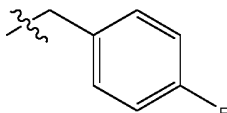
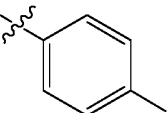
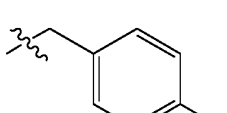
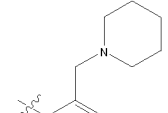
Table 4:

			
Compound Nr	Exp. Nr.		
4.01 *	4.01		
4.02	4.01		
4.03	4.01		
4.04	4.01		
4.05	4.01		
4.06	4.01		
4.07	4.01		
4.08	4.01		
4.09	4.01		

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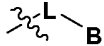
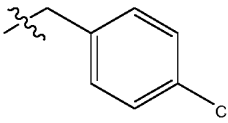
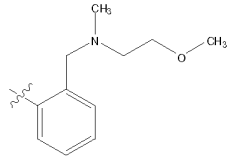
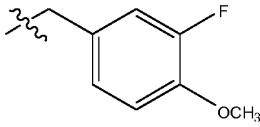
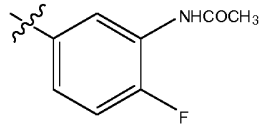
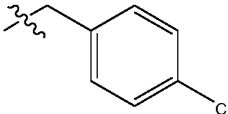
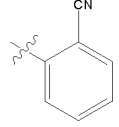
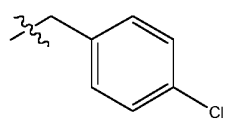
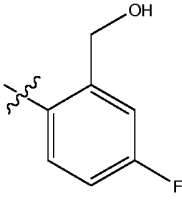
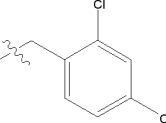
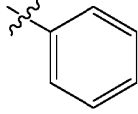
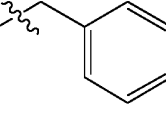
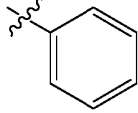
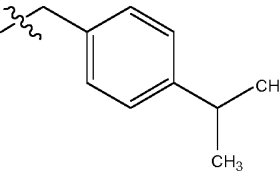
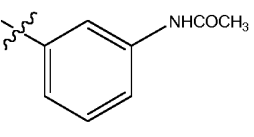
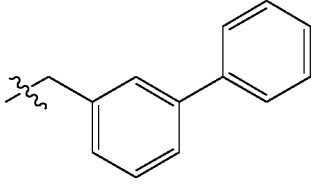
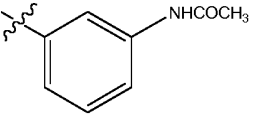
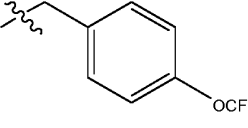
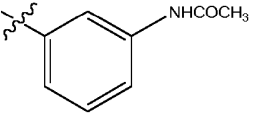
Compound Nr	Exp. Nr.		
4.10	4.01		
4.11	4.01		
4.12	4.01		
4.13	4.01		
4.14	4.01		
4.15	4.01		
4.16	4.01		
4.17	4.01		
4.18	4.01		
4.19	4.01		

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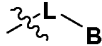
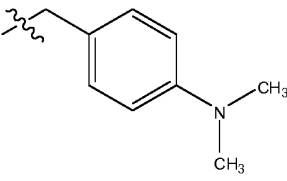
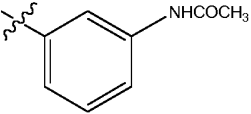
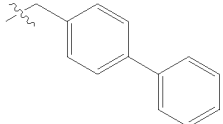
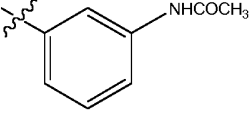
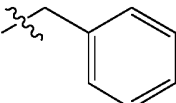
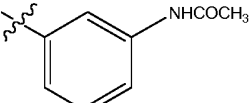
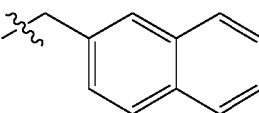
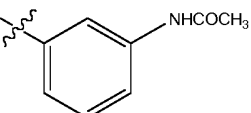
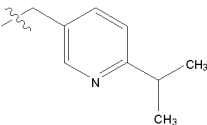
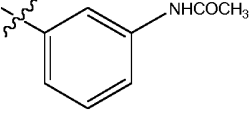
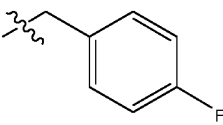
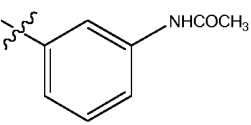
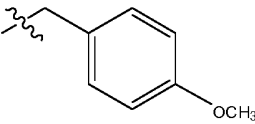
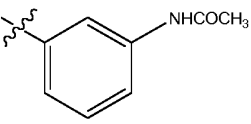
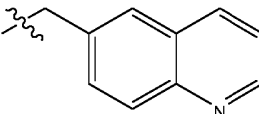
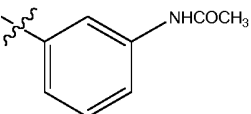
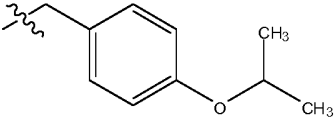
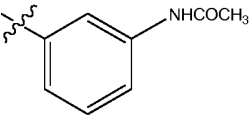
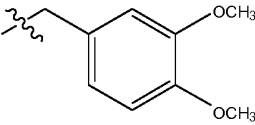
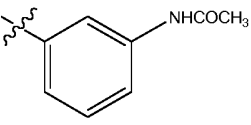
Compound Nr	Exp. Nr.		
4.20	4.01		
4.21	4.01		
4.22	4.01		
4.23	4.01		
4.24	4.01		
4.25	4.01		
4.26	4.01		
4.27	4.01		
4.28	4.01		
4.29	4.01		

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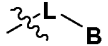
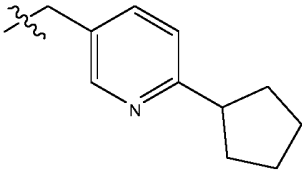
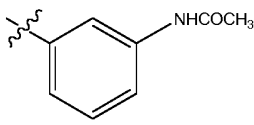
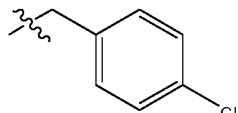
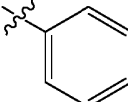
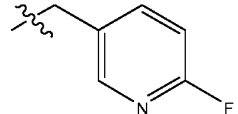
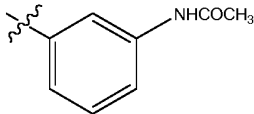
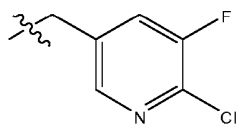
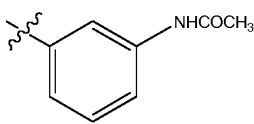
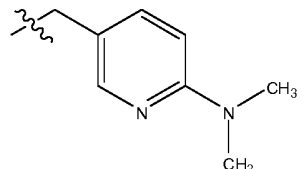
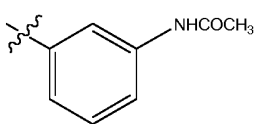
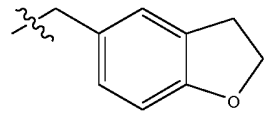
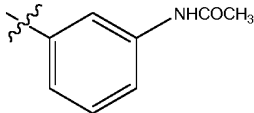
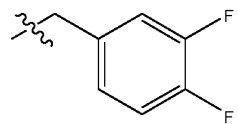
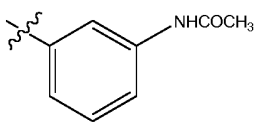
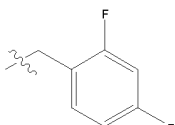
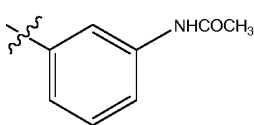
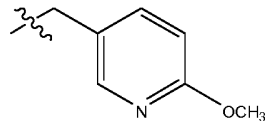
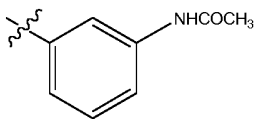
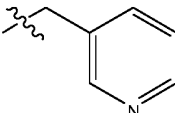
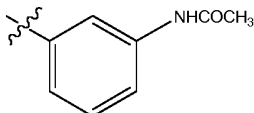
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Compound Nr	Exp. Nr.		
4.30	4.01		
4.31	4.01		
4.32	4.01		
4.33	4.01		
4.34	4.01		
4.35	4.01		
4.36*	4.36		
4.37	4.36		
4.38	4.36		

(continued)

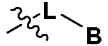
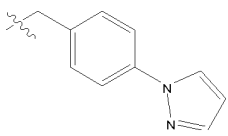
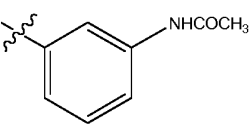
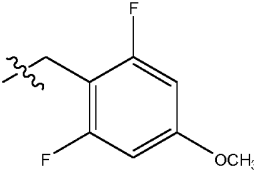
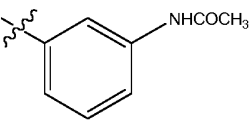
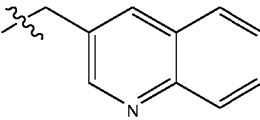
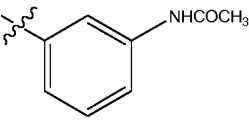
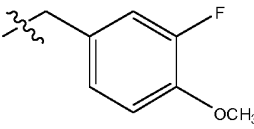
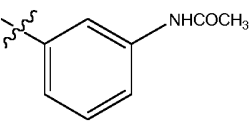
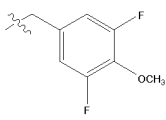
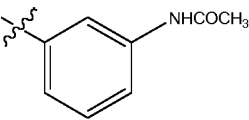
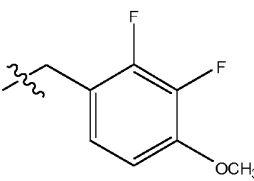
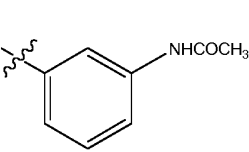
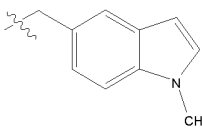
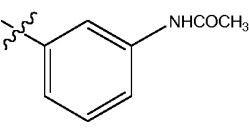
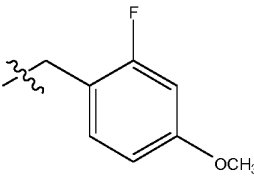
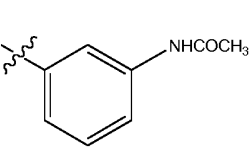
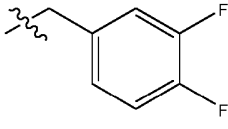
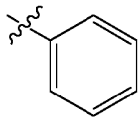
Compound Nr	Exp. Nr.		
4.39	4.36		
4.40	4.36		
4.41	4.36		
4.42	4.36		
4.43	4.36		
4.44	4.36		
4.45	4.36		
4.46	4.36		
4.47	4.36		
4.48	4.36		

(continued)

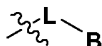
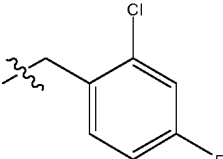
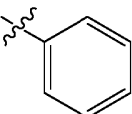
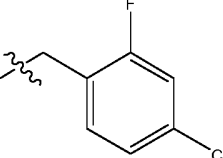
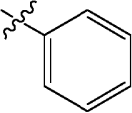
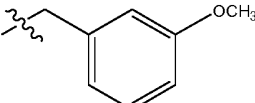
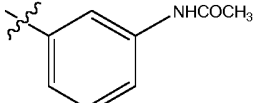
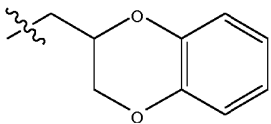
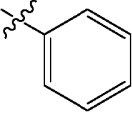
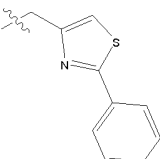
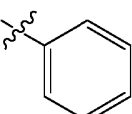
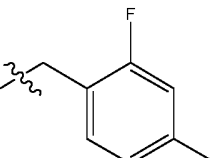
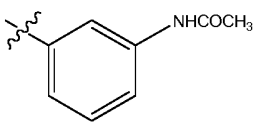
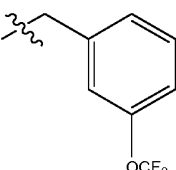
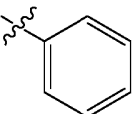
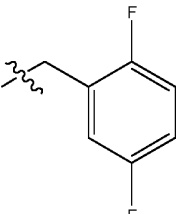
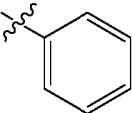
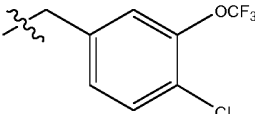
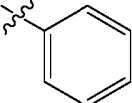
Compound Nr	Exp. Nr.		
4.49	4.36		
4.50	4.36		
4.51	4.36		
4.52	4.36		
4.53	4.36		
4.54	4.36		
4.55	4.36		
4.56	4.36		
4.57	4.36		
4.58	4.36		

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(continued)

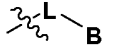
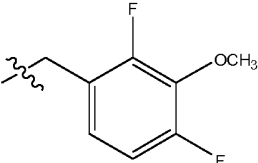
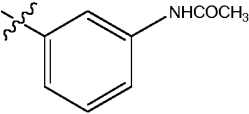
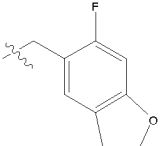
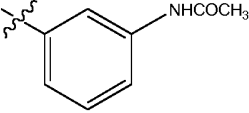
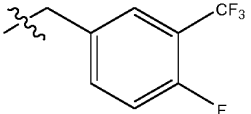
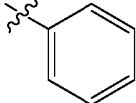
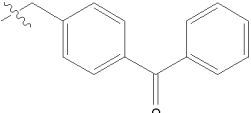
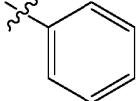
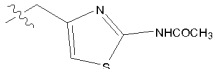
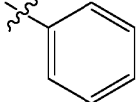
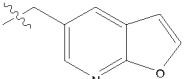
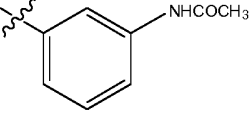
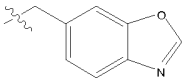
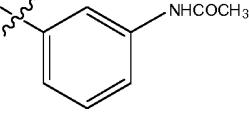
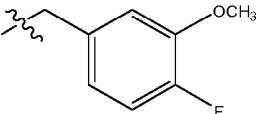
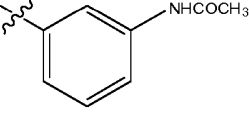
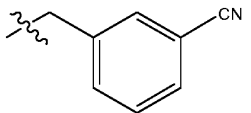
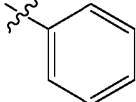
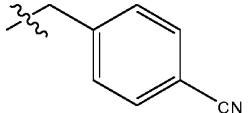
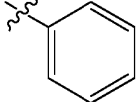
Compound Nr	Exp. Nr.		
4.59	4.36		
4.60	4.36		
4.61	4.36		
4.62	4.36		
4.63	4.36		
4.64	4.36		
4.65	4.36		
4.66	4.36		
4.67	4.36		

(continued)

Compound Nr	Exp. Nr.		
4.68	4.36		
4.69	4.36		
4.80	4.36		
4.90	4.36		
4.91	4.36		
4.92	4.36		
4.93	4.36		
4.94	4.36		
4.95	4.36		

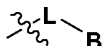
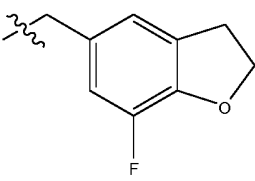
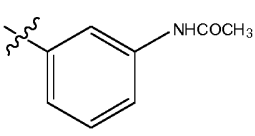
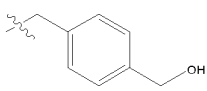
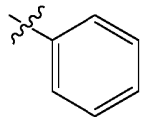
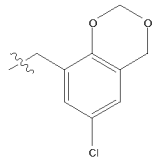
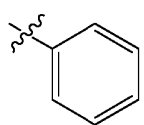
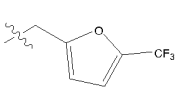
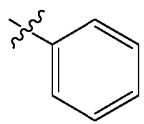
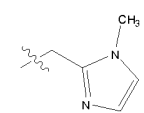
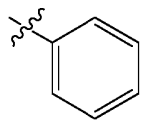
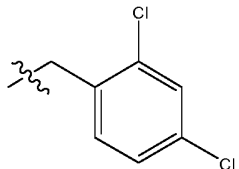
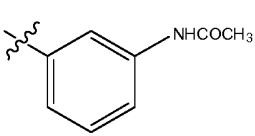
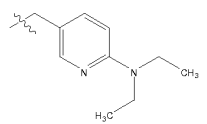
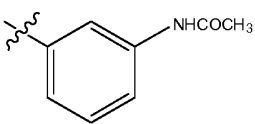
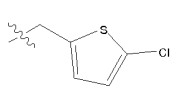
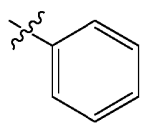
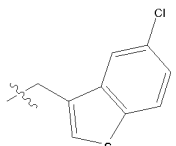
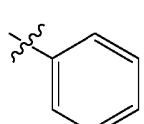
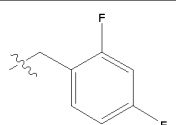
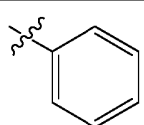
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(continued)

Compound Nr	Exp. Nr.		
4.96	4.36		
4.97	4.36		
4.98	4.36		
4.99	4.36		
4.100	4.36		
4.101	4.36		
4.102	4.36		
4.103	4.36		
4.104	4.36		
4.105	4.36		

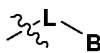
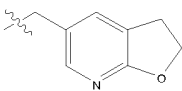
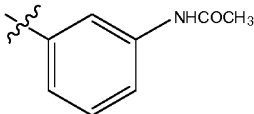
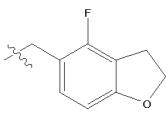
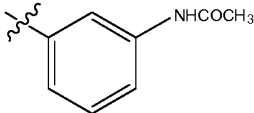
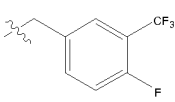
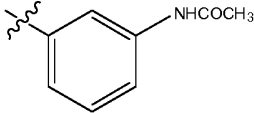
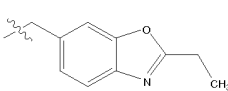
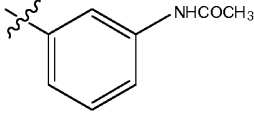
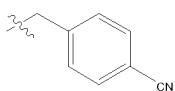
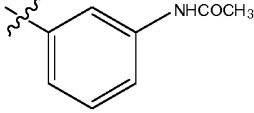
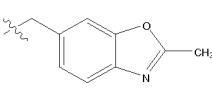
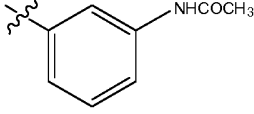
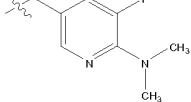
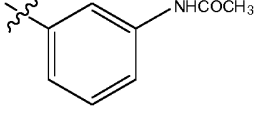
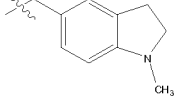
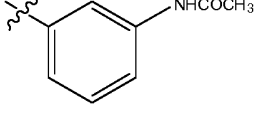
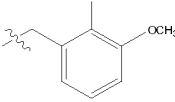
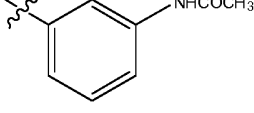
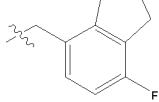
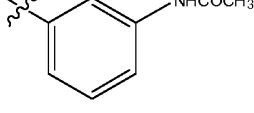
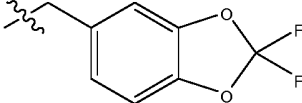
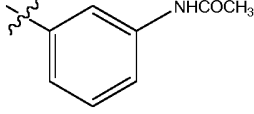
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(continued)

Compound Nr	Exp. Nr.		
4.106	4.36		
4.107	4.36		
4.108	4.36		
4.109	4.36		
4.110	4.36		
4.111	4.36		
4.112	4.36		
4.113	4.36		
4.114	4.36		
4.115	4.36		

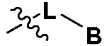
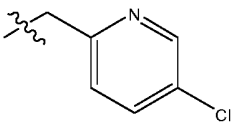
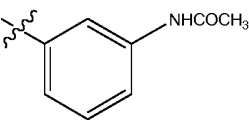
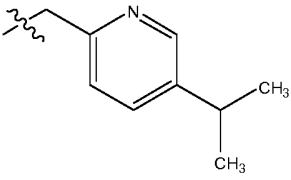
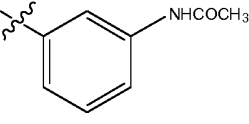
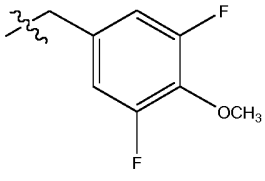
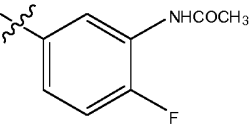
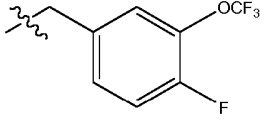
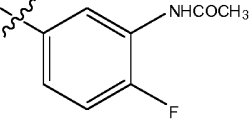
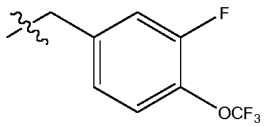
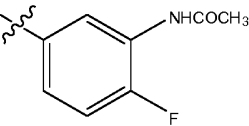
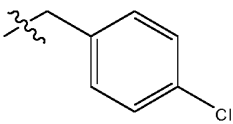
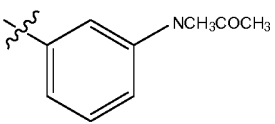
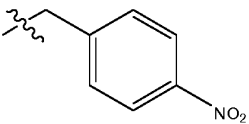
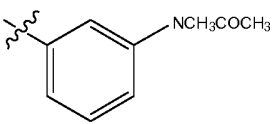
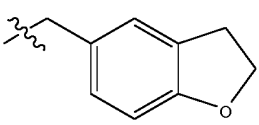
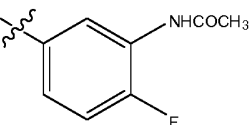
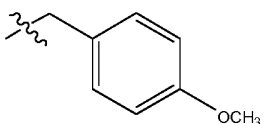
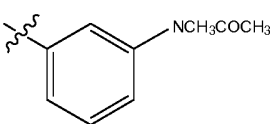
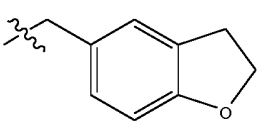
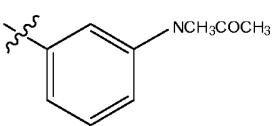
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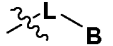
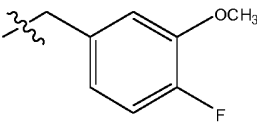
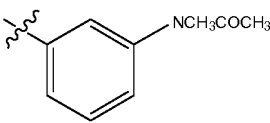
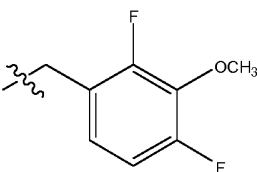
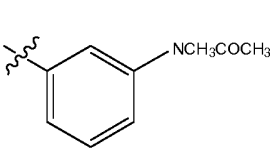
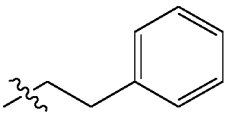
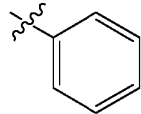
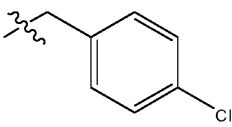
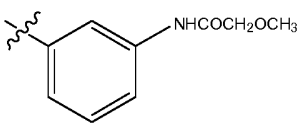
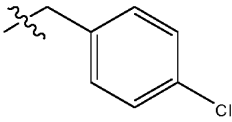
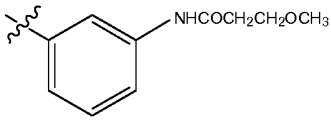
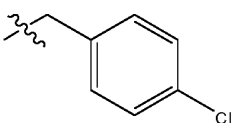
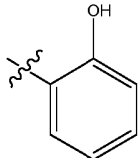
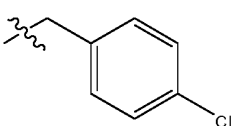
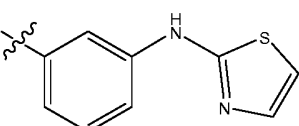
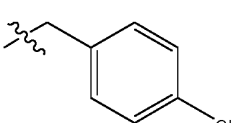
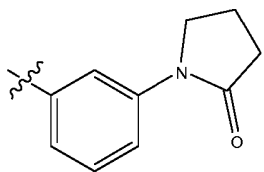
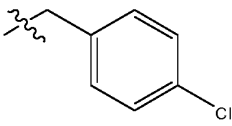
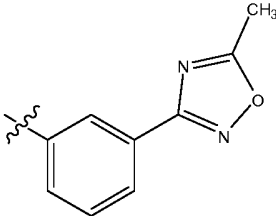
Compound Nr	Exp. Nr.		
4.116	4.36		
4.117	4.36		
4.118	4.36		
4.119	4.36		
4.120	4.36		
4.121	4.36		
4.122	4.36		
4.124	4.36		
4.125	4.36		
4.126	4.36		
4.127 *	4.127		

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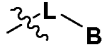
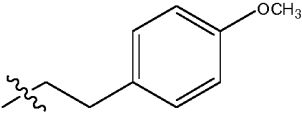
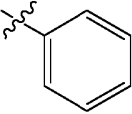
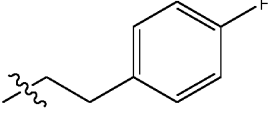
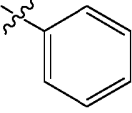
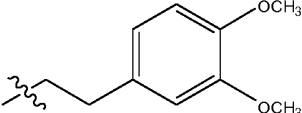
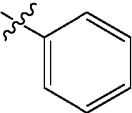
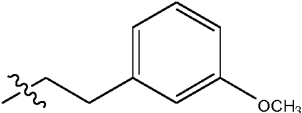
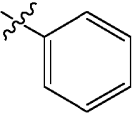
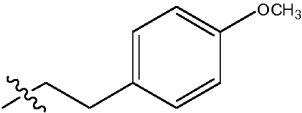
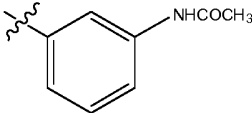
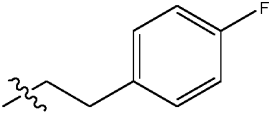
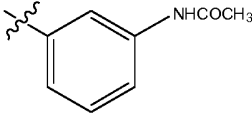
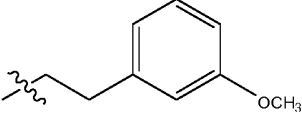
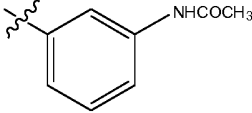
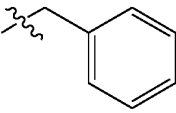
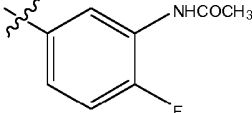
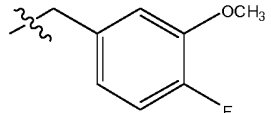
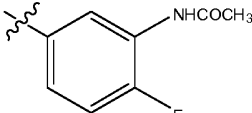
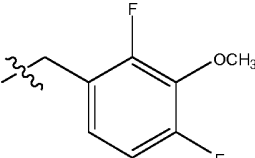
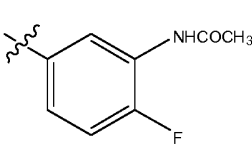
(continued)

Compound Nr	Exp. Nr.		
4.128	4.127		
4.129	4.127		
4.130	4.36		
4.131	4.36		
4.132	4.36		
4.133*	4.133		
4.134	4.133		
4.135	4.36		
4.136	4.133		
4.137	4.133		

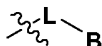
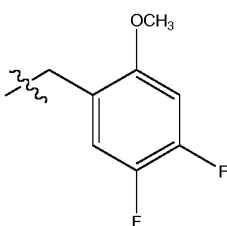
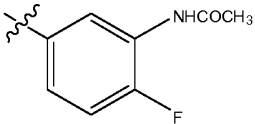
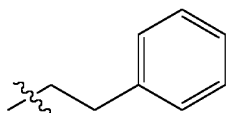
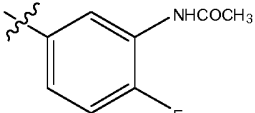
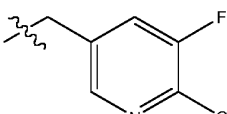
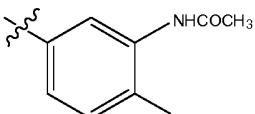
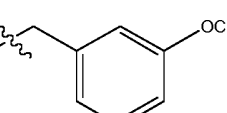
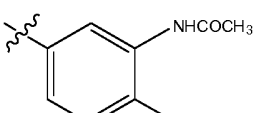
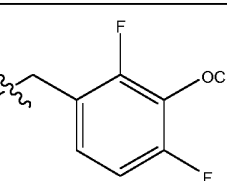
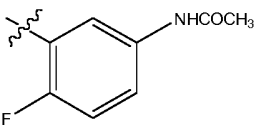
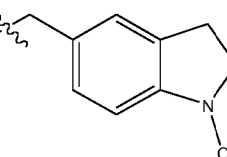
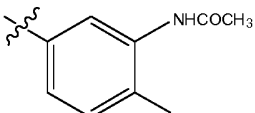
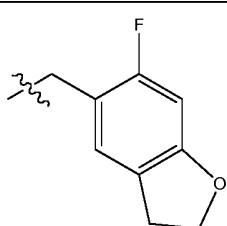
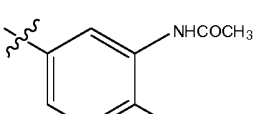
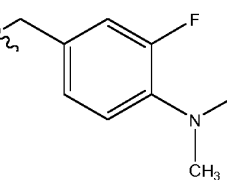
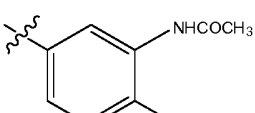
(continued)

Compound Nr	Exp. Nr.		
4.138	4.133		
4.139	4.133		
4.140	4.36		
4.141*	4.141		
4.142	4.141		
4.143*	4.143		
4.144*	4.144		
4.145*	4.145		
4.146*	4.146		

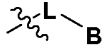
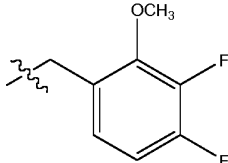
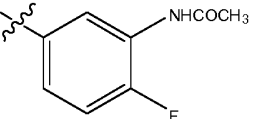
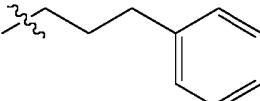
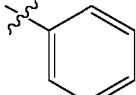
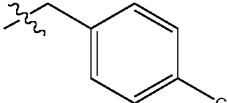
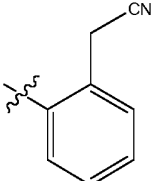
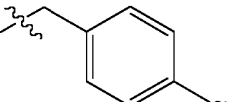
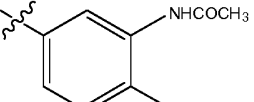
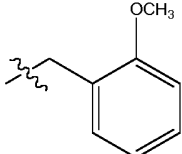
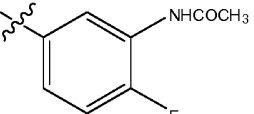
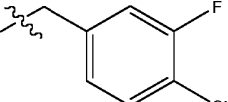
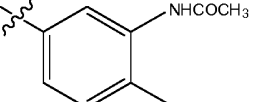
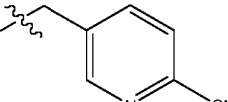
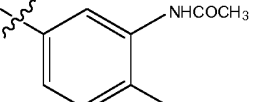
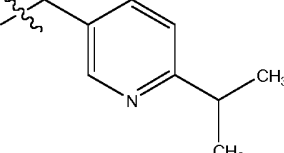
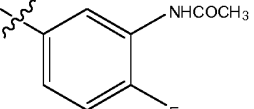
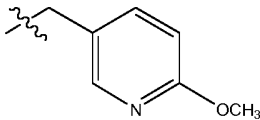
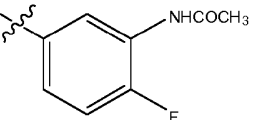
(continued)

Compound Nr	Exp. Nr.		
4.147	4.36		
4.148	4.36		
4.149	4.36		
4.150	4.36		
4.151	4.36		
4.152	4.36		
4.153	4.36		
4.154	4.36		
4.155	4.36		
4.156	4.36		

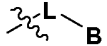
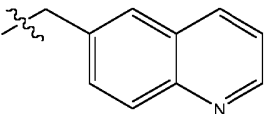
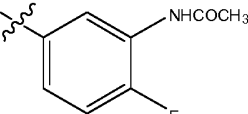
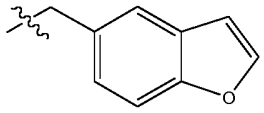
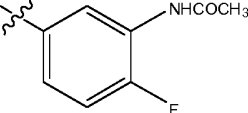
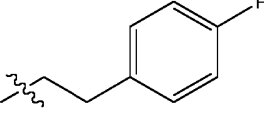
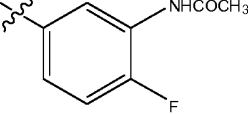
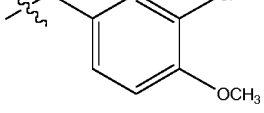
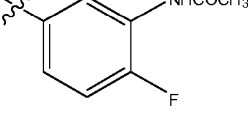
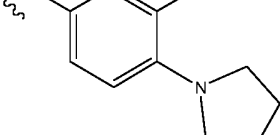
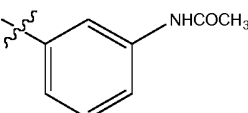
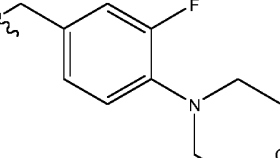
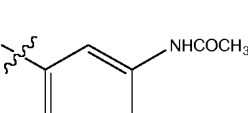
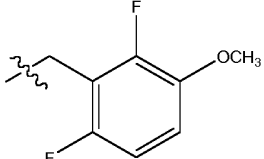
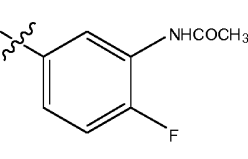
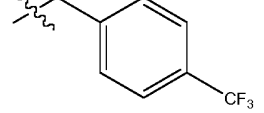
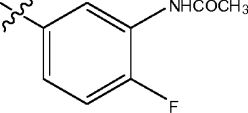
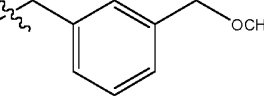
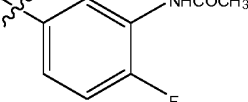
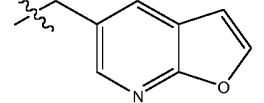
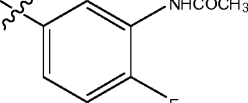
(continued)

Compound Nr	Exp. Nr.		
4.157	4.36		
4.158	4.36		
4.159	4.36		
4.160	4.36		
4.161	4.36		
4.162	4.36		
4.163	4.36		
4.164	4.36		

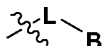
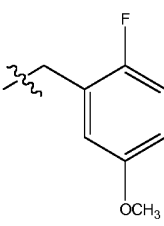
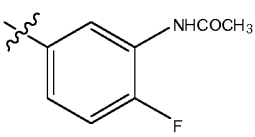
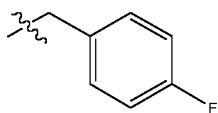
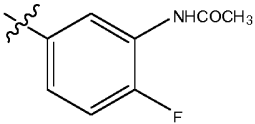
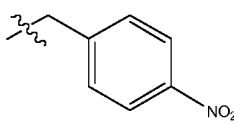
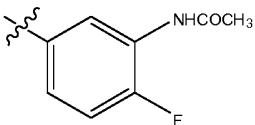
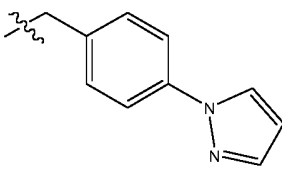
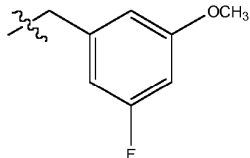
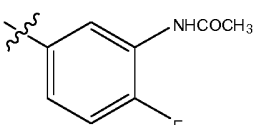
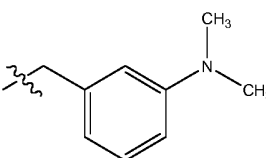
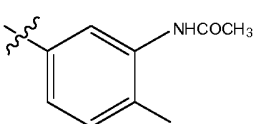
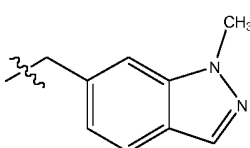
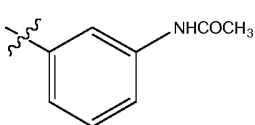
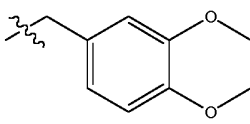
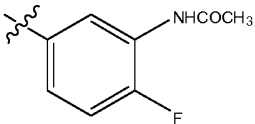
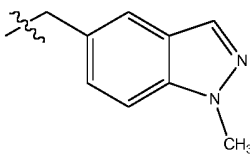
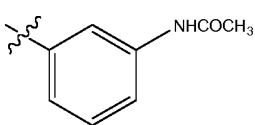
(continued)

Compound Nr	Exp. Nr.		
4.165	4.36		
4.166	4.36		
4.167*	4.167		
4.168	4.36		
4.169	4.36		
4.170	4.36		
4.171	4.36		
4.172	4.36		
4.173	4.36		

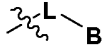
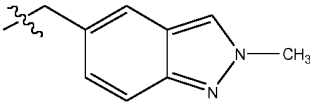
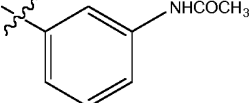
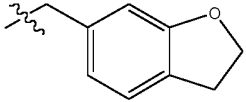
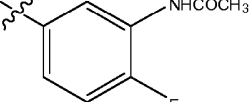
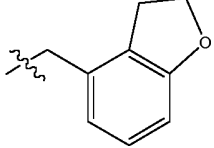
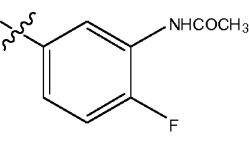
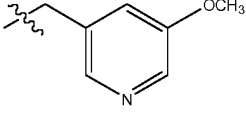
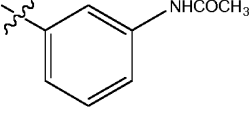
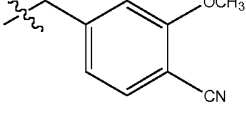
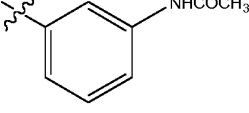
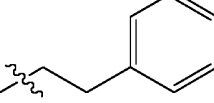
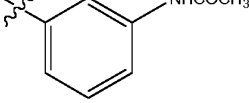
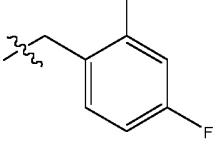
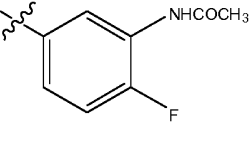
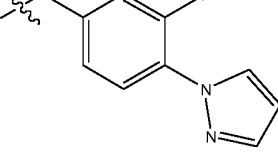
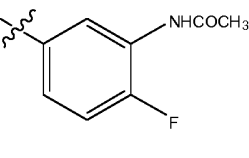
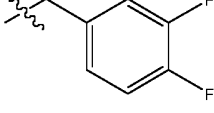
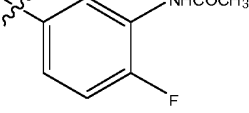
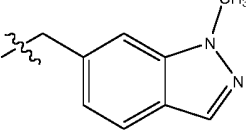
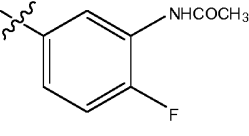
(continued)

Compound Nr	Exp. Nr.		
4.174	4.36		
4.175	4.36		
4.176	4.36		
4.177	4.36		
4.178	4.36		
4.179	4.36		
4.180	4.36		
4.181	4.36		
4.182	4.36		
4.183	4.36		

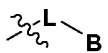
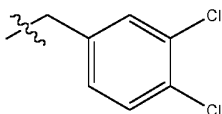
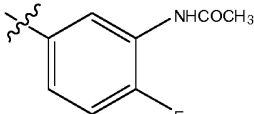
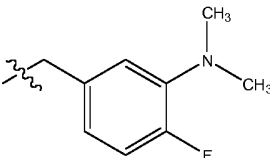
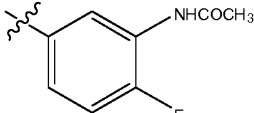
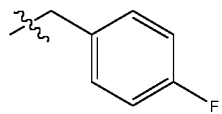
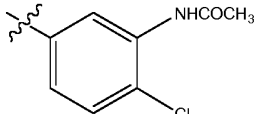
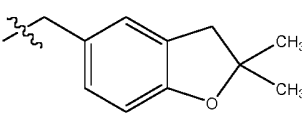
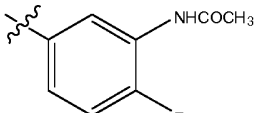
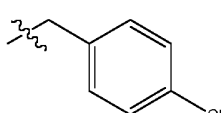
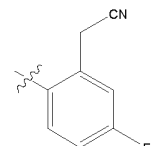
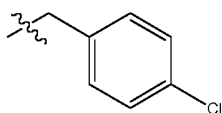
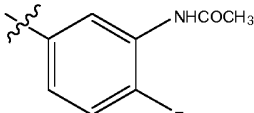
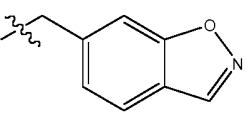
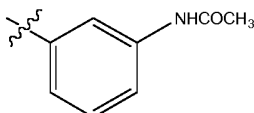
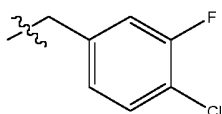
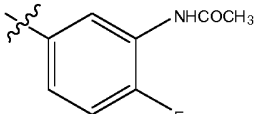
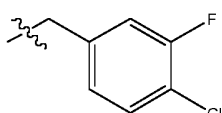
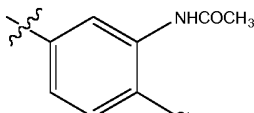
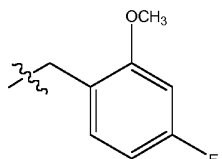
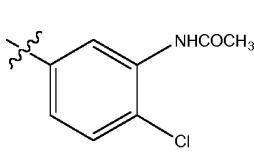
(continued)

Compound Nr	Exp. Nr.		
4.184	4.36		
4.185	4.36		
4.186	4.36		
4.187	4.36		
4.188	4.36		
4.189	4.36		
4.190	4.36		
4.191	4.36		
4.192	4.36		

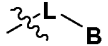
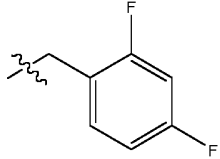
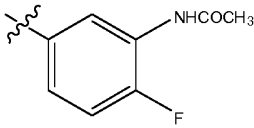
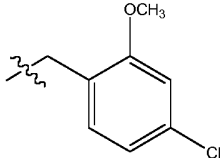
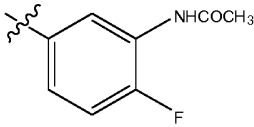
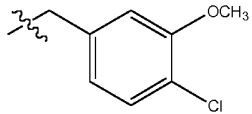
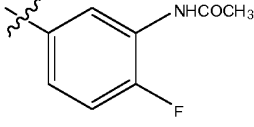
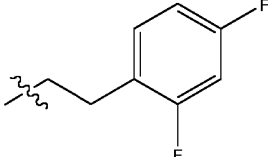
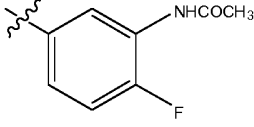
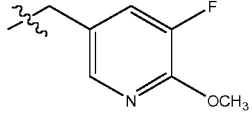
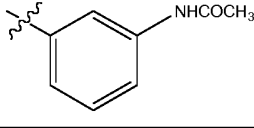
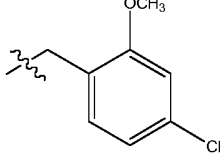
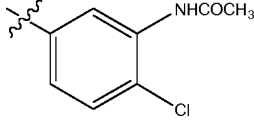
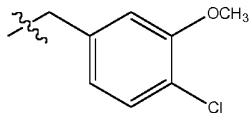
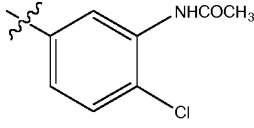
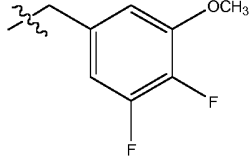
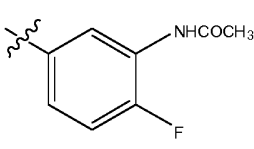
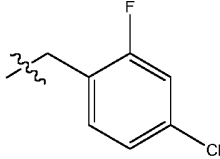
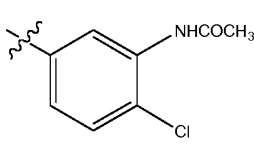
(continued)

Compound Nr	Exp. Nr.		
4.193	4.36		
4.194	4.36		
4.195	4.36		
4.196	4.36		
4.197	4.36		
4.198	4.36		
4.199	4.36		
4.200	4.36		
4.201	4.36		
4.202	4.36		

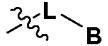
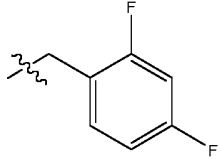
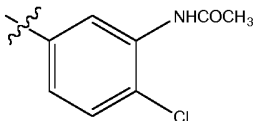
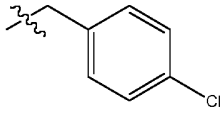
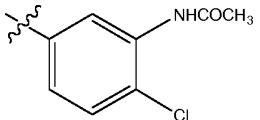
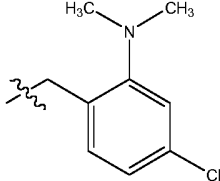
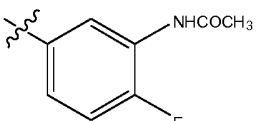
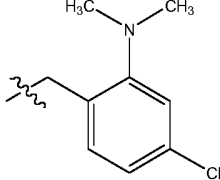
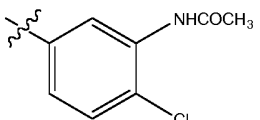
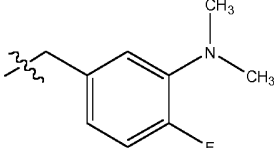
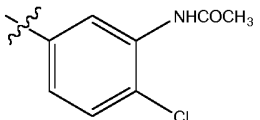
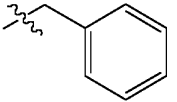
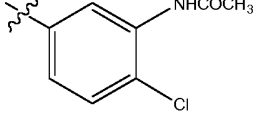
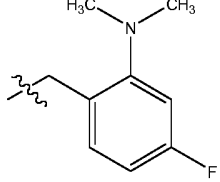
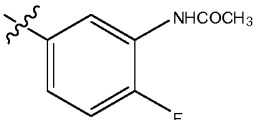
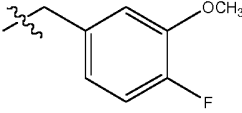
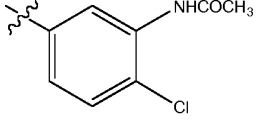
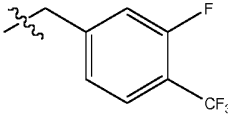
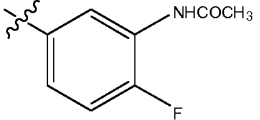
(continued)

Compound Nr	Exp. Nr.		
4.203	4.127		
4.204	4.36		
4.205	4.36		
4.206	4.36		
4.207	4.167		
4.208	4.36		
4.209	4.36		
4.210	4.127		
4.211	4.36		
4.212	4.36		

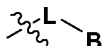
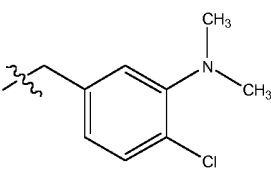
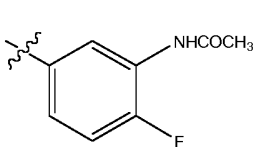
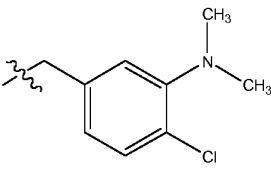
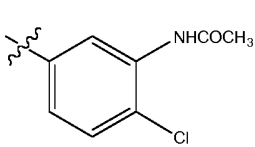
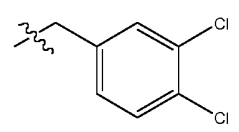
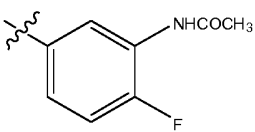
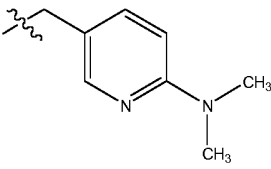
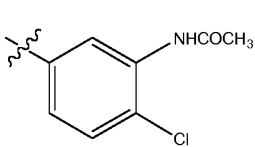
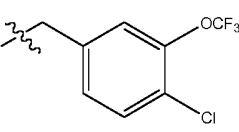
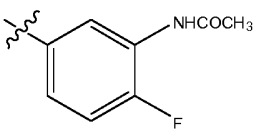
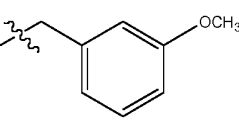
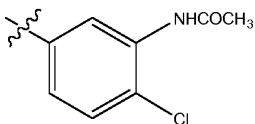
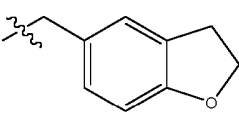
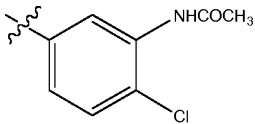
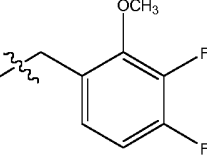
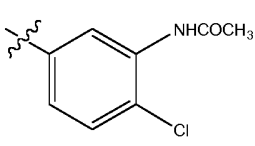
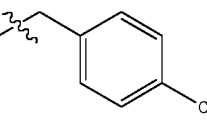
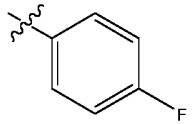
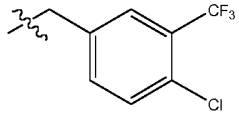
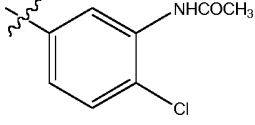
(continued)

Compound Nr	Exp. Nr.		
4.213	4.36		
4.214	4.36		
4.215	4.36		
4.216	4.36		
4.217	4.36		
4.219	4.36		
4.219	4.36		
4.220	4.36		
4.221	4.36		

(continued)

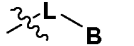
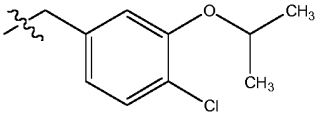
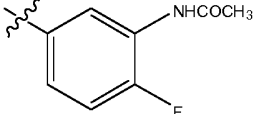
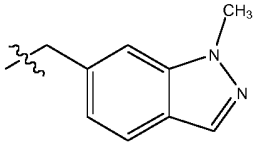
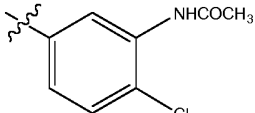
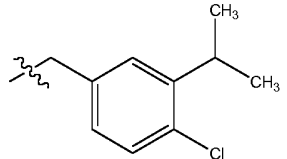
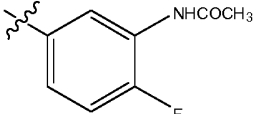
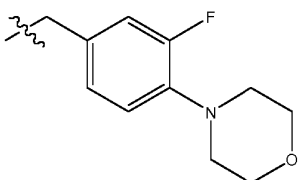
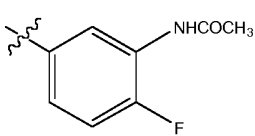
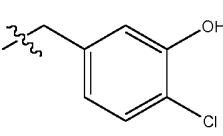
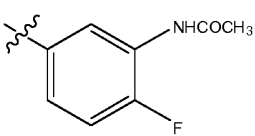
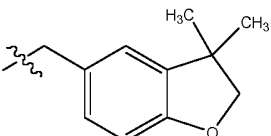
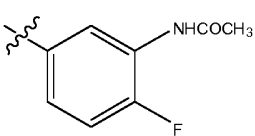
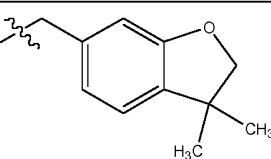
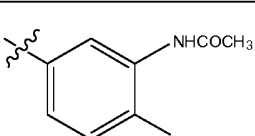
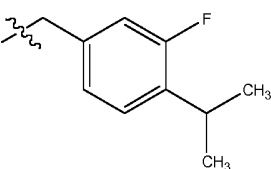
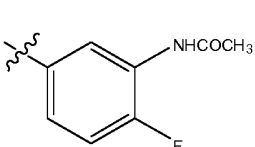
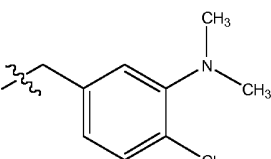
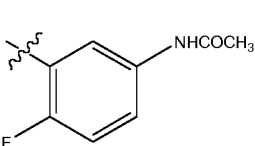
Compound Nr	Exp. Nr.		
4.222	4.36		
4.223	4.36		
4.224	4.36		
4.225	4.36		
4.226	4.36		
4.227	4.36		
4.228	4.36		
4.229	4.36		
4.230	4.211		

(continued)

Compound Nr	Exp. Nr.		
4.231	4.127		
4.232	4.36		
4.233	4.36		
4.234	4.36		
4.235	4.36		
4.236	4.36		
4.237	4.36		
4.238	4.36		
4.239	4.36		
4.240	4.36		

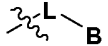
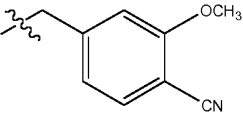
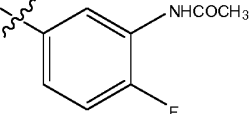
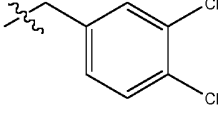
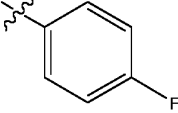
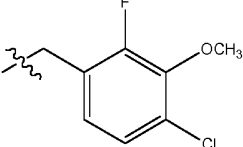
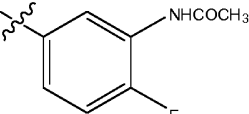
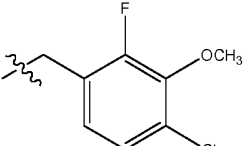
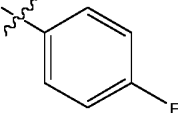
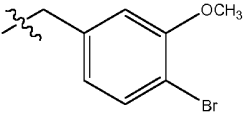
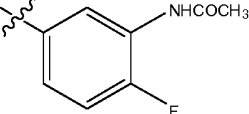
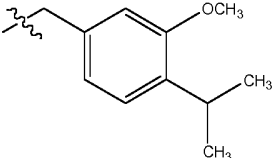
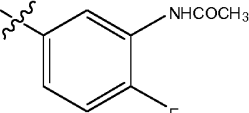
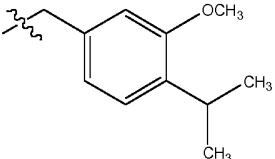
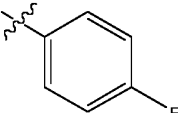
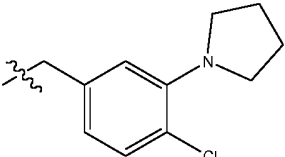
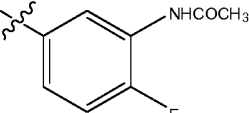
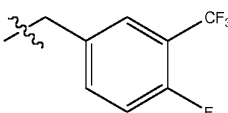
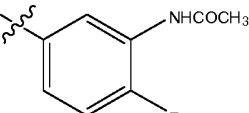
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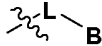
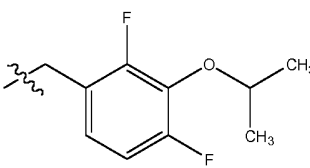
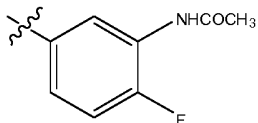
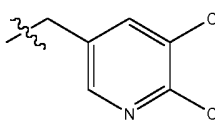
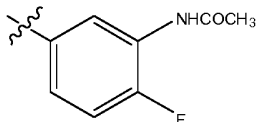
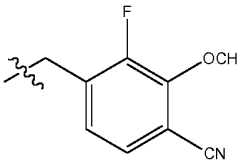
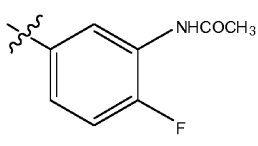
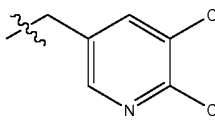
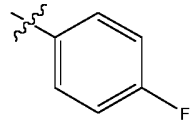
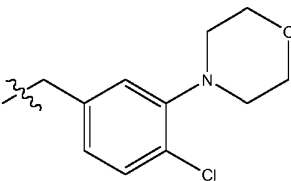
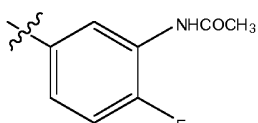
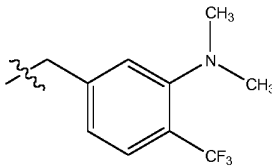
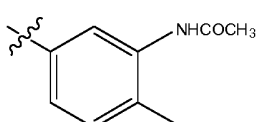
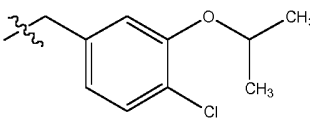
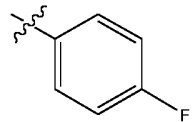
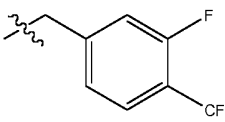
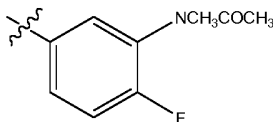
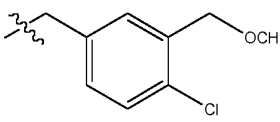
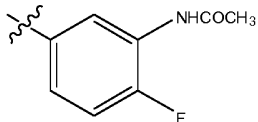
Compound Nr	Exp. Nr.		
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4.242	4.36		
4.243	4.36		
4.244	4.36		
4.245	4.36		
4.246	4.36		
4.247	4.36		
4.248	4.36		
4.249	4.36		

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(continued)

Compound Nr	Exp. Nr.		
4.250	4.36		
4.251	4.36		
4.252	4.36		
4.253	4.36		
4.254	4.36		
4.255	4.36		
4.256	4.36		
4.257	4.36		
4.258	4.36		

(continued)

Compound Nr	Exp. Nr.		
4.259	4.36		
4.260	4.36		
4.261	4.36		
4.262	4.36		
4.263	4.36		
4.264	4.36		
4.265	4.36		
4.266	4.133		
4.267	4.36		

(continued)

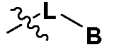
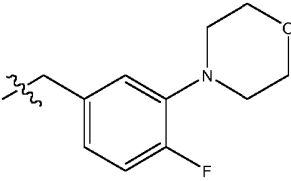
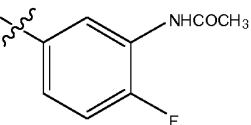
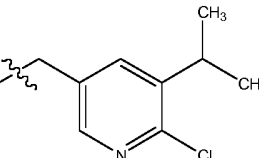
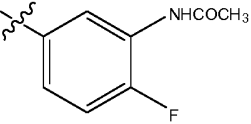
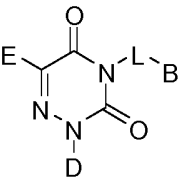
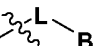
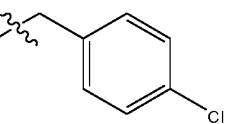
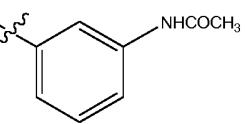
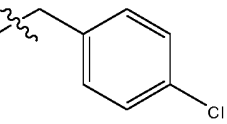
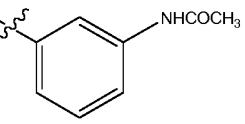
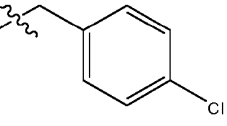
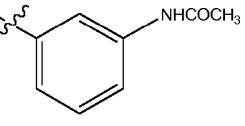
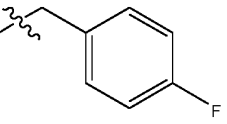
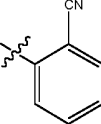
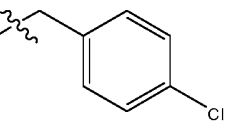
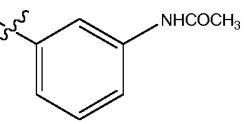
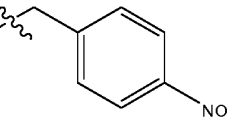
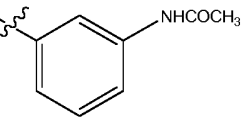
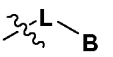
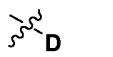
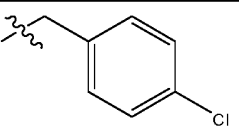
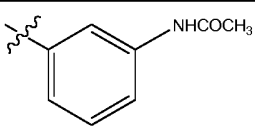
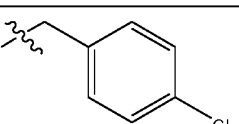
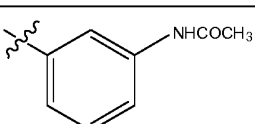
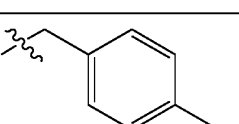
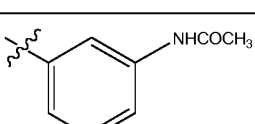
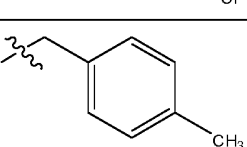
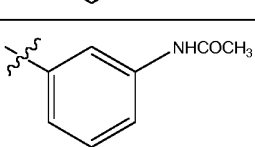
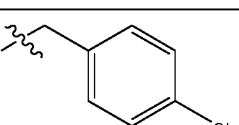
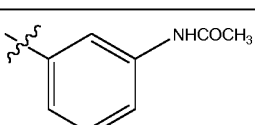
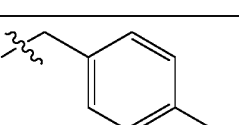
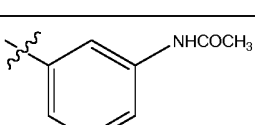
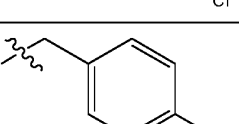
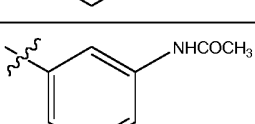
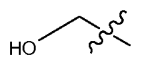
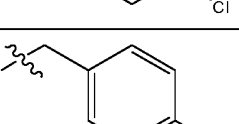
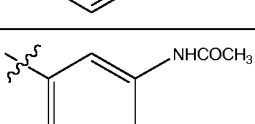
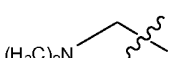
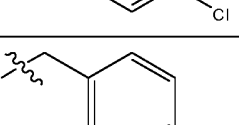
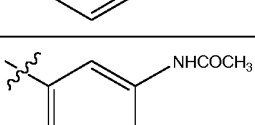
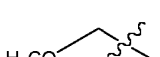
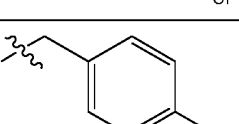
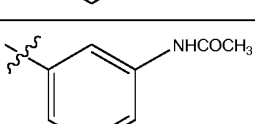
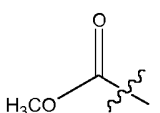
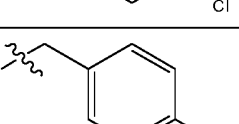
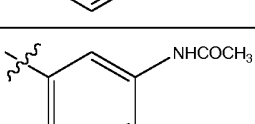
Compound Nr	Exp. Nr.		
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4.269	4.36		

Table 5:

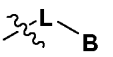
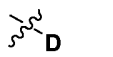
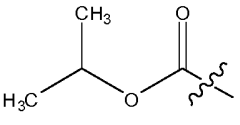
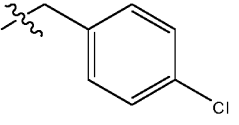
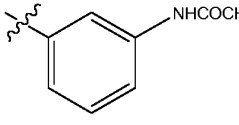
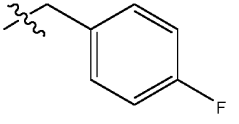
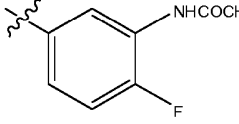
 E is not H, CN, CH ₃ or OCH ₃				
Compound Nr	Exp. Nr.	E		
5.01 *	5.01	iPr		
5.02	5.01	Et		
5.03	5.01	nBu		
5.04	4.01	Br-		
5.05	4.01	(CH ₃) ₂ N(CH ₂) ₂ O-		
5.06	4.01	CH ₃ O(CH ₂) ₂ O-		

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(continued)

Compound Nr	Exp. Nr.	E		
5.07	4.01	CH ₃ O(CH ₂) ₂ O-		
5.08	4.01	CH ₃ NH-		
5.09	4.01	CH ₃ CH ₂ O-		
5.10	4.01	Br-		
5.11	4.01	(CH ₃) ₂ CHO-		
5.12	4.01	Br-		
5.13*	5.13	(CH ₃) ₂ N-		
5.14*	5.14			
5.15*	5.15			
5.16*	5.16			
5.17*	5.17			

(continued)

Compound Nr	Exp. Nr.	E		
5.18*	5.18			
5.19	4.01	CH ₃ O(CH ₂) ₂ O-		

PHYSICOCHEMICAL DATA

[0140] ¹H NMR spectra were recorded on a Bruker 500MHz. Chemical shifts are expressed in parts of million (ppm, δ units). Coupling constants are in units of hertz (Hz) Splitting patterns describe apparent multiplicities and are designated as s (singlet), d (doublet), t (triplet), q (quadruplet), quint (quintuplet), m (multiplet). LCMS were recorded on a Waters Micromass ZQ 2996 system by the following conditions.

Method LC₁:

[0141] Column 3.0*50mm stainless steel packed with 3.5μm XTerra RP C-18; flow rate 1mL/min; mobile phase: A phase = 0.1% formic acid in water, B phase = 0.07% formic acid in acetonitrile. 0-0.5min (A: 95%, B: 5%), 0.5-6.0min (A: 0%, B: 100%), 6.0-6.5min (A: 95%, B: 5%), 6.5-7min (A: 95%, B: 5%); UV detection Diode Array: 200-400nm; Injection volume: 10μL.

Method LC₂:

[0142] Column 4.6*30mm stainless steel packed with 1.8μm Zorbax SB C-18; flow rate 1.5mL/min; mobile phase: A phase = 0.05% formic acid in water, B phase = 0.05% formic acid in acetonitrile. 0-3.5min (A: 0%, B: 10%), 3.5-3.7min (A: 0%, B: 100%), 3.8-4.5min (A: 90%, B: 10%); Oven temperature: 30°C ± 1°C; UV detection Diode Array: 200-400nm; Injection volume: 10μL.

[0143] All mass spectra were taken under electrospray ionisation (ESI) methods.

[0144] MS :

Source

- Voltage * capillarity= 3.31 (kV)
- * cone= 30 (V)
- * extraction= 6 (V)
- * RF lens= 0.9 (V)
- Temperature * source temperature= 140°C
- * desolvation temperature= 300°C

Gas flow (N₂):

- Desolvation= 250 (L/hour)
- Cone= 50 (L/hour)

[0145] Most of the reactions were monitored by thin-layer chromatography on 0.25mm Macherey-Nagel silica gel plates (60F-2254), visualized with UV light. Flash column chromatography was performed on silica gel (220-440 mesh, Fluka). Melting point determination was performed on a Buchi B-540 apparatus. Compounds 1.01 to 1.07 inclusive with E = -CN are not compounds of general Formula III as defined in the attached claims.

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Table 6:

Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
1.01	196	-	394.0	1	4.35	Yellow solid
1.02	127-128	-	412.0	1	4.43	White solid
1.03	118-119	-	436.0	1	4.50	Yellow solid
1.04	228-229	-	423.0	1	4.05	Yellow solid
1.05	176-177	448.0	446.0	1	4.51	Yellow solid
1.06	135-136	432.0	430.0	1	4.31	Yellow solid
1.07	85-91	-	-	2	2.78	Yellow solid
2.01	171-172	371.0	369.0	1	4.02	Yellow solid
2.02	-	-	405.0	1	4.11	White solid
2.03	-	344.0	-	1	4.51	Beige oil
2.04	229-230	382.0	380.0	1	3.64	White solid
2.05	184-185	362.0	360.0	1	3.47	White solid
2.06	189-190	387.0	385.0	1	4.08	White solid
2.07	163-164	379.0	377.0	1	3.60	White solid
2.08	135-138	355.3	-	1	3.71	Yellow solid
2.09	80-81	337.0	335.0	1	3.61	Yellow solid
2.10	80-82	405.2	-	1	4.06	Yellow solid
2.11	92-93	395.3	-	1	3.58	Yellow solid
2.12	96-97	387.2	-	1	4.04	Yellow solid
2.13	85-86	367.2	-	1	3.62	Yellow solid
2.14	168-169	415.0	413.0	1	4.0	Yellow solid
2.15	160-161	371.0	369.0	1	3.90	White solid
2.16	195-196	351.0	349.0	1	3.85	White solid
2.17	135-136	379.0	377.0	1	4.32	White solid
2.18	-	365.0	363.0	1	4.05	Colorless oil
2.19	-	372.0	370.0	1	3.29	White solid
2.20	184	389.2		1	3.98	Yellow solid
3.01	168	384.0	-	1	4.08	White solid
3.02	204-206	351.1	-	2	2.23	White solid
3.03	147-148	402.9	400.9	2	2.49	White solid
3.04	180-181	370.0	368.0	2	1.91	White solid
3.05	191-193	394.1	-	2	1.49	White solid
3.06	162	369.1	-	2	2.29	White solid
3.07	128-130	387	-	2	2.31	White solid
3.08	165	381.1	-	2	2.23	White solid
3.09	155	393.0	-	2	2.74	White solid
3.10	98-99	328.1	-	2	2.94	White solid
3.11	182-184	401.1	-	2	2.61	White solid

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(continued)

	Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
5	3.12	169-170	420.2	418.2	2	1.65	White solid
	3.13	80-84	387.1	385.1	2	2.23	White solid
	3.14	90-92	411.1	-	2	2.02	White solid
	3.15	172-174	387.0	-	2	2.29	White solid
10	3.16	190-191	409.1	407.1	2	2.52	White solid
	3.17	150-151	401.9	399.1	2	2.50	White solid
	3.18	172	365.0	-	1	4.01	White solid
15	3.19	185-186	383.0	381.0	2	2.44	White solid
	3.20	134-135	399.0	397.9	2	2.26	White solid
	4.01	190-191	400.9	-	1	3.98	White solid
	4.02	150-151	310.0	-	1	3.67	White solid
20	4.03	190-191	381.0	-	1	3.89	White solid
	4.04	120	434.9	-	1	4.10	Beige solid
	4.05	194-196	418.9	-	1	4.03	White solid
25	4.06	75	391.9	-	1	4.55	White solid
	4.07	120	434.9	-	1	4.11	White solid
	4.08	150-151	373.8	-	1	4.50	White solid
	4.09	234-236	429.9	-	2	2.42	White solid
30	4.10	110	382.0	-	2	1.14	Beige solid
	4.11	159-160	399.0	397.0	2	2.35	White solid
	4.12	-	-	399.9	2	2.10	White solid
35	4.13	215-216	402	400	2	1.93	White solid
	4.14	120	419.8	-	2	2.15	Yellow solid
	4.15	80	391.9	-	2	2.77	White solid
	4.16	-	387.9	-	2	2.82	Green oil
40	4.17	-	387.9	-	2	2.77	Colourless oil
	4.18	-	401.9	-	2	2.97	Yellow oil
	4.19	-	427.9	-	2	3.03	Colourless oil
45	4.20	75	404.0	-	2	2.77	Yellow solid
	4.21	80	443.0	-	2	2.45	White solid
	4.22	-	402.0	-	2	2.79	Yellow oil
	4.23	-	444.0	-	2	2.91	Colourless oil
50	4.24	-	400.0	-	2	2.67	Colourless oil
	4.25	-	401.1	-	2	1.74	Colourless oil
	4.26	-	427.1	-	2	1.87	Colourless oil
55	4.27	135	415.0	-	2	2.18	White solid
	4.28	153-155	346.1	-	2	2.70	White solid
	4.29	-	441.1	-	2	2.08	Colourless oil

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(continued)

	Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
5	4.30	-	445.2	-	2	1.97	Colourless oil
	4.31	181	433.0	-	2	2.21	White solid
	4.32	-	369.0	-	2	2.64	Yellow oil
	4.33	-	391.9	-	2	2.41	White oil
10	4.34	192-194	377.9	-	2	3.00	White solid
	4.35	143-144	310.0	-	2	2.62	White solid
	4.36	-	409.0	407.0	2	2.62	Yellow solid
15	4.37	119	443.0	441.0	2	2.64	Brown solid
	4.38	131	451.0	449.0	2	2.53	Brown solid
	4.39	130-131	410	408	2	1.70	Yellow solid
	4.40	>290	443	441	2	2.64	Brown solid
20	4.41	174-175	-	365.1	2	2.15	White solid
	4.42	-	415.0	-	2	2.46	White solid
	4.43	158	410.0	-	2	1.59	White solid
25	4.44	195-196	385.1	383.1	2	2.21	White solid
	4.45	159-161	397.0	-	2	2.13	White solid
	4.46	120	418.0	-	2	1.61	White solid
	4.47	177	425.0	-	2	2.44	White solid
30	4.48	174-175	441.1	-	2	1.94	White solid
	4.49	148-150	436.1	-	2	1.61	White solid
	4.50	143-144	344.0	-	2	2.83	White solid
35	4.51	154-156	386.1	384.1	2	1.81	White solid
	4.52	118-120	420.0	418.0	2	2.10	White solid
	4.53	110-112	411.1	409.1	2	1.20	White solid
	4.54	142	409.0	-	2	2.13	White solid
40	4.55	209-212	403.1	-	2	2.25	White solid
	4.56	216-217	403.1	-	2	2.23	White solid
	4.57	164	398.1	-	2	1.86	White solid
45	4.58	168	-	366.1	2	1.13	White solid
	4.59	249	-	431.1	2	2.10	Orange Solid
	4.60	234	433.0	-	2	2.22	White solid
50	4.61	214	418.1	-	2	1.70	White solid
	4.62	170	415.0	-	2	2.16	White solid
	4.63	125	433.0	-	2	2.29	White solid
	4.64	168	433.0	-	2	2.23	White solid
55	4.65	128-130	420.1	418.1	2	2.36	White solid
	4.66	197	415.0	-	2	2.19	White solid
	4.67	110-112	346.1	-	2	2.71	White solid

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(continued)

	Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
5	4.68	172-174	346.0	-	2	2.67	White solid
	4.69	143-145	362.0	-	2	2.85	White solid
	4.80	-	397.0	-	2	2.14	Colourless oil
	4.90	-	368.0	-	2	2.69	White solid
10	4.91	144-145	393.0	-	2	2.83	White solid
	4.92	197-198	419.0	418.0	2	2.38	White solid
	4.93	70-72	394.0	-	2	2.93	White solid
15	4.94	119-120	346.0	-	2	2.64	White solid
	4.95	121-122	427.9	-	2	3.11	White solid
	4.96	148	433.0	-	2	2.24	White solid
	4.97	181	427.0	-	2	2.16	White solid
20	4.98	140-142	396.0	-	2	2.88	White solid
	4.99	65-67	414.0	-	2	2.83	White solid
	4.100	251-253	374.1	-	2	1.92	White solid
25	4.101	138	408.0	-	2	1.82	White solid
	4.102	209	408.0	-	2	1.81	White solid
	4.103	109	415.0	-	2	2.16	White solid
	4.104	65-66	335.1	-	2	2.45	White solid
30	4.105	164-166	335.1	-	2	2.43	White solid
	4.106	181	417	-	2	2.18	White solid
	4.107	166-168	340.1	-	2	2.03	White solid
35	4.108	78-80	402.0	-	2	2.67	White solid
	4.109	112-114	368.0	-	2	2.68	White solid
	4.110	158-160	314.1	-	2	1.00	White solid
	4.111	196	397.0	-	2	2.14	White solid
40	4.112	187	439.2	-	2	1.17	White solid
	4.113	131-133	350.0	-	2	2.82	White solid
	4.114	83-85	400.0	-	2	3.11	White solid
45	4.115	172-174	346	-	2	2.67	White solid
	4.116	204	410	-	2	1.63	White solid
	4.117	186-188	427	-	2	2.14	White solid
50	4.118	170-172	453	451	2	2.46	White solid
	4.119	105	436.1	-	2	2.06	Brown solid
	4.120	186	392	-	2	2.01	White solid
	4.121	120	422	-	2	1.88	Yellow solid
55	4.122	162-163	429.0	427.0	2	1.96	White solid
	4.124	137-138	-	420.2	2	2.00	Yellow foam
	4.125	137	415	-	2	2.11	White solid

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(continued)

	Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
5	4.126	175	427.0	-	2	2.22	White solid
	4.127	184	447.0	-	2	2.47	White solid
	4.128	175	402.0	-	2	1.93	White solid
	4.129	140	410.1	-	2	1.88	White solid
10	4.130	174-175	451	449	2	2.34	White solid
	4.131	165-166	487	485	2	-	White solid
	4.132	165-167	487	485	2	-	White solid
15	4.133	157	414.0	-	1	4.09	White solid
	4.134	163	426.0	-	2	2.22	Yellow solid
	4.135	138	427	425	2	2.13	White solid
	4.136	60	411.1	-	2	2.19	White solid
20	4.137	-	423.0	-	2	2.20	Colourless oil
	4.138	-	429	-	2	2.22	Colourless oil
	4.139	-	447	-	2	2.28	Colourless oil
25	4.140	141-142	324	-	2	2.70	White solid
	4.141	143	431.0	-	2	2.48	White solid
	4.142	146	443.0	-	2	2.43	White solid
	4.143	164-165	-	-	2	2.45	White solid
30	4.144	176-178	-	-	2	2.82	White solid
	4.145	91-93	427.0	-	2	2.64	White solid
	4.146	163-165	426.0	-	2	2.88	White solid
35	4.147	125-127	354	-	2	2.66	White solid
	4.148	123-125	342	-	2	2.71	White solid
	4.149	142-144	384.1	-	2	2.45	White solid
	4.150	138-140	354	-	2	2.67	White solid
40	4.151	172-174	411.1	-	2	2.18	White solid
	4.152	153-155	399.1	-	2	2.25	White solid
	4.153	136-138	411.1	-	2	2.20	White solid
45	4.154	104	385.0	-	2	2.14	White solid
	4.155	173	432.9	-	2	2.20	White solid
	4.156	156-158	451	-	2	2.25	Beige solid
	4.157	196-198	451	-	2	2.34	Beige solid
50	4.158	155-157	399	-	2	2.30	White solid
	4.159	171	437.9	-	2	2.14	White solid
	4.160	156-157	415.9	413.0	2	2.16	White solid
55	4.161	110	451	-	2	2.28	White solid
	4.162	210-212	440	-	2	2.05	Yellow foam
	4.163	207	445	-	2	2.24	White solid

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(continued)

	Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
5	4.164	176-177	446.1	-	2	2.13	White solid
	4.165	180-182	450.9	-	2	2.36	White solid
	4.166	-	338.1	-	2	2.83	Colourless oil
	4.167	-	383	-	2	2.63	White oil
10	4.168	215-217	410	-	2	2.10	White solid
	4.169	169-171	415	-	2	2.19	White solid
	4.170	134	428	-	2	2.15	White solid
15	4.171	141	411	-	2	1.83	White solid
	4.172	170-171	428.1	-	2	1.43	White solid
	4.173	160-161	416	-	2	1.90	White solid
	4.174	147-148	436	-	2	1.48	White solid
20	4.175	122	425	-	2	2.28	White solid
	4.176	169-171	417	-	2	2.32	White solid
	4.177	116-118	416.9	-	2	2.33	White solid
25	4.178	193-194	454	-	2	2.60	Pink solid
	4.179	198-199	469.9	-	2	2.12	White solid
	4.180	194-195	450.9	-	2	2.14	Yellow solid
	4.181	86-89	452.9	-	2	2.46	White solid
30	4.182	138	429	-	2	2.13	White solid
	4.183	209	425	-	2	1.88	White solid
	4.184	173-176	432.9	-	2	2.20	Beige solid
35	4.185	108-111	403	-	2	2.22	White solid
	4.186	200-202	430	-	2	2.19	White solid
	4.187	273-275	450.9	-	2	2.12	Beige solid
	4.188	158-159	432.9	-	2	2.28	Beige solid
40	4.189	161	428	426.9	2	1.75	Yellow solid
	4.190	191-193	420.9	-	2	1.90	White solid
	4.191	168-170	443	-	2	2.12	Yellow solid
45	4.192	188-190	420.9	-	2	1.90	Orange solid
	4.193	188-190	420.9	-	2	1.72	Yellow solid
	4.194	169	427	-	2	2.16	White solid
	4.195	211	427	-	2	2.15	White solid
50	4.196	94	398	396	2	1.53	Yellow solid
	4.197	230-231	422	420	2	2.08	White solid
	4.198	194-195	381	-	2	2.24	White solid
55	4.199	219	433.9	431.9	2	2.25	White solid
	4.200	207-208	468.9	-	2	2.17	White solid
	4.201	142-144	420.9	-	2	2.30	White solid

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(continued)

	Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
5	4.202	239-241	439	-	2	1.93	Pink solid
	4.203	143-146	454.8	452.9	2	2.52	White solid
	4.204	136-137	446	443.9	2	2.11	Beige solid
	4.205	132	419.9	416.9	2	2.36	White solid
10	4.206	184	455	-	2	2.41	White solid
	4.207	136-137	400.9	-	2	2.68	Beige solid
	4.208	120	418.9	-	2	2.36	White solid
15	4.209	160	408.9	-	2	1.83	White solid
	4.210	162-163	437	435	2	2.41	White solid
	4.211	190-191	453	452.9	2	2.56	White solid
	4.212	226-227	449.9	447.9	2	2.38	White solid
20	4.213	197	454.8	452.9	2	2.52	Yellow solid
	4.214	191	448.9	-	2	2.38	White solid
	4.215	195-196	448.9	-	2	2.33	White solid
25	4.216	163-164	435	-	2	2.31	White solid
	4.217	166	416.0	414.0	2	2.02	White solid
	4.218	232-233	464.9	-	2	2.53	White solid
	4.219	220-221	464.9	462.9	2	2.49	White solid
30	4.220	177	451	449	2	2.32	Brown solid
	4.221	197-198	454.9	452.9	2	2.55	White solid
	4.222	184-185	437.0	435.9	2	2.39	White solid
35	4.223	185-186	470.9	467.0	2	2.62	White solid
	4.224	147	463.9	462	2	2.44	White solid
	4.225	154	480	478	2	2.6	White solid
	4.226	171	462	460	2	2.33	Yellow solid
40	4.227	158-159	401.0	399.1	2	2.31	White solid
	4.228	108	446	-	2	2.20	White solid
	4.229	179	449	-	2	2.35	Beige solid
45	4.230	180-181	471.1	469.0	2	2.51	White solid
	4.231	201-202	464	462	2	2.22	White solid
	4.232	203	480	478	2	2.44	White solid
	4.233	161	487.9	485.0	2	2.62	Beige solid
50	4.234	160-161	444.1	442.0	2	1.56	White solid
	4.235	150-151	503.0	501.9	2	2.67	Beige solid
	4.236	179	431	-	2	2.30	Beige solid
55	4.237	236-237	443	441	2	2.30	White solid
	4.238	205-206	467	465	2	2.49	White solid
	4.239	129-130	362.0	-	2	2.83	White solid

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(continued)

	Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
5	4.240	219	503/505/506	-	2	2.73	White solid
	4.241	188	477	-	2	2.65	White solid
	4.242	218	456.9	455	2	2.06	White solid
	4.243	182	463	461	2	2.78	White solid
10	4.244	191	488	-	2	2.09	White solid
	4.245	192	435	-	2	2.01	White solid
	4.246	119	455	-	2	2.40	White solid
15	4.247	99	455	-	2	2.46	White solid
	4.248	96	445	-	2	2.68	White solid
	4.249	113	462	-	2	2.25	White solid
	4.250	234.5	440	-	2	2.23	White solid
20	4.251	161	-	-	2	3.01	White solid
	4.252	180	467	-	2	2.40	White solid
	4.253	114-115	410/412	-	2	2.84	White solid
25	4.254	205-206	494.1	-	2	2.40	White solid
	4.255	167-168	457.2	455.2	2	2.72	White solid
	4.256	87-88	400.2	-	2	3.16	White solid
	4.257	120	488.1	-	2	2.69	White solid
30	4.258	150-151	471	-	2	2.69	White solid
	4.259	137	479	-	2	2.55	Beige solid
	4.260	100-105	455.0	-	2	2.25	White solid
35	4.261	158	458	-	2	2.16	Beige solid
	4.262	65	-	-	2	2.69	White solid
	4.263	154	504	-	2	2.33	White solid
	4.264	184-185	496.1	-	2	2.60	White solid
40	4.265	-	420	-	2	3.09	Beige solid
	4.266	69	485/486	-	2	2.59	White solid
	4.267	155	463	-	2	2.39	White solid
45	4.268	148	488	-	2	2.14	Beige solid
	4.269	179	462.1	460.0	2	2.38	White solid
	5.01	178	413.0	-	2	2.82	White solid
50	5.02	184	399.0	-	2	2.63	White solid
	5.03	164	427.0	-	2	2.96	White solid
	5.04	76-78	-	-	2	2.71	White solid
	5.05	103	457.9	-	2	1.75	White solid
55	5.06	145	453.0	-	2	2.17	White solid
	5.07	148	444.8	-	2	2.36	White solid
	5.08	186-187	399.9	-	1	3.99	Yellow solid

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(continued)

Compound Nr	Melting point (°C)	[MH ⁺]	[MH ⁻]	LC Method	Rt (min)	Appearance
5.09	105	414.9	-	1	4.18	White solid
5.10	197-198	430.9	428.9	1	4.22	Beige solid
5.11	190-191	428.9	-	1	4.34	Beige solid
5.12	202	450.7	-	1	4.3	Yellow solid
5.13	189-190	414.0	-	1	4.39	Yellow solid
5.14	-	400.8	-	2	2.07	White solid
5.15	143	427.9	-	2	1.74	Beige solid
5.16	97	414.8	-	2	2.36	Yellow solid
5.17	193	428	-	1	3.95	Beige solid
5.18	183	457	-	2	2.65	Beige solid
5.19	184-186	447.4	-	2	2.25	Pink solid

Table 7: NMR-data

Compound Nr	NMR-data
1.01	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.21 (1H, bs), 7.89-7.88 (1H, m), 7.60-7.57 (1H, m), 7.47-7.39 (5H, m), 7.16-7.13 (1H, m), 4.99 (2H, m), 2.05 (3H, s)
1.02	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.97 (1H, bs), 8.22-8.19 (1H, m), 7.60-7.57 (1H, m), 7.46-7.39 (5H, m), 7.27-7.21 (1H, m), 4.98 (2H, m), 2.11 (3H, s)
1.03	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.76-7.75 (1H, m), 7.50-7.47 (1H, m), 7.35-7.18 (11H, m), 7.05-7.02 (1H, m), 6.99 (1H, s), 1.96 (3H, s)
1.04	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.97 (1H, s), 8.22-8.19 (3H, m), 7.72-7.69 (2H, m), 7.41-7.39 (1H, m), 7.27-7.22 (1H, m), 5.13 (2H, s), 2.11 (3H, s)
1.05	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.97 (1H, s), 8.23-8.20 (1H, dd), 7.69-7.68 (1H, d), 7.63-7.61 (1H, d), 7.46-7.39 (2H, m), 7.27-7.21 (1H, m), 5.00 (2H, s), 2.12 (3H, s)
1.06	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.97 (1H, s), 8.23-8.18 (1H, m), 7.54-7.39 (3H, m), 7.32-7.21 (2H, m), 5.01 (2H, s), 2.11 (3H, s)
1.07	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.56 (2H, m), 7.41 (2H, m), 7.3 (1H, s), 7.17 (1H, dd), 6.72 (1H, d), 4.92 (2H, s), 4.51 (2H, t), 3.15 (2H, t)
2.01	¹ H-NMR (CDCl ₃), δ (ppm): 7.82 (1H, s), 7.54 (2H, s), 7.46-7.44 (2H, m), 7.37-7.35 (2H, m), 7.32-7.29 (2H, m), 7.21-7.18 (1H, m), 5.08 (2H, s), 2.13 (3H, s)
2.02	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.00 (1H, sl), 7.82 (1H, s), 7.40 (6H, m), 7.23 (2H, m), 4.98 (2H, s), 3.02 (3H, s)
2.03	¹ H-NMR (CDCl ₃), δ (ppm): 7.46 (1H, m), 7.42 (1H, m), 7.39 (1H, m), 7.36-7.33 (1H, m), 7.24-7.19 (3H, m), 7.01-6.94 (2H, m), 5.04 (1H, s), 3.75 (3H, s)
2.04	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 8.21-8.19 (2H, d, 8.7Hz), 7.86-7.85 (2H, m), 7.68-7.65 (2H, d, 8.7Hz), 7.55-7.52 (1H, d, 8.2Hz), 7.42-7.36 (1H, t, 8.1Hz), 7.19 (1H, d, 8.2Hz), 5.12 (2H, s), 2.05 (3H, s)
2.05	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.85-7.80 (4H, m), 7.60-7.55 (3H, m), 7.41-7.36 (1H, t, 7.94Hz), 7.18-7.16 (1H, dm, 7.94Hz), 5.07 (2H, s), 2.05 (3H, s)
2.06	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, bs), 7.91-7.83 (6H, m), 7.56-7.49 (4H, m), 7.41-7.35 (1H, t), 7.20-7.16 (1H, m), 5.16 (2H, s), 2.03 (3H, s)

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Compound Nr		NMR-data
5	2.07	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 8.07-8.04 (2H, m), 7.89-7.85 (2H, m), 7.66-7.62 (1H, m), 7.55-7.52 (1H, m), 7.42-7.37 (1H, t), 7.21-7.18 (1H, m), 5.11 (2H, s), 2.05 (3H, s)
	2.08	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.05 (1H, s), 7.75 (1H, s), 7.71 (1H, s), 7.47 (2H, d, J=7.13Hz), 7.32 (4H, m), 7.07 (1H, t, J=7.13Hz), 4.98 (2H, s), 1.98 (3H, s)
10	2.09	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.84-7.81 (2H, m), 7.57-7.54 (1H, dm, 8.19Hz), 7.41-7.27 (6H, m), 7.19-7.15 (1H, dm, 8.19Hz), 5 (2H, s), 2.05 (3H, s)
	2.10	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.84-7.83 (1H, m), 7.81 (1H, s), 7.56-7.51 (3H, m), 7.41-7.33 (3H, m), 7.18-7.15 (1H, dm), 4.96 (2H, s), 2.05 (3H, s)
15	2.11	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, s), 7.94-7.82 (4H, m), 7.57-7.47 (3H, m), 7.42-7.35 (1H, m), 7.18-7.15 (1H, m), 5.06 (2H, s), 3.83 (3H, s), 2.04 (3H, s)
	2.12	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.15 (1H, s), 8.22-8.19 (1H, m), 8.01-7.96 (1H, m), 7.91-7.83 (3H, m), 7.65-7.52 (3H, m), 7.47-7.36 (2H, m), 7.22-7.18 (1H, m), 5.49 (2H, s), 2.05 (3H, s)
20	2.13	¹ H-NMR (CDCl ₃), δ (ppm): 7.77 (1H, bs), 7.55-7.34 (6H, m), 7.24-7.18 (1H, m), 6.86-6.81 (2H, m), 5.08 (2H, s), 3.78 (3H, s), 2.15 (3H, s)
	2.14	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.84-7.83 (1H, m), 7.81 (1H, s), 7.56-7.51 (3H, m), 7.41-7.33 (3H, m), 7.18-7.15 (1H, dm), 4.96 (2H, s), 2.05 (3H, s)
25	2.15	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.85-7.81 (2H, m), 7.56-7.53 (1H, d, 8.7Hz), 7.46 (1H, s), 7.42-7.35 (4H, m), 7.18-7.16 (1H, d, 8.7Hz), 4.99 (2H, s), 2.05 (3H, s)
	2.16	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.82-7.77 (2H, m), 7.56-7.51 (1H, m), 7.41-7.36 (1H, m), 7.27-7.22 (2H, m), 7.17-7.10 (3H, m), 4.95 (2H, s), 2.27 (3H, s), 2.06 (3H, s)
30	2.17	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.83-7.79 (2H, m), 7.57-7.54 (1H, m), 7.42-7.36 (1H, m), 7.31-7.29 (2H, m), 7.23-7.14 (3H, m), 4.96 (2H, d), 2.97-2.94 (1H, m), 2.05 (3H, s), 1.19-1.16 (6H, m)
	2.18	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.83-7.80 (2H, m), 7.57-7.54 (1H, d, 8.7Hz), 7.42-7.36 (1H, t, 8.7Hz), 7.18-7.14 (2H, m), 7.08 (2H, s), 4.92 (2H, s), 2.19-2.18 (6H, m), 2.05 (3H, s)
35	2.19	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 8.46-8.45 (1H, s), 7.89-7.81 (3H, m), 7.55-7.48 (2H, m), 7.41-7.36 (1H, t, 8.5Hz), 7.18-7.15 (1H, d, 8.5Hz), 5.01 (2H, s), 2.05 (3H, s)
	2.20	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, s), 7.85-7.81 (2H, m), 7.57-7.13 (6H, m), 5 (2H, s), 2.06-2.03 (3H, m)
40	3.01	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, bs), 7.78 (1H, t, 2Hz), 7.38 (1H, m), 7.25 (2H, d, 8Hz), 7.14 (3H, m), 4.97 (2H, s), 2.68 (3H, s), 2.19 (3H, s), 2.04 (3H, s)
	3.02	¹ H-NMR (DMSO-d ₆), δ (ppm): 0.14 (1H, bs), 7.8 (1H, t), 7.58-7.56 (1H, d), 7.41-7.28 (7H, m), 7.18-7.16 (1H, d), 5.03 (2H, s), 2.2 (3H, s), 2.05 (3H, s)
45	3.03	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, s), 7.81 (1H, s), 7.60-7.10 (6H, m), 5.01 (2H, s), 2.20 (3H, s), 2.05 (3H, s)
	3.04	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, s), 8.27 (1H, s), 7.99 (1H, t), 7.81 (1H, s), 7.55 (1H, d), 7.37 (1H, t), 7.16 (2H, m), 5.03 (2H, s), 2.19 (3H, s), 2.04 (3H, s)
50	3.05	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 8.5-7.1 (7H, m), 5.0 (2H, s), 3.0-2.9 (1H, m), 3.2 (3H, s) 2.2 (3H, s), 2.1 (3H, s), 1.2 (6H, s)
	3.06	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 7.8-7.0 (8H, m), 5.0 (2H, s), 4.3 (2H, t), 3.7 (2H, t), 3.4 (3H, s), 2.2 (3H, s), 2.0 (3H, s)
55	3.07	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, bs), 8.10 (1H, m), 7.25 (6H, m), 4.98 (2H, s), 2.18 (3H, s), 2.10 (3H, s)
	3.08	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 7.8-6.9 (8H, m), 5.0 (2H, s), 3.7 (3H, s), 2.2 (3H, s), 2.1 (3H, s)

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	Compound Nr	NMR-data
5	3.09	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 7.8-7.1 (8H, m), 5.0 (2H, s), 3.7 (3H, s), 2.2 (3H, s), 2.1 (3H, s), 1.2 (6H, d)
	3.10	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.56-7.28 (9H, m), 5.01 (2H, s), 2.19 (3H, s)
	3.11	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 8.0-7.2 (11H, m), 5.0 (2H, s), 2.2 (3H, s), 2.1 (3H, s)
10	3.12	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, s), 8.50 (1H, s), 7.79 (1H, s), 7.67 (1H, d), 7.56 (1H, d), 7.37 (1H, t), 7.23 (1H, d), 7.15 (1H, s), 4.99 (2H, s), 3.12 (1H, q), 2.19 (3H, s), 2.04 (3H, s), 2.00-1.86 (2H, m), 1.82-1.52 (6H, m)
15	3.13	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, s), 7.81 (1H, s), 7.59-7.51 (1H, d), 7.50-7.30 (3H, m), 7.29-7.20 (1H, m), 7.19-7.12 (1H, d), 4.99 (2H, s), 2.20 (3H, s), 2.04 (3H, s)
	3.14	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 7.8 (1H, s), 7.6-7.5 (1H, d), 7.4 (1H, t), 7.2 (1H, s), 7.0 (1H, s), 6.9 (2H, s), 5.0 (2H, s), 3.7 (6H, s), 2.2 (3H, s), 2.1 (3H, s)
20	3.15	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 7.8 (1H, s), 7.6-7.5 (1H, d), 7.5-7.4 (2H, m), 7.2-7.1 (1H, t) 7.1 (1H, d), 7.0 (1H, t), 5.0 (2H, s), 3.7 (6H, s), 2.2 (3H, s), 2.1 (3H, s)
	3.16	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, s), 7.78 (1H, s), 7.56 (1H, d), 7.37 (1H, t), 7.29 (2H, d), 7.15 (1H, d), 6.85 (2H, d), 4.93 (2H, s), 4.57 (1H, q), 2.18 (3H, s), 2.04 (3H, s), 1.23 (6H, d)
25	3.17	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, s), 8.22 (1H, d), 7.98 (1H, d), 7.91-7.80 (2H, m), 7.67-7.30 (6H, m), 7.20 (1H, d), 5.51 (2H, s), 2.24 (3H, s), 2.05 (3H, s)
	3.18	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.80 (1H, t, 2Hz), 7.55 (1H, d, 8Hz), 7.37 (5H, m), 7.15 (1H, dd, 1Hz, 8Hz), 5.00 (2H, s), 2.19 (3H, s), 2.05 (3H, s)
30	3.19	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, bs), 8.10-8.06 (1H, m), 7.40-7.35 (1H, m), 7.27-7.24 (3H, m), 7.14-7.11 (2H, m), 4.96 (2H, s), 2.27 (3H, s), 2.18 (3H, s), 2.10 (3H, s)
	3.20	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, s), 8.11 (1H, s), 7.42-7.18 (3H, m), 6.98-6.89 (2H, m), 6.85 (1H, d), 4.98 (2H, s), 3.73 (3H, s), 2.19 (3H, s), 2.10 (3H, s)
35	4.01	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.87 (1H, s), 7.56-7.53 (1H, m), 7.43-7.34 (5H, m), 7.25-7.21 (1H, m), 5 (2H, s), 3.82 (3H, s), 2.05 (3H, s)
	4.02	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.18 (1H, bs), 8.20-8.18 (2H, d), 7.92 (1H, s), 7.68-7.64 (2H, d), 7.58-7.55 (1H, d), 7.39-7.33 (1H, t), 7.27-7.23 (1H, d), 5.12 (2H, s), 2.05 (3H, s)
40	4.03	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.86 (1H, s), 7.55 (1H, d), 7.37 (1H, t), 7.29-7.19 (3H, m), 7.12 (2H, d), 4.97 (2H, s), 3.81 (3H, s), 2.26 (3H, s), 2.05 (3H, s)
	4.04	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, s), 7.9 (1H, s), 7.73-7.53 (5H, m), 7.41-7.35 (1H, m), 7.28-7.23 (1H, m), 5.11 (2H, s), 3.84 (3H, s), 2.06 (3H, s)
45	4.05	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.89 (1H, s), 7.58-7.22 (6H, m), 5.01 (2H, s), 3.83 (3H, s), 2.05 (3H, s)
	4.06	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.61-7.55 (1H, m), 7.48-7.36 (3H, m), 7.22-7.17 (2H, m), 7.09-7.03 (1H, m), 5.02 (2H, s), 3.78 (3H, s), 3.74 (3H, s)
50	4.07	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.96 (1H, s), 7.51-7.31 (5H, m), 7.19-7.15 (1H, s), 5 (2H, s), 3.83 (3H, s), 3.02 (3H, s)
	4.08	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.49-7.32 (6H, m), 7.19-7.16 (1H, m), 7.08-7.02 (1H, m), 5.01 (2H, s), 3.77 (3H, s), 3.73 (3H, s)
55	4.09	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.60-7.67 (1H, m), 7.42-7.24 (7H, m), 7.09-7.04 (1H, m), 7.29-7.22 (2H, s), 3.81 (3H, s), 3.14-3.04 (2H, m), 1.04 (3H, t)
	4.10	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 8.48-8.44 (1H, m), 7.88-7.84 (1H, m), 7.68-7.63 (1H, m), 7.57-7.52 (1H, m), 7.37 (1H, t), 7.25-7.18 (2H, m), 4.99 (2H, s), 3.81 (3H, s), 2.43 (3H, s), 2.05(3H,s)

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Compound Nr	NMR-data
5	4.11 ¹ H-NMR (MeOD-d ₄), δ (ppm): 8.18-8.15 (1H, m), 7.35-7.31 (2H, m), 7.26-7.24 (2H, d, 7.9Hz), 7.14-7.11 (2H, d, 7.9Hz), 4.96 (2H, s), 3.8 (3H, s), 2.27 (3H, s), 2.1 (3H, s)
	4.12 ¹ H-NMR (MeOD-d ₆), δ (ppm): 8.6 (2H, bd), 8.4 (1H, s), 7.3 (2H, d), 7.1 (1H, d), 5.0 (2H, s), 3.9 (3H, s), 2.1 (3H, s)
10	4.13 ¹ H-NMR (DMSO-d ₆), δ (ppm): 8.43 (1H, s), 7.90-7.83 (2H, m), 7.58-7.45 (2H, m), 7.40-7.32 (1H, m), 7.23-7.21 (1H, m), 5.02 (2H, s), 3.81 (3H, s), 2.05 (3H, s)
	4.14 ¹ H-NMR (DMSO-d ₆), δ (ppm): 8.67 (1H, s), 8.51 (1H, s), 8.38 (1H, s), 7.57-7.50 (1H, m), 7.49-7.43 (1H, m), 7.29-7.24 (1H, m), 5.02 (2H, s), 3.86 (3H, s), 2.09 (3H, s)
15	4.15 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.49-7.33 (5H, m), 7.12 (1H, dd), 6.9 (1H, td), 5 (2H, s), 3.79 (3H, s), 3.73 (3H, s)
	4.16 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.54-7.28 (8H, m), 5.01 (2H, s), 4.46 (2H, s), 3.82 (3H, s), 3.31 (3H, s)
	4.17 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.53-7.38 (8H, m), 5.01 (2H, s), 4.38 (2H, s), 3.76 (3H, s), 3.18 (3H, s)
20	4.18 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.45-7.36 (7H, m), 7.26-7.22 (1H, m), 5 (2H, s), 3.82 (3H, s), 3.55 (2H, t), 3.24 (3H, s), 2.85 (2H, t)
	4.19 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.70-7.52 (4H, m), 7.43-7.34 (4H, m), 5.02 (2H, s), 3.76 (3H, s)
25	4.20 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.44-7.35 (4H, m), 7.19-7.15 (2H, m), 7.03-6.97 (1H, m), 5.02 (2H, s), 3.86 (3H, s), 3.75 (3H, s), 3.7 (3H, s)
	4.21 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.45-7.38 (8H, m), 5.06-4.97 (2H, m), 3.75 (3H, s), 3.46-3.36 (2H, m), 3.33-3.22 (4H, m), 2.13-1.99 (4H, m)
30	4.22 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.41-7.33 (8H, m), 5 (2H, s), 3.75 (3H, s), 3.44 (2H, t), 3.14 (3H, s), 2.75 (2H, t)
	4.23 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.46-7.39 (7H, m), 7.25-7.21 (1H, m), 5.01 (2H, s), 3.85-3.75 (7H, m), 2.91 (2H, s), 1.22 (3H, s)
35	4.24 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.44-7.36 (7H, m), 7.18-7.15 (1H, m), 5 (2H, s), 3.85 (2H, s), 3.83 (3H, s), 2.17 (3H, s)
	4.25 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.51-7.33 (8H, m), 5.01 (2H, s), 3.76 (3H, s), 3.33 (2H, s), 1.92 (6H, s)
	4.26 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.47-7.31 (8H, m), 5 (2H, s), 3.75 (3H, s), 3.5 (2H, bs), 2.18 (4H, bs), 1.47 (4H, bs)
40	4.27 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, bs), 8.15 (1H, d), 7.35 (4H, m), 6.87 (2H, d), 4.95 (2H, s), 3.80 (3H, s), 3.73 (3H, s), 2.11 (3H, s)
	4.28 ¹ H-NMR (CDCl ₃), δ (ppm): 7.6 (4H, m), 7.2 (2H, t), 7.0 (2H, t), 5.1 (2H, s), 3.9 (3H, s)
45	4.29 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.46-7.36 (8H, m), 5.08-4.94 (2H, m), 3.75 (3H, s), 3.33 (2H, s), 2.06-1.93 (4H, m), 1.23-1.06 (6H, m)
	4.30 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.48-7.36 (8H, m), 5 (2H, s), 3.75 (3H, s), 3.14 (2H, t), 3.08 (3H, s), 2.3 (2H, s), 1.93 (3H, s)
	4.31 ¹ H-NMR (DMSO-d ₆), δ (ppm):
50	4.32 ¹ H-NMR (DMSO-d ₆), δ (ppm): 8.01 (1H, d), 7.88 (1H, t), 7.75 (1H, d), 7.65 (1H, t), 7.44-7.36 (4H, m), 5.03 (2H, s), 3.83 (3H, s)
	4.33 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.46-7.31 (6H, m), 7.24-7.16 (1H, m), 4.99 (2H, s), 4.47-4.42 (2H, m), 3.75 (3H, s)
55	4.34 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.7-7.3 (8H, m), 5.0 (2H, s), 3.9 (3H, s)
	4.35 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.58 (2H, d), 7.47 (2H, t), 7.42-7.20 (6H, m), 5.03 (2H, s), 3.82 (3H, s)

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Compound Nr	NMR-data
5	4.36 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.87 (1H, s), 7.57-7.54 (1H, m), 7.40-7.34 (1H, t), 7.30-7.18 (5H, m), 4.98 (2H, s), 3.81 (3H, s), 2.9-2.81 (1H, q), 2.05 (3H, s), 1.18-1.16 (6H, m)
	4.37 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 7.89-7.88 (1H, m), 7.66-7.62 (3H, m), 7.58-7.54 (2H, m), 7.50-7.34 (6H, m), 7.25-7.23 (1H, m), 5.09 (2H, s), 3.82 (3H, s), 2.05 (3H, s)
10	4.38 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 7.89-7.88 (1H, m), 7.56-7.50 (3H, m), 7.40-7.34 (3H, m), 7.25-7.23 (1H, m), 5.04 (2H, s), 3.82 (3H, s), 2.05 (3H, s)
	4.39 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 7.85-7.84 (1H, s), 7.56-7.54 (1H, m), 7.39-7.33 (1H, m), 7.23-7.19 (3H, m), 6.66-6.64 (2H, m), 4.9 (2H, s), 3.8 (3H, s), 2.85 (6H, s), 2.05 (3H, s)
15	4.40 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, bs), 7.89-7.87 (1H, m), 7.65-7.60 (4H, m), 7.57-7.54 (1H, m), 7.48-7.43 (4H, m), 7.41-7.33 (2H, m), 7.27-7.24 (1H, m), 5.07 (2H, s), 3.83 (3H, s), 2.05 (3H, s)
	4.41 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.86 (1H, s), 7.55 (1H, d), 7.45-7.20 (7H, m), 5.02 (2H, s), 3.81 (3H, s), 2.05 (3H, s)
20	4.42 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 8.0-7.3 (11H, m), 5.2 (2H, s), 3.8 (3H, s), 2.0 (3H, s)
	4.43 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 8.51 (1H, d, 2Hz), 7.88 (1H, t, 1Hz), 7.7 (1H, dd, 2Hz, 8Hz), 7.54 (1H, d, 8Hz), 7.37 (1H, t, 8Hz), 7.23 (1H, d, 8Hz), 5 (2H, s), 3.81 (3H, s), 2.99 (1H, q, 7Hz), 2.05 (3H, s), 1.21 (3H, s), 1.19 (3H, s)
25	4.44 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.87 (1H, s), 7.55 (1H, d), 7.47-7.30 (3H, m), 7.23 (1H, d), 7.15 (2H, t), 5.00 (2H, s), 3.81 (3H, s), 2.05 (3H, s)
	4.45 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 7.9-6.9 (8H, m), 5.0 (2H, s), 3.8 (3H, s), 3.7 (3H, s), 2.1 (3H, s)
30	4.46 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 8.9-7.3 (10H, m), 5.2 (2H, s), 3.8 (3H, s), 2.0 (3H, s)
	4.47 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, bs), 7.87 (1H, s), 7.55 (1H, d), 7.37 (1H, t), 7.27 (3H, m), 6.84 (1H, d), 4.94 (2H, s), 4.57 (1H, h), 3.81 (3H, s), 2.06 (3H, s), 1.23 (6H, d)
	4.48 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 7.9 (1H, s), 7.6-7.5 (1H, d), 7.4 (1H, t), 7.3-7.2 (1H, s), 7.0 (1H, s), 6.9 (2H, s), 5.0 (2H, s), 3.8 (3H, s) 3.7 (6H, s), 2.1 (3H, s)
35	4.49 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s) 8.5-7.2 (11H, m), 5.0 (2H, s), 3.8 (3H, s), 3.2-3.1 (1H, m), 2.1 (3H, s), 2.0-1.9 (2H, m), 1.7-1.6 (6H, m)
	4.50 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.58 (2H, d), 7.53-7.32 (7H, m), 5.01 (2H, s), 3.82 (3H, s)
40	4.51 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, s), 8.28 (1H, s), 8.00 (1H, t), 7.89 (1H, s), 7.55 (1H, d), 7.38 (1H, t), 7.24 (1H, d), 7.17 (1H, d), 5.05 (2H, s), 3.82 (3H, s), 2.06 (3H, s)
	4.52 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 8.35 (1H, s), 7.96 (1H, d), 7.90 (1H, s), 7.53 (1H, d), 7.37 (1H, t), 7.23 (1H, d), 5.07 (2H, s), 3.83 (3H, s), 2.05 (3H, s)
45	4.53 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 8.11 (1H, s), 7.85 (1H, s), 7.60-7.46 (2H, m), 7.36 (1H, t), 7.22 (1H, d), 6.57 (1H, d), 4.87 (2H, s), 3.80 (3H, s), 2.98 (6H, s), 2.05 (3H, s)
	4.54 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.87 (1H, t), 7.54 (1H, d), 7.37 (1H, t), 7.23 (2H, m), 7.12 (1H, d), 6.69 (1H, d), 4.93 (2H, s), 4.49 (2H, t), 3.82 (3H, s), 3.17 (2H, t), 2.06 (3H, s)
50	4.55 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.4 (1H, s), 8.1 (1H, s), 7.8-7.5 (6H, m), 5.0 (2H, s), 4.1 (3H, s), 2.3 (3H, s)
	4.56 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 7.9 (1H, s), 7.6-7.5 (1H, d), 7.5 (1H, d), 7.5-7.4 (1H, d), 7.4 (1H, t), 7.3-7.2 (1H, d), 7.0 (1H, t), 5.0 (2H, s), 3.8 (3H, s), 2.1 (3H, s)
55	4.57 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 8.2 (1H, s), 7.9 (1H, s), 7.7 (1H, dd), 7.6-7.5 (1H, s), 7.4 (1H, t), 7.2 (1H, s), 6.8 (1H, s), 5.0 (2H, s), 3.8 (3H, s) 3.8 (3H, s), 2.1 (3H, s)
	4.58 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 8.6 (1H, d), 8.5 (1H, dd), 7.9 (1H, t), 7.8 (1H, dt), 7.6-7.5 (1H, d), 7.4 (2H, m), 7.3-7.2 (1H, d), 5.0 (2H, s), 3.8 (3H, s), 2.1 (3H, s)

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Compound Nr		NMR-data
5	4.59	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 8.5 (1H, d), 7.9 (1H, t), 7.8-7.7 (3H, m), 7.6-7.5 (3H, m), 7.4 (1H, t), 7.3-7.2 (1H, d), 5.0 (2H, s), 3.8 (3H, s), 2.0 (3H, s)
	4.60	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.83 (1H, dd), 7.55 (1H, d), 7.37 (1H, t), 7.19 (1H, dd), 6.69 (2H, d), 5.02 (2H, s), 3.81 (3H, s), 3.76 (3H, s), 2.05 (3H, s)
10	4.61	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 9.0-8.9 (1H, d), 8.3 (1H, d), 8.0 (2H, m), 7.9 (1H, t), 7.8-7.7 (1H, td), 7.6-7.5 (2H, m), 7.4-7.3 (1H, t), 7.3-7.2 (1H, d), 5.2 (2H, s), 3.8 (3H, s), 2.0 (3H, s)
	4.62	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, bs), 7.88 (1H, s), 7.56 (1H, d), 7.38 (1H, t), 7.16 (4H, m), 4.95 (2H, s), 3.82 (6H, s), 2.06 (3H, s)
15	4.63	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.89 (1H, t), 7.53 (1H, d), 7.37 (1H, t), 7.22 (3H, m), 4.95 (2H, s), 3.88 (3H, s), 3.82 (3H, s), 2.05 (3H, s)
	4.64	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.88 (1H, t), 7.54 (1H, dd), 7.37 (1H, t), 7.2 (2H, m), 6.96 (1H, dt), 5.02 (2H, s), 3.85 (3H, s), 3.83 (3H, s), 2.05 (3H, s)
20	4.65	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.85 (1H, s), 7.56 (2H, s), 7.50-7.13 (5H, m), 6.38 (1H, s), 5.10 (2H, s), 3.81 (3H, s), 3.76 (3H, s), 2.05 (3H, s)
	4.66	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.87 (1H, t), 7.55 (1H, d), 7.33 (3H, m), 6.83 (1H, dd), 6.73 (1H, dd), 4.99 (2H, s), 3.83 (3H, s), 3.74 (3H, s), 2.05 (3H, s)
25	4.67	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.2-7.6 (8H, m), 5 (2H, s), 3.8 (3H, s)
	4.68	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.6-7.1 (8H, m), 5.0 (2H, s), 3.9 (3H, s)
	4.69	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.6-7.2 (8H, m), 5 (2H, s), 3.8 (3H, s)
30	4.80	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.87 (1H, s), 7.55 (1H, d), 7.39 (1H, t), 7.23 (2H, m), 6.86 (3H, m), 4.99 (2H, s), 3.82 (3H, s), 3.73 (3H, s), 2.05 (3H, s)
	4.90	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.5-7.3 (5H, m), 6.8 (4H, m), 4.5 (2H, m), 4.3 (1H, m), 4.1 (2H, m), 3.9 (3H, s)
	4.91	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.9 (2H, m), 7.5 (2H, m), 7.4-7.2 (7H, m), 5.3 (2H, s), 3.9 (3H, s)
35	4.92	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.88 (1H, s), 7.56-7.54 (1H, m), 7.47-7.35 (3H, m), 7.25-7.23 (2H, m), 5.03 (2H, s), 3.84 (3H, s), 2.05 (3H, s)
	4.93	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.6-7.3 (8H, m), 5.1 (2H, s), 3.8 (3H, s)
	4.94	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.60-7.57 (2H, d, 7.42Hz), 7.50-7.45 (2H, t), 7.38-7.12 (4H, m), 5.04 (2H, s), 3.84 (3H, s)
40	4.95	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.66-7.33 (8H, m), 5.05 (2H, s), 3.83 (3H, s)
	4.96	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, bs), 7.9 (1H, s), 7.55 (1H, d), 7.41 (1H, t), 7.2 (1H, d), 7.1 (2H, m), 5 (2H, s), 3.9 (3H, s), 3.8 (3H, s), 2 (3H, s)
	4.97	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, bs), 7.9 (1H, s), 7.6 (1H, d), 7.4 (1H, t), 7.2 (2H, t), 6.6 (1H, d), 5 (2H, s), 4.6 (2H, t), 3.8 (3H, s), 3.1 (2H, t), 2.1 (3H, t)
45	4.98	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.8-7.3 (8H, m), 5.1 (2H, s), 3.8 (3H, s)
	4.99	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.74-7.65 (5H, m), 7.61-7.45 (8H, m), 7.39-7.36 (1H, m)
	4.100	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.60-7.57 (2H, d, 7.42Hz), 7.50-7.45 (2H, t), 7.38-7.12 (4H, m), 5.04 (2H, s), 3.84 (3H, s)
50	4.101	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 8.36 (1H, d), 8.12 (2H, dd), 7.87 (1H, s), 7.54 (1H, d), 7.36 (1H, t), 7.23 (1H, d), 7.01 (1H, d), 5.14 (2H, s), 3.81 (3H, s), 2.04 (3H, s)
	4.102	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 8.74 (1H, s), 7.89 (1H, s), 7.78 (2H, m), 7.55 (1H, d), 7.43 (2H, m), 7.25 (1H, m), 5.17 (2H, s), 3.84 (3H, s), 2.06 (3H, s)

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Compound Nr		NMR-data
5	4.103	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.89 (1H, d), 7.53 (1H, d), 7.37 (1H, t), 7.13 (3H, m), 6.9 (1H, m), 4.98 (2H, s), 3.82 (6H, s), 2.05 (3H, s)
	4.104	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.86 (1H, bs), 7.76-7.72 (2H, t), 7.60-7.45 (6H, m), 7.38-7.36 (1H, m), 5.07 (2H, s), 3.84 (3H, s)
10	4.105	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.81-7.79 (2H, m), 7.59-7.56 (4H, m), 7.50-7.44 (2H, m), 7.38-7.33 (1H, m), 5.09 (2H, s), 3.83 (3H, s)
	4.106	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.88 (1H, s), 7.54 (1H, d), 7.37 (1H, t), 7.23 (1H, dd), 7.05 (2H, m), 4.92 (2H, s), 4.6 (2H, t), 3.81 (3H, s), 3.21 (2H, t), 2.05 (3H, s)
15	4.107	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.59-7.44 (4H, m), 7.38-7.24 (5H, m), 5.17-5.13 (2H, t, 5.63Hz), 5.01 (2H, s), 4.46-4.44 (2H, d, 5.63Hz), 3.82 (3H, s)
	4.108	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.61-7.59 (2H, d), 7.51-7.45 (2H, t), 7.38-7.33 (1H, t), 7.2 (1H, d), 7.09 (1H, d), 5.34 (2H, s), 4.92-4.88 (4H, d), 3.85 (3H, s)
20	4.109	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.61-7.59 (2H, d), 7.51-7.45 (2H, t), 7.38-7.33 (1H, t), 7.2 (1H, d), 7.09 (1H, d), 5.34 (2H, s), 4.92-4.88 (4H, d), 3.85 (3H, s)
	4.110	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.58-7.56 (2H, m), 7.50-7.45 (2H, m), 7.39-7.34 (1H, m), 7.05 (1H, d), 6.72 (1H, d), 5.07 (2H, bs), 3.85 (3H, s)
25	4.111	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.87 (1H, s), 7.55 (1H, d), 7.39 (1H, t), 7.23 (2H, m), 6.86 (3H, m), 4.99 (2H, s), 3.82 (3H, s), 3.73 (3H, s), 2.05 (3H, s)
	4.112	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 8.1 (1H, d), 7.9 (1H, t), 7.8 (1H, dt), 7.6-7.5 (1H, d), 7.5 (1H, dd), 7.4-7.3 (1H, t), 7.2 (1H, d), 6.5 (1H, d), 4.9 (2H, s), 3.8 (3H, s), 3.5-3.4 (4H, q), 2.1 (3H, s), 1.1-1.0 (6H, t)
30	4.113	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.58-7.37 (5H, m), 7.01-6.98 (2H, d), 5.09 (2H, s), 3.81 (3H, s)
	4.114	¹ H-NMR (DMSO-d ₆), δ (ppm): 8.17 (1H, d), 8.05-8.02 (1H, d, 8.44Hz), 7.87 (1H, s), 7.60-7.57 (2H, d, 7.68Hz), 7.51-7.34 (4H, m), 5.23 (2H, s), 3.83 (3H, s)
	4.115	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.0-7.6 (8H, m), 5 (2H, s), 3.8 (3H, s)
35	4.116	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.91 (2H, dt), 7.61 (1H, d), 7.54 (1H, d), 7.37 (1H, t), 7.22 (1H, dt), 4.93 (2H, s), 4.54 (2H, t), 3.81 (3H, s), 3.19 (2H, t), 2.05 (3H, s)
	4.117	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.86 (1H, t), 7.54 (1H, d), 7.37 (1H, t), 7.22 (1H, dd), 7.12 (1H, dd), 6.56 (1H, d), 4.98 (2H, s), 4.59 (2H, t), 3.82 (3H, s), 3.21 (2H, t), 2.05 (3H, s)
40	4.118	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.91-7.89 (1H, s), 7.82-7.75 (2H, m), 7.55-7.44 (2H, m), 7.40-7.35 (1H, t), 7.25-7.22 (1H, d), 5.06 (2H, s), 3.82 (3H, s), 2.05 (3H, s)
	4.119	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.89-7.85 (1H, m), 7.69-7.67 (1H, m), 7.63-7.53 (2H, m), 7.37 (2H, t), 7.25-7.22 (1H, m), 5.13 (2H, s), 3.82 (3H, s), 2.93 (2H, q), 2.05 (3H, s), 1.32 (3H, t)
45	4.120	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.89 (1H, s), 7.8 (2H, d), 7.56 (3H, m), 7.36 (1H, t), 7.22 (1H, d), 5.09 (2H, s), 3.83 (3H, s), 2.05 (3H, s)
	4.121	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.87 (1H, s), 7.67 (1H, s), 7.61-7.52 (2H, m), 7.39-7.33 (2H, m), 7.25-7.21 (1H, m), 5.13 (2H, s), 3.82 (3H, s), 2.59 (2H, s), 2.05 (2H, s)
50	4.122	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 7.99 (1H, s), 7.87 (1H, s), 7.55 (1H, d), 7.48-7.32 (2H, m), 7.23 (1H, d), 4.91 (2H, s), 3.81 (3H, s), 2.98 (6H, d), 2.05 (3H, s)
	4.124	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, s), 7.86 (1H, t), 7.55 (1H, d), 7.37 (1H, t), 7.23 (1H, d), 7.06 (2H, m), 6.42 (1H, d), 4.88 (2H, s), 3.8 (3H, s), 3.2 (2H, t), 2.82 (2H, t), 2.66 (3H, s), 2.05 (3H, s)
55	4.125	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.86 (1H, s), 7.55 (1H, d), 7.35 (1H, dd), 7.23 (1H, d), 7.07 (2H, m), 6.87 (1H, dd), 5.05 (2H, s), 3.82 (6H, s), 2.04 (3H, s)

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	Compound Nr	NMR-data
5	4.126	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.88 (1H, t), 7.53 (1H, d), 7.37 (1H, t), 7.23 (1H, d), 7.04 (1H, dd), 6.6 (1H, t), 4.91 (2H, s), 4.64 (2H, t), 3.83 (3H, s), 3.23 (2H, t), 2.05 (3H, s)
	4.127	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 7.89 (1H, t), 7.54 (1H, d), 7.39 (3H, m), 7.24 (1H, d), 5.03 (2H, s), 3.83 (3H, s), 2.06 (3H, s)
10	4.128	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.5 (1H, s), 8.9 (1H, d), 8.3 (2H, m), 7.9 (1H, d), 7.8-7.7 (1H, t), 7.6 (1H, d), 5.5 (2H, s), 4.2 (3H, s), 2.4 (3H, s)
	4.129	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, s), 8.37 (1H, d), 7.89 (1H, t), 7.64 (1H, dd), 7.54 (1H, d), 7.35 (2H, dd), 7.24 (1H, d), 5.12 (2H, s), 3.85 (3H, s), 2.9 (1H, q), 2.04 (3H, s)
15	4.130	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.20-8.17 (1H, m), 7.36-7.32 (2H, m), 7.20-7.17 (2H, m), 4.95 (2H, s), 3.88 (3H, s), 3.81 (3H, s), 2.1 (3H, s)
	4.131	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.20-8.16 (1H, m), 7.62-7.59 (1H, m), 7.48-7.46 (2H, m), 7.39-7.29 (2H, m), 5.02 (2H, s), 3.81 (3H, s), 2.1 (3H, s)
20	4.132	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.21-8.18 (1H, m), 7.59-7.51 (2H, m), 7.36-7.33 (3H, m), 5.03 (2H, s), 3.81 (3H, s), 2.1 (3H, s)
	4.133	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.5 (8H, m), 5.01 (2H, s), 3.85 (3H, s), 3.18 (3H, s), 2.5 (3H, s)
	4.134	¹ H-NMR (DMSO-d ₆), δ (ppm): 8.2-7.4 (8H, m), 5.2 (2H, s), 3.9 (3H, s), 3.2 (1H, s), 1.8 (3H, s)
25	4.135	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.17-8.15 (1H, m), 7.35-7.32 (2H, m), 7.25 (1H, s), 7.13-7.10 (1H, m), 6.70-6.67 (1H, m), 4.92 (2H, s), 4.52-4.46 (2H, t, 8.71Hz), 3.79 (3H, s), 3.16-3.10 (2H, t, 8.71Hz), 2.10 (3H, s)
	4.136	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.6-7.5 (3H, m), 7.3(3H, d), 6.9(2H, d), 5.0 (2H, s), 3.9 (3H, s), 3.7 (3H, s), 3.2 (3H, s), 1.8 (3H, s)
30	4.137	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.53 (3H, m), 7.33 (1H, d), 7.25 (2H, s), 7.12 (1H, d), 6.69 (1H, d), 4.94 (2H, s), 4.48 (2H, t), 3.94 (3H, s), 3.13 (5H, m), 1.83 (3H, s)
	4.138	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.64-7.51 (3H, m), 7.38-7.31 (1H, m), 7.22-7.11 (2H, m), 7.95-7.89 (1H, m), 5 (2H, s), 3.85 (3H, s), 3.82 (3H, s), 3.18 (3H, bs), 1.83 (3H, bs)
35	4.139	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.60-7.50 (3H, m), 7.36-7.31 (1H, m), 7.17-7.04 (2H, m), 5.03 (2H, s), 3.92 (3H, s), 3.86 (3H, s), 3.18 (3H, bs), 1.83 (3H, bs)
	4.140	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.6-7.2 (10H, m), 4.1 (2H, t), 4.1-4.0 (3H, s), 3.8 (3H, s), 2.9 (2H, t)
40	4.141	¹ H-NMR (CDCl ₃), δ (ppm): 10.1 (1H, bs), 8.01 (1H, t), 7.65 (1H, d), 7.4 (5H, m), 7.33 (1H, dd), 5.01 (2H, s), 4.02 (2H, s), 3.81 (3H, s), 3.38 (3H, s)
	4.142	¹ H-NMR (CDCl ₃), δ (ppm): 10.14 (1H, bs), 7.91 (1H, s), 7.58 (1H, d), 7.4 (5H, m), 7.25 (1H, dd), 5.01 (2H, s), 3.83 (3H, s), 3.62 (2H, t), 3.24 (3H, s), 2.54 (2H, t)
45	4.143	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.4 (4H, s), 7.30-7.23 (2H, m), 6.95-6.82 (2H, m), 5 (2H, s), 3.74 (3H, s)
	4.144	¹ H-NMR (CDCl ₃), δ (ppm): 10.4 (1H, s), 8.0 (1H, s), 7.6-6.9 (9H, m), 5.0 (2H, s), 3.8 (3H, s)
	4.145	¹ H-NMR (CDCl ₃), δ (ppm): 7.9 (1H, s), 7.6-7.3 (7H, m), 5.2 (2H, s), 3.9 (3H, s), 1.5-0.9 (6H, m)
50	4.146	¹ H-NMR (CDCl ₃), δ (ppm): 8.27 (1H, bs), 7.98 (1H, d), 7.82-7.78 (1H, m), 7.68 (1H, t), 7.44-7.36 (4H, m), 5.03 (2H, s), 3.85 (3H, s), 2.68 (3H, s)
	4.147	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.6-7.2 (5H, m), 7.2-7.1(2H,d), 6.9(2H,d), 4.0 (2H, t), 3.8 (3H, s), 3.7(3H, s), 2.8(2H, t)
55	4.148	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.6-7.1 (9H, m), 4.1 (2H, t), 3.8 (3H, s), 2.9 (2H, t)
	4.149	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.6-7.3 (6H, m), 6.9-6.7 (2H, m), 4.1 (2H, t), 4.1-4.0 (3H, s), 3.8 (3H, s), 3.8 (3H, s), 2.8 (2H, t)

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Compound Nr	NMR-data
5	4.150 ¹ H-NMR (DMSO-d ₆), δ (ppm): 8.1- 7.7 (6H, m), 7.3 (3H, m), 4.6 (2H, t), 4.5 (3H, s), 4.3 (3H, s), 3.4 (2H, t)
4.151	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, s), 7.88 (1H, s), 7.58-7.51 (1H, m), 7.37 (1H, t), 7.24-7.13 (3H, m), 6.88 (2H, d), 3.98 (2H, t), 3.81 (3H, s), 3.72 (3H, s), 2.8 (3H, t), 2.06 (3H, s)
10	4.152 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 7.9 (1H, s), 7.6-7.1 (7H, m), 4.0 (2H, t), 3.8 (3H, s), 2.9 (2H, t), 2.1 (3H, s)
4.153	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 7.9 (1H, s), 7.6 (1H, d), 7.4 (1H, t), 7.2 (2H, m), 6.8 (3H, m), 4.1 (2H, t), 3.8 (3H, s), 3.7 (3H, s), 2.9 (2H, t), 2.1 (3H, s)
15	4.154 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.18-8.15 (1H, m), 7.38-7.27 (7H, m), 5.01 (2H, s), 3.81 (3H, s), 2.1 (3H, s)
4.155	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.17 (1H, dd), 7.31 (2H, m), 7.15 (2H, m), 6.91 (1H, m), 4.97 (2H, s), 3.81 (3H, s), 3.8 (3H, s), 2.1 (3H, s)
20	4.156 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.17 (1H, dd), 7.38-7.31 (2H, m), 7.14-7.07 (2H, m), 5 (2H, s), 3.91 (3H, s), 3.81 (3H, s), 2.1 (3H, s)
4.157	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.22-8.18 (1H, m), 7.43-7.31 (3H, m), 7.21-7.11 (1H, m), 4.87 (2H, s), 3.83 (6H, s), 2.1 (3H, s)
25	4.158 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.9 (1H, s), 8.2-8.1 (1H, d), 7.4-7.2 (7H, m), 4.0 (2H, t), 3.8 (3H, s), 2.9-2.8 (2H, t), 2.1 (3H, s)
4.159	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, s), 8.35 (1H, s), 8.19 (1H, d), 7.96 (1H, d), 7.43-7.26 (2H, m), 5.07 (2H, s), 3.82 (3H, s), 2.10 (3H, s)
30	4.160 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.19-8.14 (1H, m), 7.38-7.31 (2H, m), 7.23 (1H, t), 6.95-6.82 (3H, m), 4.98 (2H, s), 3.81 (3H, s), 3.73 (3H, s), 2.1 (3H, s)
4.161	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.18 (1H, s), 7.95-7.90 (1H, m), 7.57-7.51 (1H, m), 7.32 (1H, t), 7.14-7.08 (2H, m), 5.01 (2H, s), 3.92 (3H, s), 3.78 (3H, s), 2.05 (3H, s)
35	4.162 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.14 (1H, d), 7.38-7.32 (2H, m), 7.06-7.04 (2H, m), 6.41 (1H, d), 4.86 (2H, s), 3.78 (3H, s), 3.2 (2H, t), 2.81 (2H, t), 2.65 (3H, s), 2.10 (3H, s)
4.163	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.18 (1H, dd), 7.25 (3H, m), 6.63 (1H, dd), 4.94 (2H, s), 4.55 (2H, t), 3.81 (3H, s), 3.09 (3H, t), 2.1 (3H, s)
40	4.164 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.16 (1H, m), 7.34-7.09 (5H, m), 6.89 (1H, t), 4.91 (2H, s), 3.79 (3H, s), 2.73 (6H, s), 2.1 (3H, s)
4.165	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.19 (1H, d), 7.35-7.08 (4H, m), 4.99 (2H, s), 3.96 (3H, s), 3.82 (3H, s), 2.1 (3H, s)
4.166	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.57-7.15 (10H, m), 3.88 (2H, t), 3.8 (3H, s), 2.65 (2H, t), 1.91 (2H, q)
45	4.167 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.58-7.36 (4H, m), 7.44-7.36 (4H, m), 5 (2H, s), 4.02 (2H, s), 3.8 (3H, s)
4.168	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.9 (1H, s), 8.2 (1H, s), 7.8 (2H, d), 7.6 (2H, d), 7.3 (2H, bs), 5.1 (2H, s), 3.8 (3H, s), 2.1 (3H, s)
50	4.169 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.21-8.16 (1H, s), 7.39-7.21 (4H, m), 7.09 (1H, d), 7 (1H, d), 6.86 (1H, t), 4.96 (2H, s), 3.83 (6H, s), 2.1 (3H, s)
4.170	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.19 (1H, dd), 7.88 (1H, t), 7.59 (1H, dd), 7.35 (3H, m), 5.08 (2H, s), 3.82 (3H, s), 2.1 (3H, s)
55	4.171 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.79 (1H, s), 8.18 (1H, d), 8.12-7.97 (2H, m), 7.43-7.22 (2H, m), 5.12 (2H, s), 3.82 (3H, s), 2.10 (3H, s)

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	Compound Nr	NMR-data
5	4.172	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.5 (1H, d), 8.22 (1H, d), 7.69 (1H, dd), 7.35-7.31 (2H, m), 7.23 (1H, d), 4.98 (2H, s), 3.79 (3H, s), 2.98 (1H, m), 2.1 (3H, s), 1.19 (6H, d)
	4.173	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.5 (1H, d), 8.22 (1H, d), 7.69 (1H, dd), 7.35-7.31 (2H, m), 7.23 (1H, d), 4.98 (2H, s), 3.79 (3H, s), 2.98 (1H, m), 2.1 (3H, s)
10	4.174	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.88 (1H, dd), 8.34 (1H, d), 8.19 (1H, d), 8.00-7.96 (2H, m), 7.77 (1H, dd), 7.52 (1H, dd), 7.35 (2H, m), 5.21 (2H, s), 3.83 (3H, s), 2.10 (3H, s)
	4.175	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.16 (1H, d), 7.98 (1H, d), 7.67 (1H, d), 7.54 (1H, d), 7.36 (3H, m), 6.93 (1H, d), 5.1 (2H, s), 3.8 (3H, s), 2.1 (3H, s)
15	4.176	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, s), 8.17-8.13 (1H, m), 7.39-7.25 (4H, m), 7.13 (2H, t), 4 (2H, t), 3.8 (3H, s), 2.86 (2H, t), 2.11 (3H, s)
	4.177	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.17 (1H, s), 7.45-7.44 (1H, m), 7.36-7.31 (3H, m), 7.10-7.08 (1H, m), 4.93 (2H, s), 3.83 (3H, s), 3.8 (3H, s), 2.1 (3H, s)
20	4.178	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 7.86 (1H, s), 7.55 (1H, d), 7.37 (1H, t), 7.22 (1H, d), 7.09-7.02 (2H, m), 6.65 (1H, t), 4.89 (2H, s), 3.8 (3H, s), 3.36 (4H, s), 2.05 (3H, s), 1.87 (4H, m)
	4.179	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 7.87 (1H, s), 7.54 (1H, d), 7.37 (1H, t), 7.24-7.11 (2H, m), 6.97 (1H, t), 4.94 (2H, s), 3.81 (3H, s), 3.72 (4H, m), 2.96 (4H, m), 2.04 (3H, s)
25	4.180	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.12 (1H, m), 7.37-7.27 (2H, m), 7.11-6.99 (2H, m), 5.09 (2H, s), 3.8 (6H, s), 2.09 (3H, s)
	4.181	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.18 (1H, m), 7.37-7.27 (2H, m), 7.69 (2H, d), 7.6 (2H, d), 7.35-7.32 (2H, m), 5.09 (2H, s), 3.81 (3H, s), 2.1 (3H, s)
30	4.182	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.9 (1H, bs), 8.2 (1H, d), 7.3 (6H, m), 5.8 (2H, s), 5 (2H, s), 4.4 (2H, s), 3.8 (3H, s), 3.3 (3H, s), 2.1 (3H, s)
	4.183	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.9 (1H, bs), 8.34 (1H, d), 8.13 (3H, m), 7.31 (2H, m), 7.01 (1H, dd), 5.13 (2H, s), 3.79 (3H, s), 2.1 (3H, s)
35	4.184	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.18 (1H, d), 7.37-7.32 (2H, m), 7.12 (1H, t), 6.90-6.84 (2H, m), 5 (2H, s), 3.82 (3H, s), 3.7 (3H, s), 2.1 (3H, s)
	4.185	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.16 (1H, dd), 7.38 (4H, m), 7.17 (3H, m), 4.99 (2H, s), 3.8 (3H, s), 2.1 (3H, s)
40	4.186	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.20-8.17 (3H, d), 7.67-7.64 (2H, d), 7.36-7.34-7.32 (2H, t), 5.13 (2H, s), 3.82 (3H, s), 2.1 (3H, s)
	4.187	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.93 (1H, s), 8.48 (1H, s), 8.21-8.13 (1H, m), 7.84-7.70 (3H, m), 7.52-7.47 (2H, m), 7.37-7.31 (2H, m), 6.53 (1H, s), 5.04 (2H, s), 3.81 (3H, s), 2.1 (3H, s)
45	4.188	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.18 (1H, d), 7.35-7.32 (2H, m), 6.79-6.71 (3H, m), 4.96 (2H, s), 3.81 (3H, s), 3.75 (3H, s), 2.1 (3H, s)
	4.189	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, s), 8.18 (1H, s), 7.53-7.22 (2H, m), 7.10 (1H, t), 6.89-6.53 (3H, m), 4.95 (2H, s), 3.80 (3H, s), 2.87 (6H, s), 2.10 (3H, s)
50	4.190	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 8 (1H, s), 7.89 (1H, m), 7.69 (1H, d), 7.59-7.54 (2H, m), 7.37 (1H, t), 7.25 (1H, d), 7.14 (1H, d), 5.16 (2H, s), 4.02 (3H, s), 3.84 (3H, s), 2.04 (3H, s)
	4.191	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.98 (1H, s), 8.20-8.11 (1H, m), 7.38-7.26 (2H, m), 6.90-6.075 (3H, m), 4.88 (2H, s), 4.2 (4H, s), 3.79 (3H, s), 2.1 (3H, s)
55	4.192	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 8.01 (1H, d), 7.86 (1H, t), 7.75 (1H, m), 7.60-7.53 (2H, m), 7.43 (2H, dd), 7.36 (2H, t), 7.23 (1H, m), 5.12 (2H, s), 4.01 (3H, s), 3.81 (3H, s), 2.04 (3H, s)
	4.193	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 8.28 (1H, s), 7.86 (1H, t), 7.67 (1H, s), 7.57-7.52 (2H, m), 7.36 (1H, t), 7.25 (2H, m), 5.06 (3H, s), 4.14 (3H, s), 3.81 (3H, s), 2.04 (3H, s)

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Compound Nr	NMR-data
5	4.194 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.16 (1H, dd), 7.33 (2H, m), 7.14 (1H, d), 6.78 (2H, m), 4.93 (2H, s), 4.48 (2H, t), 3.79 (3H, s), 3.12 (2H, t), 2.09 (3H, s)
	4.195 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.17 (1H, dd), 7.33 (2H, m), 7.01 (1H, t), 6.74 (1H, d), 6.67 (1H, d), 4.94 (2H, s), 4.53 (2H, t), 3.81 (3H, s), 3.24 (2H, t), 2.09 (3H, s)
10	4.196 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, bs), 8.21-8.20 (2H, m), 7.89 (1H, m), 7.56-7.53 (1H, m), 7.40-7.35 (2H, m), 7.26-7.23 (1H, m), 5.03 (2H, s), 3.82 (3H, s), 3.81 (3H, s), 2.05 (3H, s)
	4.197 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.11 (1H, bs), 7.90-7.89 (1H, s), 7.55-7.52 (1H, dm, 8.96Hz), 7.51-7.49 (1H, d, 8.19Hz), 7.40-7.35 (1H, t, 8.19Hz), 7.26-7.22 (1H, dm, 7.96Hz), 7.13-7.12 (1H, d, 1.79Hz), 6.89-6.86 (1H, dd, 8.19Hz, 1.79Hz), 4.99 (2H, s), 3.83 (6H, s), 2.05 (3H, s)
15	4.198 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, s), 7.90-7.88 (1H, m), 7.56-7.54 (1H, m), 7.41-7.20 (6H, m), 4.06-4.00 (2H, t), 3.81 (3H, s), 2.91-2.85 (2H, t), 2.06 (3H, s)
	4.199 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.19 (1H, d), 7.43-7.26 (2H, m), 7.17 (1H, t), 6.93 (1H, dd), 6.68 (1H, td), 4.91 (2H, s), 3.84 (3H, s), 3.82 (3H, s), 2.10 (3H, s)
20	4.200 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.20-8.17 (2H, m), 7.78 (1H, d), 7.73 (1H, t), 7.49 (1H, dd), 7.36-7.33 (3H, m), 6.55 (1H, t), 5.06 (2H, s), 3.82 (3H, s), 2.1 (3H, s)
	4.201 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.18 (1H, m), 7.45-7.24 (5H, m), 4.98 (2H, s), 3.8 (3H, s), 2.1 (3H, s)
25	4.202 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.19 (1H, m), 8 (1H, s), 7.7 (1H, d), 7.59 (1H, s), 7.34 (2H, m), 7.13 (1H, d), 5.15 (2H, s), 4.02 (3H, s), 3.82 (3H, s), 2.09 (3H, s)
	4.203 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.19 (1H, m), 7.63-7.35 (5H, m), 4.99 (2H, s), 3.81 (3H, s), 2.1 (3H, s)
30	4.204 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, bs), 8.20-8.15 (1H, m), 7.36-7.33 (3H, m), 7.06-6.98 (2H, m), 6.87-6.83 (1H, m), 4.94 (2H, s), 3.8 (3H, s), 2.75 (6H, s), 2.1 (3H, s)
	4.205 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, bs), 8.03-8.01 (1H, m), 7.60-7.57 (1H, d, 8.71Hz), 7.45-7.39 (3H, m), 7.18-7.12 (2H, tm, 8.71Hz), 4.99 (2H, s), 3.82 (3H, s), 2.11 (3H, s)
35	4.206 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.16 (1H, t), 7.33 (2H, m), 7.19 (1H, s), 7.10 (1H, dd), 6.61 (1H, d), 4.90 (2H, s), 3.79 (3H, s), 2.96 (2H, s), 2.10 (3H, s), 1.37 (6H, s)
	4.207 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.60-7.55 (1H, m), 7.50-7.30 (6H, m), 5 (2H, s), 4 (2H, s), 3.8 (3H, s)
40	4.208 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.87 (1H, s), 9.18-9.14 (1H, m), 7.42-7.29 (7H, m), 4.99 (2H, s), 3.8 (3H, s), 2.1 (3H, s)
	4.209 ¹ H-NMR (DMSO-d ₆), δ (ppm): 11.08 (1H, s), 10.13 (1H, s), 7.9 (1H, s), 7.55 (2H, t), 7.38 (1H, t), 7.26-7.22 (1H, m), 6.99-6.92 (2H, m), 4.98 (2H, s), 3.83 (3H, s), 2.05 (3H, s)
45	4.210 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.19 (1H, d), 7.54 (1H, t), 7.45 (1H, dd), 7.40-7.20 (3H, m), 5.00 (2H, s), 3.81 (3H, s), 2.10 (3H, s)
	4.211 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, s), 8.03 (1H, d), 7.65-7.34 (4H, m), 7.26 (1H, d), 5.01 (2H, s), 3.83 (3H, s), 2.11 (3H, s)
50	4.212 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, s), 8.04 (1H, d), 7.59 (1H, d), 7.42 (1H, dd), 7.17 (1H, t), 6.93 (1H, dd), 6.68 (1H, td), 4.91 (2H, s), 3.84 (6H, s), 2.11 (3H, s)
	4.213 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.17 (1H, d), 7.47-7.24 (4H, m), 7.03 (1H, t), 5.01 (2H, s), 3.81 (3H, s), 2.1 (3H, s)
55	4.214 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.17 (1H, d), 7.34 (2H, m), 7.17 (1H, d), 7.09 (1H, s), 6.92 (1H, d), 4.9 (2H, s), 3.86 (3H, s), 3.82 (3H, s), 2.1 (3H, s)
	4.215 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.17 (1H, d), 7.36-7.32 (3H, m), 7.15 (1H, d), 6.93 (1H, dd), 4.99 (2H, s), 3.83 (3H, s), 3.8 (3H, s), 2.1 (3H, s)

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Compound Nr	NMR-data
5	4.216 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, bs), 8.14-8.11 (1H, m), 7.39-7.32 (2H, m), 7.28-7.16 (2H, m), 7.06-6.99 (1H, m), 4.06-4.01 (2H, t, 7.68Hz), 3.79 (3H, s), 2.93-2.89 (2H, t, 7.68Hz), 2.11 (3H, s)
	4.217 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.12 (1H, s), 8.03 (1H, d), 7.88 (1H, t), 7.66 (1H, dd), 7.54 (1H, d), 7.37 (1H, t), 7.23 (1H, d), 4.98 (2H, s), 3.92 (3H, s), 3.81 (3H, s), 2.05 (3H, s)
10	4.218 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.63 (1H, bs), 8.05-8.03 (1H, m), 7.60-7.57 (1H, d, 8.7Hz), 7.43-7.39 (1H, dd, 8.7Hz, 2.56Hz), 7.19-7.16 (1H, d, 8.19Hz), 7.10-7.09 (1H, d, 1.79Hz), 6.93-6.90 (1H, dd, 8.19Hz, 2.56Hz), 4.91 (2H, s), 3.86 (3H, s), 3.84 (3H, s), 2.11 (3H, s)
15	4.219 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.63 (1H, bs), 8.03 (1H, d, 2.56Hz), 7.60-7.57 (1H, d, 8.96Hz), 7.43-7.39 (1H, dd, 8.96Hz, 2.56Hz), 7.36-7.33 (1H, d, 8.19Hz), 7.16-7.15 (1H, d, 1.79Hz), 6.95-6.92 (1H, dd, 8.19Hz, 1.79Hz), 5 (2H, s), 3.84 (3H, s), 3.83 (3H, s), 2.11 (3H, s)
	4.220 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.21-8.19 (1H, m), 7.39-7.30 (2H, m), 7.05-6.98 (2H, m), 4.97 (2H, s), 3.86 (3H, s), 3.82 (3H, s), 2.1 (3H, s)
20	4.221 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, s), 8.02 (1H, s), 7.58 (1H, d), 7.52-7.35 (3H, m), 7.24 (1H, d), 5.02 (2H, s), 3.84 (3H, s), 2.11 (3H, s)
	4.222 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, s), 8.02 (1H, s), 7.59 (1H, d), 7.53-7.36 (2H, m), 7.25 (1H, t), 7.04 (1H, t), 5.02 (2H, s), 3.83 (3H, s), 2.11 (3H, s)
25	4.223 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, s), 8.03 (1H, s), 7.80-7.52 (5H, m), 7.41 (1H, dd), 5.10 (2H, s), 3.84 (3H, s), 2.11 (3H, s)
	4.224 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.87 (1H, bs), 8.18 (1H, d), 7.35-7.32 (2H, m), 7.16-7.10 (2H, m), 7 (1H, dd), 5.02 (2H, s), 3.83 (3H, s), 2.66 (6H, s), 2.09 (3H, s)
30	4.225 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.63 (1H, bs), 8.03 (1H, d), 7.58 (1H, d), 7.41 (1H, dd), 7.16-7.11 (2H, m), 7 (1H, dd), 5.02 (2H, s), 3.85 (3H, s), 2.67 (6H, s), 2.1 (3H, s)
	4.226 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.63 (1H, bs), 8.02-8.01 (1H, m), 7.60-7.57 (1H, m), 7.42-7.39 (1H, m), 7.05-6.97 (2H, m), 6.87-6.82 (1H, m), 4.94 (2H, s), 3.82 (3H, s), 2.75 (6H, s), 2.11 (3H, s)
35	4.227 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, s), 8.02 (1H, d), 7.59 (1H, d), 7.47-7.21 (6H, m), 5.02 (2H, s), 3.83 (3H, s), 2.11 (3H, s)
	4.228 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.19 (1H, t), 7.34 (2H, m), 7.11 (1H, dd), 6.95 (1H, dd), 6.77 (1H, dt), 5.02 (2H, s), 3.83 (3H, s), 2.66 (6H, s), 2.10 (3H, s)
40	4.229 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, s), 8.05-8.02 (1H, m), 7.58 (1H, d), 7.43-7.39 (1H, m), 7.19-7.10 (2H, m), 6.95-6.88 (1H, m), 4.97 (2H, s), 3.82 (3H, s), 3.81 (3H, s), 2.11 (3H, s)
	4.230 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.21 (1H, s), 7.74 (1H, t), 7.56 (1H, d), 7.49-7.26 (3H, m), 5.09 (2H, s), 3.83 (3H, s), 2.10 (3H, s)
45	4.231 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.17 (1H, d), 7.36-7.29 (3H, m), 7.17 (1H, s), 6.96 (1H, dd), 4.96 (2H, s), 3.8 (3H, s), 2.71 (6H, s), 2.1 (3H, s)
	4.232 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, bs), 8.02 (1H, s), 7.5 (1H, d), 7.4 (1H, dd), 7.3 (1H, d), 7.18 (1H, s), 6.96 (1H, d), 4.97 (2H, s), 3.82 (3H, s), 2.71 (6H, s), 2.11 (3H, s)
50	4.233 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.19 (1H, s), 7.88 (1H, s), 7.69 (2H, s), 7.43-7.23 (2H, m), 5.06 (2H, s), 3.81 (3H, s), 2.10 (3H, s)
	4.234 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.63 (1H, s), 8.51 (1H, d), 8.02 (1H, d), 7.70 (1H, dd), 7.58 (1H, d), 7.41 (1H, dd), 7.23 (1H, d), 4.99 (2H, s), 3.81 (3H, s), 2.98 (1H, q), 2.11 (3H, s), 1.20 (6H, d)
55	4.235 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.18 (1H, s), 7.73-7.55 (2H, m), 7.52-7.23 (3H, m), 5.04 (2H, s), 3.81 (3H, s), 2.11 (3H, s)
	4.236 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, s), 8.04-8.01 (1H, m), 7.57 (1H, d), 7.43-7.39 (1H, m), 7.23 (1H, t), 6.95-6.82 (3H, m), 4.98 (2H, s), 3.83 (3H, s), 3.73 (3H, s), 2.11 (3H, s)

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Compound Nr	NMR-data
5 4.237	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.64 (1H, bs), 8.03-8.00 (1H, m), 7.60-7.57 (1H, m), 7.43-7.39 (1H, m), 7.25 (1H, s), 7.13-7.10 (1H, m), 6.70-6.67 (1H, m), 4.92 (2H, s), 4.52-4.46 (2H, t, 8.71Hz), 3.81 (3H, s), 3.16-3.10 (2H, t, 8.71Hz), 2.11 (3H, s)
4.238	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.66 (1H, bs), 8.06-8.01 (1H, m), 7.60-7.57 (1H, m), 7.44-7.39 (1H, m), 7.12-7.05 (2H, m), 4.99 (2H, s), 3.96 (3H, s), 3.84 (3H, s), 2.11 (3H, s)
10 4.239	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.69-7.53 (2H, m), 7.47-7.24 (6H, m), 5.00 (2H, s), 3.82 (3H, s)
4.240	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.63 (1H, bs), 8.03 (1H, s), 7.88 (1H, s), 7.69 (2H, s), 7.58 (1H, d), 7.39 (1H, dd), 5.06 (2H, s), 3.82 (3H, s), 2.11 (3H, s)
15 4.241	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.17 (1H, dd), 7.30 (3H, m), 7.17 (1H, dd), 7.91 (1H, dd), 4.97 (2H, s), 4.64 (1H, hex), 3.80 (3H, s), 2.10 (3H, s), 1.28 (6H, d)
4.242	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.63 (1H, bs), 8.02 (1H, s), 7.69 (1H, d), 7.57 (2H, m), 7.42 (1H, d), 7.14 (1H, d), 5.15 (2H, s), 4.01 (3H, s), 3.84 (3H, s), 2.1 (3H, s)
20 4.243	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.87 (1H, bs), 8.17 (1H, d), 7.42-7.32 (4H, m), 7.17 (1H, dd), 4.99 (2H, s), 3.8 (3H, s), 3.31 (1H, m), 2.1 (3H, s), 1.19 (6H, d)
4.244	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.16 (1H, d), 7.35-6.97 (5H, m), 4.93 (2H, s), 3.81 (3H, s), 3.72 (4H, s), 2.96 (4H, s), 2.1 (3H, s)
25 4.245	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.17 (1H, bs), 9.88 (1H, bs), 8.16 (1H, dd), 7.33 (3H, m), 6.98 (1H, d), 6.81 (1H, dd), 4.90 (2H, s), 3.80 (3H, s), 2.10 (3H, s)
4.246	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.16 (1H, dd), 7.32 (2H, m), 7.21 (1H, d), 7.10 (1H, dd), 6.69 (1H, dd), 4.93 (2H, s), 4.18 (2H, s), 3.79 (3H, s), 2.10 (3H, s), 1.25 (6H, s)
30 4.247	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.17 (1H, dd), 7.34 (2H, m), 7.12 (1H, d), 6.86 (1H, dd), 6.77 (1H, s), 4.93 (2H, s), 4.18 (2H, s), 3.79 (3H, s), 2.10 (3H, s), 1.39 (3H, s), 1.25 (3H, s)
4.248	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.20-8.14 (1H, s), 7.39-7.25 (3H, m), 7.17-7.10 (2H, m), 4.97 (2H, s), 3.8 (3H, s), 3.16-3.07 (1H, m), 1.18 (6H, d)
35 4.249	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.18 (1H, s), 7.95-7.90 (1H, m), 7.58-7.52 (1H, m), 7.36-7.29 (2H, m), 7.17-7.14 (1H, m), 6.97-6.93 (1H, m), 4.97 (2H, s), 3.77 (3H, s), 2.71 (6H, s), 2.05 (3H, s)
4.250	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.19 (1H, d), 7.66 (1H, d), 7.36-7.32 (3H, m), 7.23 (1H, s), 7.06 (1H, d), 5.06 (2H, s), 3.91 (3H, s), 3.82 (3H, s), 2.1 (3H, s)
4.251	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.68-7.58 (4H, m), 7.40-7.28 (3H, m), 5 (2H, s), 3.83 (3H, s)
40 4.252	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.17 (1H, dd), 7.38-7.15 (4H, m), 5.02 (2H, s), 3.88 (3H, s), 3.81 (3H, s), 2.09 (3H, s)
4.253	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.62 (1H, d), 7.59 (1H, d), 7.32 (2H, t), 7.26 (1H, dd), 7.15 (1H, t), 5.04 (3H, s), 3.88 (3H, s), 3.83 (3H, s)
45 4.254	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.89 (1H, s), 8.18 (1H, s), 7.68-7.22 (3H, m), 7.12 (1H, m), 6.88 (1H, d), 4.99 (2H, s), 3.82 (6H, d), 2.10 (3H, s)
4.255	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.17 (1H, d), 7.45-7.26 (2H, m), 7.12 (1H, d), 6.96 (1H, s), 6.88 (1H, d), 4.97 (2H, s), 3.80 (3H, s), 3.76 (3H, s), 3.19 (1H, q), 2.10 (3H, s), 1.12 (6H, d)
50 4.256	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.69-7.57 (2H, m), 7.40-7.26 (2H, t), 7.12 (1H, d), 6.97 (1H, s), 6.89 (1H, d), 4.98 (2H, s), 3.82 (3H, s), 3.76 (3H, s), 3.19 (1H, q), 1.12 (6H, d)
4.257	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.20-8.14 (1H, m), 7.40-7.27 (5H, m), 7.21 (1H, d), 6.97 (1H, bs), 6.76 (2H, d), 4.94 (2H, s), 3.8 (3H, s), 3.31-3.25 (4H, m), 2.1 (3H, s), 1.89-1.83 (4H, m)
55 4.258	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.18 (1H, dd), 7.76 (2H, m), 7.48 (1H, t), 7.32 (2H, m), 5.05 (2H, s), 3.81 (3H, s), 2.1 (3H, s)

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Compound Nr	NMR-data
5	4.259 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.15 (1H, dd), 7.33 (2H, m), 7.06 (2H, m), 5.00 (2H, s), 4.35 (1H, m), 3.81 (3H, s), 2.10 (3H, s), 1.26 (6H, d)
4.260	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.44 (1H, d), 8.20 (1H, d), 8.14 (1H, d), 7.43-7.25 (2H, m), 5.04 (2H, s), 3.82 (3H, s), 2.11 (3H, s)
10	4.261 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.19 (1H, dd), 7.54 (1H, dd), 7.31 (3H, m), 5.08 (2H, s), 4.06 (3H, d), 3.83 (3H, s), 2.10 (3H, s)
4.262	¹ H-NMR (DMSO-d ₆), δ (ppm): 8.44 (1H, d), 8.14 (1H, d), 7.68-7.55 (2H, m), 7.32 (2H, t), 5.06 (2H, s), 3.83 (3H, s)
15	4.263 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.18 (1H, dd), 7.33 (3H, m), 7.18 (1H, s), 7.02 (1H, d), 4.97 (2H, s), 3.80 (3H, s), 3.73 (4H, s), 2.96 (4H, s), 2.10 (3H, s)
4.264	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.19 (1H, dd), 7.55 (1H, d), 7.51 (1H, s), 7.39-7.31 (2H, m), 7.2 (1H, d), 5.04 (2H, s), 3.81 (3H, s), 2.66 (6H, s), 2.1 (3H, s)
20	4.265 ¹ H-NMR (DMSO-d ₆), δ (ppm): 7.59 (2H, m), 7.31 (3H, m), 6.93 (1H, d), 6.91 (1H, dd), 4.99 (2H, s), 4.64 (1H, hex), 3.82 (3H, s), 1.28 (6H, d)
4.266	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.74-7.43 (6H, m), 5.11 (1H, s), 3.87 (1H, s), 3.12 (1H, s), 1.79 (1H, s)
4.267	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.16 (1H, dd), 7.49 (1H, d), 7.33 (4H, m), 4.99 (2H, s), 4.45 (2H, s), 3.90 (3H, s), 3.35 (3H, s), 2.10 (3H, s)
25	4.268 ¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, bs), 8.17 (1H, dd), 7.62 (1H, m), 7.33 (2H, m), 7.06 (2H, m), 6.95 (1H, m), 4.95 (2H, s), 3.80 (4H, s), 3.72 (3H, m), 2.98 (4H, m), 2.10 (3H, s)
4.269	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.88 (1H, s), 8.30-8.13 (2H, m), 7.84 (1H, d), 7.43-7.25 (2H, m), 5.02 (2H, s), 3.80 (3H, s), 3.20 (1H, q), 2.10 (3H, s), 1.21 (6H, d)
30	5.01 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.2 (1H, bs), 7.93 (1H, s), 7.67 (1H, dt, 1Hz, 8Hz), 7.5 (5H, m), 7.3 (1H, dt, 1Hz, 8Hz), 5.11 (2H, s), 3.26 (1H, hep, 6Hz), 2.16 (3H, s), 1.27 (6H, d, 6Hz)
5.02	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, bs), 7.83 (1H, t, 2Hz), 7.56 (1H, dt, 1Hz, 8Hz), 7.38 (5H, m), 7.18 (1H, dt, 1Hz, 8Hz), 5 (2H, s), 2.62 (2H, q, 7Hz), 2.05 (3H, s), 1.13 (3H, t, 7Hz)
35	5.03 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.13 (1H, bs), 7.82 (1H, s), 7.56 (1H, d, 8Hz), 7.38 (5H, m), 7.16 (1H, d, 8Hz), 5 (2H, s), 2.59 (2H, t, 7Hz), 2.05 (3H, s), 1.55 (2H, m), 1.39 (2H, m), 0.89 (3H, t, 7Hz)
5.04	¹ H-NMR (DMSO-d ₆), δ (ppm): 8.1 (1H, d), 8.0 (1H, m), 7.7 (2H, m), 7.5 (2H, m), 7.2 (2H, t)
40	5.05 ¹ H-NMR (DMSO-d ₆), δ (ppm): 2.05 (3H, s), 2.2 (6H, s), 2.64 (2H, t, 5Hz), 4.2 (2H, t, 5Hz), 4.99 (1H, s), 7.22 (1H, dt, 1Hz, 8Hz), 7.4 (5H, m), 7.53 (1H, dt, 1Hz, 8Hz), 7.88 (1H, t, 2Hz), 10.11 (1H, bs)
5.06	¹ H-NMR (DMSO-d ₆), δ (ppm): 8.2-7.2 (8H, m), 5.1 (2H, s), 4.3 (2H, t), 3.7 (2H, t), 3.4 (3H, s), 2.0 (3H, s)
45	5.07 ¹ H-NMR (DMSO-d ₆), δ (ppm): 2.05 (3H, s), 3.67 (2H, dd, 4Hz, 2Hz), 4.24 (2H, dd, 4Hz, 2Hz), 5 (2H, s), 7.22 (1H, dt, 1Hz, 8Hz), 7.4 (5H, m), 7.53 (1H, dt, 1Hz, 8Hz), 7.85 (1H, t, 2Hz), 10.11 (1H, bs)
5.08	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.07 (1H, s), 7.85 (1H, s), 7.55-7.50 (1H, m), 7.41-7.22 (6H, m), 7.04-6.99 (1H, m), 5.02 (2H, s), 2.7 (3H, d)
50	5.09 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 7.87 (1H, s), 7.55-7.52 (1H, m), 7.42-7.33 (5H, m), 7.24-7.20 (1H, m), 4.98 (2H, s), 4.19 (1H, q), 2.05 (3H, s), 1.34 (3H, t)
5.10	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.16 (1H, s), 7.81 (1H, s), 7.59-7.57 (1H, m), 7.42-7.36 (1H, m), 7.29-7.27 (2H, m), 7.17-7.11 (3H, m), 4.97 (2H, s), 2.27 (3H, s), 2.05 (3H, s)
55	5.11 ¹ H-NMR (DMSO-d ₆), δ (ppm): 10.1 (1H, s), 7.87 (1H, s), 7.54-7.51 (1H, m), 7.43-7.33 (5H, m), 7.23-7.19 (1H, m), 4.98 (2H, s), 4.97-4.87 (1H, m), 2.05 (3H, s), 1.33 (3H, s), 1.31 (3H, s)
5.12	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.17 (1H, s), 7.83 (1H, s), 7.60-7.54 (1H, s), 7.46-7.37 (5H, m), 5 (2H, s), 2.05 (3H, s)

(continued)

Compound Nr	NMR-data
5.13	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.07 (1H, s), 7.87 (1H, s), 7.53-7.49 (1H, m), 7.38-7.23 (6H, m), 5 (2H, s), 2.95 (6H, s), 2.04 (3H, s)
5.14	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.14 (1H, bs), 7.82 (1H, t), 7.56 (1H, dd), 7.4 (5H, m), 7.17 (1H, dd), 5.31 (1H, t, 6Hz), 5 (2H, s), 4.4 (2H, d), 2.05 (3H, s)
5.15	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.27 (1H, bs), 10.09 (1H, bs), 7.98 (1H, t), 7.47 (6H, m), 7.21 (1H, dq), 5.02 (2H, s), 4.3 (2H, s), 2.87 (6H, s), 2.06 (3H, s)
5.16	¹ H-NMR (DMSO-d ₆), δ (ppm): 10.15 (1H, bs), 7.82 (1H, t), 7.57 (1H, dd), 7.4 (5H, m), 7.17 (1H, dd), 5 (2H, s), 4.34 (2H, s), 2.05 (3H, s)
5.17	¹ H-NMR (DMSO-d ₆), δ (ppm): 7.84 (1H, bs), 7.45 (5H, m), 7.3 (3H, m), 5.15 (2H, s), 3.97 (3H, s), 2.2 (3H, s)
5.18	¹ H-NMR (CDCl ₃), δ (ppm): 10.22 (1H, bs), 9.26 (1H, s), 7.93 (1H, t, 2Hz), 7.62 (1H, dt, 1Hz, 8Hz), 7.45 (5H, m), 7.22 (1H, dt, 1Hz, 8Hz), 5.07 (2H, s), 2.06 (3H, s)
5.19	¹ H-NMR (DMSO-d ₆), δ (ppm): 9.87 (1H, bs), 8.16-8.13 (1H, m), 7.45-7.40 (2H, m), 7.35-7.30 (2H, m), 7.18-7.12 (2H, m), 4.99 (2H, s), 4.23-4.20 (2H, m), 3.67-3.64 (2H, m), 3.31 (3H, s), 2.1 (3H, s)

PHARMACOLOGY

[0146] The compounds provided in this invention are positive allosteric modulators of metabotropic GABA receptors; in particular they are positive allosteric modulators of GABA_B receptors. The compounds of the present invention do not appear to bind to the GABA recognition site, the orthosteric ligand site, but instead to an allosteric site within the seven transmembrane region of the receptor. In the presence of GABA or an agonist of GABA_B receptor, the compounds of this invention increase the GABA_B response.

[0147] Hence, the present invention relates to a compound or a pharmaceutical composition for the manufacture of a medicament for treating or preventing a condition in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of GABA_B positive allosteric modulators.

[0148] The pharmacological activity of positive allosteric modulators at GABA_B receptors is demonstrated in the following paragraphs.

EXAMPLE A

[³⁵S]GTPγS binding assay

[0149] The [³⁵S]GTPγS binding is a functional membrane-based assay used to study G-protein coupled receptor (GPCR) function. This method consists of using a binding assay to assess the initial step in receptor-mediated G protein activation in membranes prepared from cells expressing recombinant GPCR or using membranes from discrete areas of the rat brain. In brief, the assay measures the activation of G proteins by catalyzing the exchange of guanosine 5'-diphosphate (GDP) by guanosine 5'-triphosphate (GTP) at the α subunit. The GTP-bound G proteins dissociate into two subunits, Gα-GTP and Gβγ, which in turn regulate intracellular enzymes and ion channels. GTP is rapidly hydrolysed by the Gα-subunit (GTPases) and the G protein is deactivated and ready for a new GTP exchange cycle (Harper, Curr. Protoc. Pharmacol. 1998, 2.6, 1-10, John Wiley & Sons, Inc.). [³⁵S]GTPγS, a non-hydrolyzed analogue of GTP, is used for this purpose.

[0150] This method is widely used to study receptor activation of G protein in membranes prepared from rat brain tissue, including GABA_B receptors (Urwyler et al, The Journal of Pharmacol. and Exp Ther. 2003, 307, 322-330; Olanas et al, Neurochemistry Intern. 2005, 46, 149-158). GABA_B receptors are expressed in the rat brain cortex (Odagaki and Yamauchi, Basic Clin. Pharmacol. Toxicol. 2004, 94, 89-98) and are coupled to Gα_{o/i}-protein, a preferential coupling for this method.

[0151] [³⁵S]GTPγS binding assay using cortical rat brain membranes preparation was used and adapted from Olanas et al (Neurochemistry Intern. 2005, 46, 149-158) for the detection of the positive allosteric modulator properties of the compounds of this invention on native rat GABA_B receptors.

[0152] Membrane preparation. Cortices were dissected out from brains of 200-300 g Sprague-Dawley rats (Charles River Laboratories, L'Arbresle, France). Tissues were homogenized in 6 volumes (vol/wt) of 10% sucrose at 4°C using

a glass-teflon homogenizer. The homogenate was centrifuged at 1,250g for 10 min, and the supernatant was centrifuged at 40,000g for 20 min (4°C). The pellet was resuspended in 25 ml water using a Polytron disrupter (Kinematica AG, Luzern, Switzerland) and centrifuged for 10 min at 3000 g. (4°C). The supernatant was centrifuged at 40,000g for 20 min (4°C). The supernatant was discarded and the pellet washed twice by resuspension in 10 volumes 5 mM HEPES-KOH, pH 7.4. The homogenate was frozen and thawed twice and centrifuged at 40,000g for 20 min. The final pellet was resuspended in 5 mM HEPES-KOH, pH 7.4 and stored at -80°C before its use. Protein concentration was determined by the Bradford method (Bio-Rad protein assay, Reinach, Switzerland) with bovine serum albumin as standard.

[0153] [³⁵S]GTPγS binding assay. Measurement of GABA_B receptor positive allosteric modulator properties in rat cortical membranes was performed as follows: rat cortical membrane (1.5 μg) were incubated in 96-well microplates for 15 min at 30°C in assay buffer (50 mM HEPES pH 7.4, 100 mM NaCl, 5 mM MgCl₂, 10 μM GDP, 10 μg/ml saponin, 1 mM CaCl₂) with increasing concentrations of positive allosteric modulator (from 1 nM to 60 μM) and a minimal concentration of GABA or Baclofen, a selective GABA_B receptor agonist, that has been determined in previous experiments to correspond to the EC₅₀, a concentration that gives 50 % of the maximal response of the agonist, and is in accordance with published data (Olianas et al, *Neurochemistry Intern.* 2005, 46, 149-158). After addition of 0.1 nM [³⁵S]GTPγS to achieve a total reaction volume of 200 μl, microplates were shaken for 1 min and further incubated at 30°C for 30 min. The incubation was stopped by rapid vacuum filtration over glass-fiber filter plates (Unifilter 96-well GF/C filter plates, Perkin-Elmer, Schwerzenbach, Switzerland) using a 96-well plate cell harvester (Filtermate, Perkin-Elmer, Downers Grove, USA). The Unifilter plate was washed three times with 300 μl of ice-cold wash buffer (20 mM HEPES pH 7.4, 100 mM NaCl). When filters were dry, 40 μl of liquid scintillation cocktail (Microscint 20) was added to each well. The amount of membrane-bound [³⁵S]GTPγS was measured using a 96-well plate reader (Top-Count, Perkin-Elmer, Downers Grove, USA). Non specific [³⁵S]GTPγS binding is determined in the presence of 10 μM of GTP.

[0154] Data analysis. The concentration-response curves of representative compounds of the present invention in the presence of EC₅₀ of GABA_B receptor agonist were generated using the Prism Graph-Pad program (Graph Pad Software Inc, San Diego, USA). The curves were fitted to a four-parameter logistic equation ($Y = \text{Bottom} + (\text{Top} - \text{Bottom}) / (1 + 10^{-(\text{LogEC}_{50} - X) \cdot \text{Hill Slope}})$) allowing determination of EC₅₀ values. Each curve was performed using duplicate sample per data point and 10 concentrations. The concentration-response curves of a selective GABA_B receptor agonist in the absence or in the presence of representative compounds of the present invention were also generated using Prism Graph-Pad program (Graph Pad Software Inc, San Diego, USA). The curves were fitted to a four-parameter logistic equation ($Y = \text{Bottom} + (\text{Top} - \text{Bottom}) / (1 + 10^{-(\text{LogEC}_{50} - X) \cdot \text{Hill Slope}})$) allowing determination of EC₅₀ values of the selective GABA_B receptor agonist. Each curve was performed using duplicate sample per data point and 10 concentrations.

[0155] Data presented in Figure A represent the ability of 10 μM of the Compound **4.02** to increase the [GTPγ³⁵S] binding induced by 15 μM of Baclofen, a GABA_B receptor agonist. Said example has no statistically significant agonistic activity when tested in the absence of 15 μM Baclofen (1% of maximal response, as a comparison, buffer is 0% of maximal response). Instead, when compounds are added together with a GABA_B receptors agonist like Baclofen, the effect measured is significantly potentiated compared to the effect of the agonist alone at the same concentration. Each bar graph is the mean and S.E.M. of duplicate data points and is representative of at least two independent experiments.

[0156] Table 8 shows representative compounds of the present invention that were clustered into four classes according to their ability (EC₅₀) to potentiate an EC₅₀ of GABA_B receptor agonist such as Baclofen. Class A: EC₅₀ < 100 nM; Class B: 100 nM < EC₅₀ < 500 nM; Class C: 500 nM < EC₅₀ < 1000 nM; Class D: EC₅₀ > 1000 nM.

Table 8: Summary of activity-data

Compound Nr	EC ₅₀
4.02	B
4.05	A
4.13	D
4.43	D
4.52	D
4.118	A
4.135	B
4.160	D
4.185	B

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(continued)

	Compound Nr	EC ₅₀
5	4.191	C
	4.192	B
	4.197	A
	4.198	B
10	4.204	A
	4.205	B
	4.207	A
	4.209	A
15	4.211	A
	4.219	D
	4.231	A
20	4.232	A
	4.233	A
	4.234	A
25	4.236	A
	4.237	D
	4.239	C
	4.241	A
30	4.242	A
	4.244	A
	4.247	A
35	4.248	A
	4.251	A
	4.253	A
	4.254	D
40	4.255	A
	4.260	A
	4.262	B
45	4.266	A
	4.268	A
	4.270	B

50 **[0157]** Thus, the positive allosteric modulators provided in the present invention are expected to increase the effectiveness of GABA or GABA_B receptor agonists at GABA_B receptors, and therefore, these positive allosteric modulators are expected to be useful for treatment of various neurological and psychiatric disorders associated with GABA dysfunction described to be treated herein and others that can be treated by such positive allosteric modulators.

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EXAMPLE B

Marble Burying test, model of anxiety in mice

[0158] Anxiety models in rodents are used as standard tests to demonstrate anxiolytic-like properties of novel compounds. Mice exhibit a tendency to bury harmless novel objects when encountered in a test cage. Marble burying behavior in mice is reduced by compounds which are efficacious anxiolytics in humans. Thus, marble burying in mice has been used as a model for the prediction of anxiolytic-like effects of compounds (Millan, M.J. et al., *Neuropharmacology* 2002, 42, 677-684).

[0159] Subjects: The present studies were performed in accordance with the animal care and use policies of Addex Pharmaceuticals and the EEC directives on the protection of animals used for experimental and other scientific purposes (86/609/EEC and subsequent revisions). Male C57BL6/j mice (20-30 g), were group housed in a temperature and humidity controlled facility on a 12 hour light /dark cycle for at least 5 days before use. Mice had access to food and water ad libitum except during marble burying experiments.

[0160] Assessment of marble burying: The effect of compounds on marble burying behaviour in mice was tested. On the day of the test, animals were marked on their tails and weighed in a separate preparation room, 1 hour before drug administration. Test compound or vehicle was administered p.o. 60 minutes prior to the test session. Marble burying behaviour was tested in a separate experimental room. For the test, mice were placed individually into clear plastic cages (16 x 22 x 14 cm) with 5 cm of sawdust and 10 marbles evenly spaced against the walls of the cage. The mice were left undisturbed in the cages for 30 minutes. After removal of the mice from the test cages, the number of marbles buried was counted. A marble was considered buried if it was 2/3 or more covered.

[0161] Compound administration: Test compounds were dissolved in a solution of 1% CMC. Test compounds were administered 60 minutes before testing by oral gavage (p.o.) in a volume of 10 mL/kg. In each assay, a vehicle and reference positive control groups were tested.

[0162] Statistical analyses: Statistical analyses were performed using GraphPad PRISM version 4.01 statistical software (GraphPad, San Diego, CA, USA). Data were analyzed using a non-parametric Kruskal-Wallis test followed by a Dunn's multiple comparisons test. The significance level was set at $p < 0.05$ when compared to the vehicle group.

[0163] Effect of compounds in the marble burying test in mice: At the doses of about 3 to 30 mg/kg, selected compounds of the invention significantly attenuate the marble burying behavior in mice.

EXAMPLE C

Vogel conflict drinking test, model of anxiety in rats

[0164] Anxiety models in rodents are used as standard tests to demonstrate anxiolytic-like properties of novel compounds. The Vogel conflict drinking model involves the conflict between thirsts and receiving mild shocks for drinking water (punished drinking). Water-deprived rats are placed in a chamber and they receive mild electric shocks every time they drink. The shocks suppress drinking and anxiolytics reverse this shock-induced suppression of drinking. The Vogel conflict drinking model was first proposed as a screening model for anxiolytics (Vogel, J.R. et al., *Psychopharmacologia* 1971, 21:1-7) and is widely accepted as a robust model for testing the anxiolytic-like properties of compounds (Millan, M.J. and Brocco M., *Eur. J. Pharm.* 2003, 463:67-96).

[0165] Subjects: The present studies were performed in accordance with the animal care and use policies of Addex Pharmaceuticals and the EEC directives on the protection of animals used for experimental and other scientific purposes (86/609/EEC and subsequent revisions). Male Sprague-Dawley rats (320 - 420 g) were housed in a temperature and humidity controlled facility on a 12 hours light /dark cycle for at least 5 days before use. Rats had access to food ad libitum except during Vogel conflict drinking model experiments. Rats had access to water ad libitum until 48 hours prior to the test session.

[0166] Assessment of Vogel conflict drinking: The effect of compounds on drinking in the Vogel conflict drinking model in rats was tested. Test chambers are housed in sound-attenuating boxes and each chamber contains a stainless steel drinking spout and a steel grid floor (MedAssociates, Georgia, Vermont, USA). Forty-eight hours prior to the test session, rats were habituated to the test chambers for 10 minutes. Water was removed from the rats immediately after the habituation session. Twenty-four hours before the test session, rats were again placed into the test chambers and allowed to drink for 4 minutes. Rats were then allowed 1 hour of access to water and then water was removed. On the test day, rats were brought to the test room at least 30 minutes before the test session. Rats were placed individually into the test chamber for a 5 minute session. Rats received a shock every 20th lick on the drinking spout. The number of punished drinks was counted automatically by the computer interface. The number of punished drinks was compared between treatment groups. An increase in the number of punished drinks in rats treated with a compound is interpreted as an anxiolytic-like effect.

[0167] Compound administration: Test compounds were suspended in 1% CMC. Test compounds were administered by oral gavage (p.o.) in a volume of 3 ml/kg 1 or 2 hours before testing. In each assay, a vehicle and reference positive control groups were tested.

[0168] Statistical analyses: Statistical analyses were performed using GraphPad PRISM version 4.01 statistical software (GraphPad, San Diego, CA, USA). Data were analyzed using one-way analysis of variance (ANOVA) followed by Bonferroni's multiple comparisons. The significance level was set at $p < 0.05$ when compared to the vehicle group.

[0169] Effect of compounds on the Vogel conflict test in rats: At the doses of about 0.3 to 3 mg/kg, selected compounds of the invention increase the number of punished licks in the Vogel conflict test in rats.

EXAMPLE D

Elevated plus maze test, model of anxiety in mice or rats

[0170] Anxiety models in rodents are used as standard tests to demonstrate anxiolytic-like properties of novel compounds. The elevated plus maze is an animal model of anxiety, which has been validated behaviourally, physiologically, and pharmacologically (Pellow et al., J. Neurosci. Methods 1985, 14: 149-167). Following placement on the maze, untreated animals (mice or rats) tend to avoid the open arms and spend most of their time in the closed arms. Anxiolytic drugs such as benzodiazepines increase open arm exploration. Anxiety in the elevated plus maze is measured by examining the time spent on the open arms as well as open arm entries. In addition, closed arm entries are measured and used as a general activity score.

[0171] Subjects: The present studies were performed in accordance with the animal care and use policies of Addex Pharmaceuticals and the EEC directives on the protection of animals used for experimental and other scientific purposes (86/609/EEC and subsequent revisions). Male Sprague-Dawley rats (200 - 300 g) or male C57/B16J mice (20-30g) were housed in a temperature and humidity controlled facility on a 12 hours light /dark cycle for at least 5 days before use. Animals had access to food *ad libitum* excepted during elevated plus maze experiments.

[0172] Assessment of the elevated plus-maze test: The elevated plus-maze consists of four plastic arms connected by a central square, the arms are all horizontal and at 90° angle from each other making the shape of a plus sign. The maze is raised to a height of 50 cm from the ground. Two of the opposite arms have high plastic walls whereas the other two opposite arms do not. These arms are therefore referred to as the enclosed and open arms, respectively. The central square is not enclosed and the animal is initially placed in this area facing an enclosed arm.

The animals had free access to all 4 arms for a 5 minute period. During this time, the number of entries into, and the time spent on each type of arm were recorded.

[0173] The results were presented as:

- Percentage of total number of entries onto open arms.
- Percentage of time spent on the open arms.
- Total number of entries into all arms.

[0174] On the day of the test, animals were marked on their tails and weighed in a separate preparation room before drug administration. Test compound or vehicle was administered p.o. 60 minutes prior to the test session.

[0175] Compound administration: Test compounds were dissolved in a solution of 1% CMC. Test compounds were administered 60 minutes before testing by oral gavage (p.o.) in a volume of 10 mL/kg. In each assay, a vehicle and reference positive control groups were tested.

[0176] Statistical analyses: Statistical analyses were performed using GraphPad PRISM version 4.01 statistical software (GraphPad, San Diego, CA, USA). Data were analyzed using one-way analysis of variance (ANOVA) followed by Bonferroni's multiple comparisons test. The significance level was set at $p < 0.05$ when compared to the vehicle group.

[0177] Effect of compounds in the elevated plus maze test: At the doses of about 0.3 to 3 mg/kg, selected compounds of the invention increase both the number of entries and the time spent on the open arms.

[0178] Thus, the positive allosteric modulators provided in the present invention are expected to increase the effectiveness of GABA or mGABA_B agonists at GABA_B receptor. Therefore, these positive allosteric modulators are expected to be useful for treating, or preventing, ameliorating, controlling or reducing the risk of various central nervous system disorders as well as other disorders associated with GABA_B dysfunction in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of GABA_B positive allosteric modulators.

[0179] In particular, the neurological and psychiatric disorders associated with GABA_B receptor dysfunction, include one or more of the following conditions or diseases: anxiety, depression, epilepsy, schizophrenia, cognitive disorders, spasticity and skeletal muscle rigidity, spinal cord injury, multiple sclerosis, neuropathic pain and craving associated with cocaine and nicotine, amyotrophic lateral sclerosis, cerebral palsy, posttraumatic stress disorders or gastro-intestinal disorders, cerebral palsy, psychosis, panic disorder

[0180] 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 medical and scientific progress.

[0181] 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 the invention or the other drugs may have utility, where the combination of the drugs together are safer or more effective than either drug alone.

FORMULATION EXAMPLES

[0182] Typical examples of recipes for the formulation of the compounds of the invention are as follows:

1) Tablets

Compound 4.02	5 to 50 mg
Di-calcium phosphate	20 mg
Lactose	30 mg
Talcum	10 mg
Magnesium stearate	5 mg
Potato starch	ad 200 mg

In this example, the Compound **4.02** can be replaced by the same amount of any of the described examples.

2) Suspension

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

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 and water.

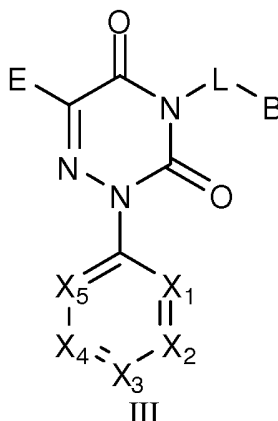
4) Ointment

Compound 4.02	5 to 1000 mg
Stearyl alcohol	3 g
Lanoline	5 g
White petroleum	15 g
Water	ad 100 g

[0183] In this example, the compound **4.02** can be replaced by the same amount of any of the described examples.

Claims

1. A compound having the Formula III



and stereoisomeric forms, mixtures of stereoisomeric forms or pharmaceutically acceptable salts, hydrates or solvates of such compounds, wherein:

B is selected from C₆-C₁₀ aryl optionally substituted with one to 5 Y, C₅-C₁₀ cycloalkenyl optionally substituted with one to 7 Y, 5 to 14 membered heteroaryl group optionally substituted by one to 8 Y, wherein said heteroaryl group comprises one, two or three heteroatoms selected from N, O, S;

Y is 1 to 5 substituents independently selected from the group consisting of hydrogen, halogen, -OCF₃, -NO₂, -CN, -CF₃, an optionally substituted-(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR¹, -(C₀-C₆)alkylNR²R³, -(C₀-C₆)alkyl-C(=NR⁴)NR²R³, -(C₀-C₆)alkylSR², -(C₀-C₆)alkylNR²C(=O)R³, -(C₀-C₆)alkylNR²S(=O)₂R³, -(C₀-C₆)alkylNR⁴C(=O)NR²R³, -(C₀-C₆)alkylS(=O)R², -(C₀-C₆)alkylS(=O)₂R², -(C₀-C₆)alkylS(=O)₂NR²R³, -(C₀-C₆)alkylC(=O)R², -(C₀-C₆)alkylC(=O)OR¹, -(C₀-C₆)alkylC(=O)NR²R³, -(C₀-C₆)alkylC(=NR²)R³, or -(C₀-C₆)alkylC(=NOR¹)R³, heteroaryl, heteroarylalkyl, arylalkyl, aryl alkylaryl, alkylheteroaryl; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)alkylaryl, -O(C₁-C₃)alkylheteroaryl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₀-C₃)alkylaryl) or -N((C₀-C₆)alkyl)((C₀-C₃)alkylheteroaryl) groups; R¹ at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl, 3 to 7 membered heterocycloalkyl, -(C₆-C₁₀)aryl, 5 or 6 membered heteroaryl, -(C₇-C₁₀)arylalkyl, -C(=O)R⁵; R⁵ are selected from hydrogen, -(C₁-C₆)alkyl, and -(C₆-C₁₀)aryl;

R², R³ and R⁴ each independently is hydrogen, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, -(C₁-C₆)alkylhalo, heterocycloalkyl, heteroaryl, heteroarylalkyl, arylalkyl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkylalkyl, -O(aryl), -O(heteroaryl), -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₃-C₇)cycloalkyl) or -N((C₀-C₆)alkyl)(aryl) substituents; wherein optionally R², R³, R⁴ substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl;

X₁, X₂, X₃, X₄ and X₅ are each independently selected from the group consisting of -CR⁶=, -N=;

R⁶ is 1 to 5 substituents independently selected from the group consisting of hydrogen, halogen, -OCF₃, -CN, -CF₃, -NO₂, an optionally substituted-(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR⁷, -(C₀-C₆)alkylNR⁸R⁹, -(C₀-C₆)alkyl-C(=NR¹⁰)NR⁸R⁹, -(C₀-C₆)alkylSR⁸, -(C₀-C₆)alkylNR⁸C(=O)R⁹, -(C₀-C₆)alkylNR⁸S(=O)₂R⁹, -(C₀-C₆)alkylNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)alkylS(=O)R⁸, -(C₀-C₆)alkylS(=O)₂R⁸, -(C₀-C₆)alkylS(=O)₂NR⁸R⁹, -(C₀-C₆)alkylC(=O)R⁸, -(C₀-C₆)alkylC(=O)OR⁷, -(C₁-C₆)alkylC(=O)NR⁸R⁹, or -(C₀-C₆)alkylC(=NOR⁷)R⁹, heteroaryl, heteroarylalkyl, arylalkyl, aryl, a 3 to 6 heterocycloalkyl; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)alkylaryl, -O(C₁-C₃)alkylheteroaryl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₀-C₃)alkylaryl) or -N((C₀-C₆)alkyl)((C₀-C₃)alkylheteroaryl) groups;

R⁷ at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl, 3 to 7 membered heterocycloalkyl, -(C₆-C₁₀)aryl, 5 or 6 membered heteroaryl, -(C₇-C₁₀)arylalkyl, -C(=O)R¹¹; R¹¹ are selected from -(C₁-C₆)alkyl, -(C₃-C₇)cycloalkyl and -(C₆-C₁₀)aryl;

R⁸, R⁹ and R¹⁰ each independently is hydrogen, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, -(C₁-C₆)alkylhalo, heterocycloalkyl, heteroaryl, heteroarylalkyl, arylalkyl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkylalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₃-C₇)cycloalkyl) or -N((C₀-C₆)alkyl)(aryl) substituents;

E is selected from the group consisting of hydrogen, halogen, -OCF₃, -NO₂, -CF₃, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR¹², -(C₀-C₆)alkylNR¹³R¹⁴, -(C₀-C₆)alkyl-C(=NR¹⁵)NR¹³R¹⁴, -(C₀-C₆)alkylSR¹², -(C₀-C₆)alkylNR¹³C(=O)R¹⁴, -(C₀-C₆)alkylNR¹³S(=O)₂R¹⁴, -(C₀-C₆)alkylNR¹⁵C(=O)NR¹³R¹⁴, -(C₀-C₆)alkylS(=O)R¹³, -(C₀-C₆)alkylS(=O)₂R¹³, -(C₀-C₆)alkylS(=O)₂NR¹³R¹⁴, -(C₀-C₆)alkylC(=O)R¹³, -(C₀-C₆)alkylC(=O)OR¹³, -(C₀-C₆)alkylC(=O)NR¹³R¹⁴, -(C₀-C₆)alkylC(=NR¹³)R¹⁴, or -(C₀-C₆)alkylC(=NOR¹³)R¹⁴; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, -CN, an optionally substituted -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)alkylaryl, -O(C₁-C₃)alkylheteroaryl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₀-C₃)alkylaryl) or -N((C₀-C₆)alkyl)((C₀-C₃)alkylheteroaryl) groups;

R¹² at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl, -C(=O)R¹⁶; R¹⁶ are selected from hydrogen, -(C₁-C₆)alkyl, -(C₃-C₇)cycloalkyl and -(C₆-C₁₀)aryl;

R¹³, R¹⁴ and R¹⁵ each independently is hydrogen, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, -(C₁-C₆)alkylhalo, heterocycloalkyl, heteroaryl, heteroarylalkyl, alkylheteroaryl, arylalkyl, alkylaryl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkylalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₃-C₇)cycloalkyl) or -N((C₀-C₆)alkyl)(aryl) substituents;

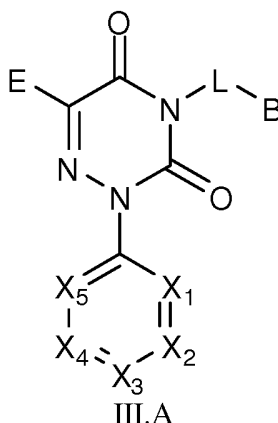
When E is -H the compounds for which X₃ is not CH= or -CF= are excluded from the invention;

L is independently selected from a -((C₁-C₅)alkyl-Q_m)- substituted with 1-6 T; m is 0 or 1;

T is 1 to 6 substituents independently selected from the group consisting of hydrogen, fluorine, -OCF₃, -NO₂, -CN, -CF₃, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR¹⁷, -(C₀-C₆)alkylNR¹⁸R¹⁹, -(C₀-C₆)alkyl-C(=NR²⁰)NR¹⁸R¹⁹, -(C₀-C₆)alkylSR¹⁸, -(C₀-C₆)alkylNR¹⁸S(=O)R¹⁹, -(C₀-C₆)alkylNR¹⁸C(=O)₂R¹⁹, -(C₀-C₆)alkylNR¹⁸S(=O)₂R¹⁹, -(C₀-C₆)alkylNR²⁰C(=O)NR¹⁸R¹⁹, -(C₀-C₆)alkylS(=O)R¹⁸, -(C₀-C₆)alkylS(=O)₂R¹⁸, -(C₀-C₆)alkylS(=O)₂NR¹⁸R¹⁹, -(C₀-C₆)alkylC(=O)R¹⁸, -(C₀-C₆)alkylC(=O)OR¹⁷, -(C₀-C₆)alkylC(=O)NR¹⁸R¹⁹, -(C₀-C₆)alkylC(=NR¹⁸)R¹⁹, or -(C₀-C₆)alkylC(=NOR¹⁷)R¹⁹, heteroaryl, heteroarylalkyl, arylalkyl, alkylaryl, alkylheteroaryl, aryl; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)alkylaryl, -O(C₁-C₃)alkylheteroaryl, -N((C₀-C₆)alkyl)((C₀-C₃)alkylaryl) or -N((C₀-C₆)alkyl)((C₀-C₃)alkylheteroaryl) groups; R¹⁷ at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl, 3 to 7 membered heterocycloalkyl, -(C₆-C₁₀)aryl, 5 or 6 membered heteroaryl, -(C₇-C₁₀)arylalkyl, -C(=O)R²¹; R²¹ is selected from hydrogen, -(C₁-C₆)alkyl, -(C₃-C₇)cycloalkyl, and -(C₆-C₁₀)aryl; R¹⁸, R¹⁹ and R²⁰ each independently is hydrogen, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, -(C₁-C₆)alkylhalo, heterocycloalkyl, heteroaryl, heteroarylalkyl, arylalkyl, alkylaryl, alkylheteroaryl, aryl; any of which is optionally substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkylalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₃-C₇)cycloalkyl) or -N((C₀-C₆)alkyl)(aryl) substituents;

Q is -C(=O)-, -C(=O)O-, -C(=O)NR²²-, -OC(=O)-, -OC(=O)NR²²-, -NR²²-, -NR²²C(=O)-, -NR²²C(=O)₂-, -NR²²C(=S)-, -S-, -S(=O)-, or -S(=O)₂-; R²² is selected from -(C₁-C₆)alkyl, and -(C₆-C₁₀)aryl; Any N may be an N-oxide.

2. A compound according to claim 1 having the Formula III.A



and stereoisomeric forms, mixtures of stereoisomeric forms or pharmaceutically acceptable salts, hydrates or solvates of such compounds, wherein:

B is selected from C₆-C₁₀ aryl optionally substituted with one to 5 Y, C₅-C₁₀ cycloalkenyl optionally substituted with one to 7 Y, 5 to 14 membered heteroaryl group optionally substituted by one to 8 Y, wherein said heteroaryl group comprises one, two or three heteroatoms selected from N, O, S;

Y is 1 to 5 substituents independently selected from the group consisting of hydrogen, halogen, -OCF₃, -NO₂, -CN, -CF₃, an optionally substituted-(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR¹, -(C₀-C₆)alkylNR²R³, -(C₀-C₆)alkyl-C(=NR⁴)NR²R³, -(C₀-C₆)alkylSR²,

-(C₀-C₆)alkylNR²C(=O)R³, -(C₀-C₆)alkylNR²S(=O)₂R³, -(C₀-C₆)alkylNR⁴C(=O)NR²R³, -(C₀-C₆)alkylS(=O)R², -(C₀-C₆)alkylS(=O)₂R², -(C₀-C₆)alkylS(=O)₂NR²R³, (C₀-C₆)alkylC(=O)R², -(C₀-C₆)alkylC(=O)OR¹, -(C₀-C₆)alkylC(=O)NR²R³, -(C₀-C₆)alkylC(=NR²)R³, or -(C₀-C₆)alkylC(=NOR¹)R³, heteroaryl, heteroarylalkyl, arylalkyl, aryl alkylaryl, alkylheteroaryl; wherein optionally two substituents are combined to the intervening

atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)alkylaryl, -O(C₁-C₃)alkylheteroaryl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₀-C₃)alkylaryl) or -N((C₀-C₆)alkyl)((C₀-C₃)alkylheteroaryl) groups;

R¹ at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl, 3 to 7 membered heterocycloalkyl, -(C₆-C₁₀)aryl, 5 or 6 membered heteroaryl, -(C₇-C₁₀)arylalkyl, -C(=O)R⁵; R⁵ are selected from hydrogen, -(C₁-C₆)alkyl, and -(C₆-C₁₀)aryl;

R², R³ and R⁴ each independently is hydrogen, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, -(C₁-C₆)alkylhalo, heterocycloalkyl, heteroaryl, heteroarylalkyl, arylalkyl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkylalkyl, -O(aryl), -O(heteroaryl), -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₃-C₇)cycloalkyl) or -N((C₀-C₆)alkyl)(aryl) substituents; wherein optionally R², R³, R⁴ substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl;

X₁, X₂, X₃, X₄ and X₅ are each independently selected from the group consisting of -CR⁶=, -N=;

R⁶ is 1 to 5 substituents independently selected from the group consisting of hydrogen, halogen, -OCF₃, -CN, -CF₃, -NO₂, an optionally substituted-(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR⁷, -(C₀-C₆)alkylNR⁸R⁹, -(C₀-C₆)alkyl-C(=NR¹⁰)NR⁸R⁹, -(C₀-C₆)alkylSR⁸, -(C₀-C₆)alkylNR⁸C(=O)R⁹, -(C₀-C₆)alkylNR⁸S(=O)₂R⁹, -(C₀-C₆)alkylNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)alkylS(=O)R⁸, -(C₀-C₆)alkylS(=O)₂R⁸, -(C₀-C₆)alkylS(=O)₂NR⁸R⁹, -(C₀-C₆)alkylC(=O)R⁸, -(C₀-C₆)alkylC(=O)OR⁷, -(C₁-C₆)alkylC(=O)NR⁸R⁹, or -(C₀-C₆)alkylC(=NOR⁷)R⁹, heteroaryl, heteroarylalkyl, arylalkyl, aryl, a 3 to 6 heterocycloalkyl; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)alkylaryl, -O(C₁-C₃)alkylheteroaryl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₀-C₃)alkylaryl) or -N((C₀-C₆)alkyl)((C₀-C₃)alkylheteroaryl) groups;

R⁷ at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl, 3 to 7 membered heterocycloalkyl, -(C₆-C₁₀)aryl, 5 or 6 membered heteroaryl, -(C₇-C₁₀)arylalkyl, -C(=O)R¹¹; R¹¹ are selected from -(C₁-C₆)alkyl, -(C₃-C₇)cycloalkyl and -(C₆-C₁₀)aryl;

R⁸, R⁹ and R¹⁰ each independently is hydrogen, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alky-

nyl, $-(C_1-C_6)alkylhalo$, heterocycloalkyl, heteroaryl, heteroarylalkyl, arylalkyl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, $-CN$, $-(C_1-C_6)alkyl$, $-O(C_0-C_6)alkyl$, $-O(C_3-C_7)cycloalkylalkyl$, $-O(aryl)$, $-O(aryl)-(C_0-C_3)alkyl$, $-O(heteroaryl)$, $-O(heteroaryl)-(C_0-C_3)alkyl$, $-N((C_0-C_6)alkyl)_2$, $-N((C_0-C_6)alkyl)((C_3-C_7)cycloalkyl)$ or $-N((C_0-C_6)alkyl)(aryl)$ substituents;

E is selected from the group consisting of hydrogen, halogen, $-OCF_3$, $-NO_2$, $-CF_3$, $-(C_1-C_6)alkyl$, $-(C_3-C_6)cycloalkyl$, $halo-(C_1-C_6)alkyl$, $-(C_0-C_6)alkylOR^{12}$, $-(C_0-C_6)alkylINR^{13}R^{14}$, $-(C_0-C_6)alkylISR^{12}$, $-(C_0-C_6)alkylINR^{13}C(=O)R^{14}$, $-(C_0-C_6)alkylINR^{13}S(=O)_2R^{14}$, $-(C_0-C_6)alkylINR^{15}C(=O)NR^{13}R^{14}$, $-(C_0-C_6)alkylS(=O)R^{13}$, $-(C_0-C_6)alkylS(=O)_2R^{13}$, $-(C_0-C_6)alkylS(=O)NR^{13}R^{14}$, $-(C_0-C_6)alkylC(=O)R^{13}$, $-(C_0-C_6)alkylC(=O)OR^{13}$, $-(C_0-C_6)alkylC(=O)NR^{13}R^{14}$; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, $-CN$, an optionally substituted $-(C_1-C_6)alkyl$, $-O(C_0-C_6)alkyl$, $-O(C_3-C_7)cycloalkyl$, $-O(aryl)$, $-O(aryl)-(C_0-C_3)alkyl$, $-O(heteroaryl)$, $-O(heteroaryl)-(C_0-C_3)alkyl$, $-O(C_1-C_3)alkylaryl$, $-O(C_1-C_3)alkylheteroaryl$, $-N((C_0-C_6)alkyl)_2$, $-N((C_0-C_6)alkyl)((C_0-C_3)alkylaryl)$ or $-N((C_0-C_6)alkyl)((C_0-C_3)alkylheteroaryl)$ groups;

R^{12} at each occurrence is selected from hydrogen, $-(C_1-C_6)alkyl$, $-(C_2-C_6)alkenyl$, $-(C_3-C_7)cycloalkyl$, $-C(=O)R^{16}$; R^{16} are selected from hydrogen, $-(C_1-C_6)alkyl$, $-(C_3-C_7)cycloalkyl$ and $-(C_6-C_{10})aryl$;

R^{13} , R^{14} and R^{15} each independently is hydrogen, $-(C_1-C_6)alkyl$, $-(C_3-C_6)cycloalkyl$, $-(C_2-C_6)alkenyl$, $-(C_2-C_6)alkynyl$, $-(C_1-C_6)alkylhalo$, heterocycloalkyl, heteroaryl, heteroarylalkyl, alkylheteroaryl, arylalkyl, alkylaryl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, $-CN$, $-(C_1-C_6)alkyl$, $-O(C_0-C_6)alkyl$, $-O(C_3-C_7)cycloalkylalkyl$, $-O(aryl)$, $-O(aryl)-(C_0-C_3)alkyl$, $-O(heteroaryl)$, $-O(heteroaryl)-(C_0-C_3)alkyl$, $-N((C_0-C_6)alkyl)_2$, $-N((C_0-C_6)alkyl)((C_3-C_7)cycloalkyl)$ or $-N((C_0-C_6)alkyl)(aryl)$ substituents;

When E is $-H$ the compounds for which X_3 is not $CH=$ or $-CF=$ are excluded from the invention;

L is independently selected from a $-(C_1-C_5)alkyl-Q_m$ - substituted with 1-6 T; m is 0 or 1;

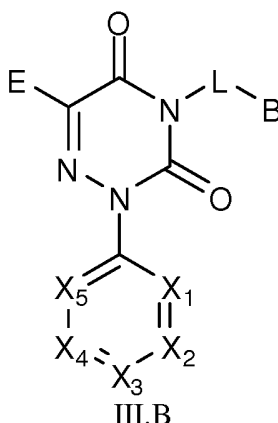
T is 1 to 6 substituents independently selected from the group consisting of hydrogen, fluorine, $-OCF_3$, $-NO_2$, $-CN$, $-CF_3$, $-(C_1-C_6)alkyl$, $-(C_3-C_6)cycloalkyl$, $-(C_2-C_6)alkenyl$, $-(C_2-C_6)alkynyl$, $halo-(C_1-C_6)alkyl$, $-(C_0-C_6)alkylOR^{17}$, $-(C_0-C_6)alkylINR^{18}R^{19}$, $-(C_0-C_6)alkylI-C(=NR^{20})NR^{18}R^{19}$, $-(C_0-C_6)alkylISR^{18}$, $-(C_0-C_6)alkylINR^{18}C(=O)R^{19}$, $-(C_0-C_6)alkylINR^{18}C(=O)_2R^{19}$, $-(C_0-C_6)alkylINR^{18}S(=O)_2R^{19}$, $-(C_0-C_6)alkylINR^{20}C(=O)NR^{18}R^{19}$, $-(C_0-C_6)alkylS(=O)R^{18}$, $-(C_0-C_6)alkylS(=O)_2R^{18}$, $-(C_0-C_6)alkylS(=O)_2NR^{18}R^{19}$, $-(C_0-C_6)alkylC(=O)R^{18}$, $-(C_0-C_6)alkylC(=O)OR^{17}$, $-(C_0-C_6)alkylC(=O)NR^{18}R^{19}$, $-(C_0-C_6)alkylC(=NR^{18})R^{19}$, or $-(C_0-C_6)alkylC(=NOR^{17})R^{19}$, heteroaryl, heteroarylalkyl, arylalkyl, alkylaryl, alkylheteroaryl, aryl; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, $-CN$, $-(C_1-C_6)alkyl$, $-O(C_0-C_6)alkyl$, $-O(C_3-C_7)cycloalkyl$, $-O(aryl)$, $-O(aryl)-(C_0-C_3)alkyl$, $-O(heteroaryl)$, $-O(heteroaryl)-(C_0-C_3)alkyl$, $-O(C_1-C_3)alkylaryl$, $-O(C_1-C_3)alkylheteroaryl$, $-N((C_0-C_6)alkyl)_2$, $-N((C_0-C_6)alkyl)((C_0-C_3)alkylaryl)$ or $-N((C_0-C_6)alkyl)((C_0-C_3)alkylheteroaryl)$ groups;

R^{17} at each occurrence is selected from hydrogen, $-(C_1-C_6)alkyl$, $-(C_2-C_6)alkenyl$, $-(C_3-C_7)cycloalkyl$, 3 to 7 membered heterocycloalkyl, $-(C_6-C_{10})aryl$, 5 or 6 membered heteroaryl, $-(C_7-C_{10})arylalkyl$, $-C(=O)R^{21}$; R^{21} is selected from hydrogen, $-(C_1-C_6)alkyl$, $-(C_3-C_7)cycloalkyl$, and $-(C_6-C_{10})aryl$;

R^{18} , R^{19} and R^{20} each independently is hydrogen, $-(C_1-C_6)alkyl$, $-(C_3-C_6)cycloalkyl$, $-(C_2-C_6)alkenyl$, $-(C_2-C_6)alkynyl$, $-(C_1-C_6)alkylhalo$, heterocycloalkyl, heteroaryl, heteroarylalkyl, arylalkyl, alkylaryl, alkylheteroaryl, aryl; any of which is optionally substituted with 1 to 5 independent halogen, $-CN$, $-(C_1-C_6)alkyl$, $-O(C_0-C_6)alkyl$, $-O(C_3-C_7)cycloalkylalkyl$, $-O(aryl)$, $-O(aryl)-(C_0-C_3)alkyl$, $-O(heteroaryl)$, $-O(heteroaryl)-(C_0-C_3)alkyl$, $-N((C_0-C_6)alkyl)_2$, $-N((C_0-C_6)alkyl)((C_3-C_7)cycloalkyl)$ or $-N((C_0-C_6)alkyl)(aryl)$ substituents; Q is $-C(=O)-$, $-C(=O)O-$, $-C(=O)NR^{22}$, $-OC(=O)-$, $-OC(=O)NR^{22}$, $-NR^{22}$, $-NR^{22}C(=O)-$, $-NR^{22}C(=O)_2-$, $-NR^{22}C(=S)-$, $-S-$, $-S(=O)-$, or $-S(=O)_2-$; R^{22} is selected from $-(C_1-C_6)alkyl$, and $-(C_6-C_{10})aryl$;

Any N may be an N-oxide.

3. A compound according to 2 having the Formula III.B



and stereoisomeric forms, mixtures of stereoisomeric forms or pharmaceutically acceptable salts, hydrates or solvates of such compounds, wherein:

B is selected from C₆-C₁₀ aryl optionally substituted with one to 5 Y, C₅-C₁₀ cycloalkenyl optionally substituted with one to 7 Y, 5 to 14 membered heteroaryl group optionally substituted by one to 8 Y, wherein said heteroaryl group comprises one, two or three heteroatoms selected from N, O, S;

Y is 1 to 5 substituents independently selected from the group consisting of hydrogen, halogen, -OCF₃, -NO₂, -CN, -CF₃, an optionally substituted-(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR¹, -(C₀-C₆)alkylNR²R³, -(C₀-C₆)alkyl-C(=NR⁴)NR²R³, -(C₀-C₆)alkylSR², -(C₀-C₆)alkylNR²C(=O)R³, -(C₀-C₆)alkylNR²S(=O)₂R³, -(C₀-C₆)alkylNR⁴C(=O)NR²R³, -(C₀-C₆)alkylS(=O)R², -(C₀-C₆)alkylS(=O)₂R², -(C₀-C₆)alkylS(=O)₂NR²R³, (C₀-C₆)alkylC(=O)R², -(C₀-C₆)alkylC(=O)OR¹, -(C₀-C₆)alkylC(=O)NR²R³, -(C₀-C₆)alkylC(=NR²)R³, or -(C₀-C₆)alkylC(=NOR¹)R³, heteroaryl, heteroarylalkyl, arylalkyl, aryl alkylaryl, alkylheteroaryl; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl,

aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)alkylaryl, -O(C₁-C₃)alkylheteroaryl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₀-C₃)alkylaryl) or -N((C₀-C₆)alkyl)((C₀-C₃)alkylheteroaryl) groups;

R¹ at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl, 3 to 7 membered heterocycloalkyl, -(C₆-C₁₀)aryl, 5 or 6 membered heteroaryl, -(C₇-C₁₀)arylalkyl, -C(=O)R⁵; R⁵ are selected from hydrogen, -(C₁-C₆)alkyl, and -(C₆-C₁₀)aryl;

R², R³ and R⁴ each independently is hydrogen, -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, -(C₁-C₆)alkylhalo, heterocycloalkyl, heteroaryl, heteroarylalkyl, arylalkyl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkylalkyl, -O(aryl), -O(heteroaryl), -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₃-C₇)cycloalkyl) or -N((C₀-C₆)alkyl)(aryl) substituents; wherein optionally R², R³, R⁴ substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl;

X₁, X₂, X₃, X₄ and X₅ are each independently selected from the group consisting of -CR⁶=, -N=;

R⁶ is 1 to 5 substituents independently selected from the group consisting of hydrogen, halogen, -OCF₃, -CN, -CF₃, -NO₂, an optionally substituted-(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₂-C₆)alkenyl, -(C₂-C₆)alkynyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR⁷, -(C₀-C₆)alkylNR⁸R⁹, -(C₀-C₆)alkylSR⁸, -(C₀-C₆)alkylNR⁸C(=O)R⁹, -(C₀-C₆)alkylNR⁸S(=O)₂R⁹, -(C₀-C₆)alkylNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)alkylS(=O)R⁸, -(C₀-C₆)alkylS(=O)₂R⁸, -(C₀-C₆)alkylS(=O)₂NR⁸R⁹, -(C₀-C₆)alkylC(=O)R⁸, -(C₀-C₆)alkylC(=O)OR⁷, -(C₁-C₆)alkylC(=O)NR⁸R⁹, heteroaryl, heteroarylalkyl, arylalkyl, aryl, a 3 to 6 heterocycloalkyl; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, -CN, -(C₁-C₆)alkyl, -O(C₀-C₆)alkyl, -O(C₃-C₇)cycloalkyl, -O(aryl), -O(aryl)-(C₀-C₃)alkyl, -O(heteroaryl), -O(heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)alkylaryl, -O(C₁-C₃)alkylheteroaryl, -N((C₀-C₆)alkyl)₂, -N((C₀-C₆)alkyl)((C₀-C₃)alkylaryl) or -N((C₀-C₆)alkyl)((C₀-C₃)alkylheteroaryl) groups;

R⁷ at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl, 3 to 7 membered heterocycloalkyl, -(C₆-C₁₀)aryl, 5 or 6 membered heteroaryl, -(C₇-C₁₀)arylalkyl, -C(=O)R¹¹; R¹¹ are

lected from $-(C_1-C_6)alkyl$, $-(C_3-C_7)cycloalkyl$ and $-(C_6-C_{10})aryl$;

R^8 , R^9 and R^{10} each independently is hydrogen, $-(C_1-C_6)alkyl$, $-(C_3-C_6)cycloalkyl$, $-(C_2-C_6)alkenyl$, $-(C_2-C_6)alkynyl$, $-(C_1-C_6)alkylhalo$, heterocycloalkyl, heteroaryl, heteroarylalkyl, arylalkyl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, $-CN$, $-(C_1-C_6)alkyl$, $-O(C_0-C_6)alkyl$, $-O(C_3-C_7)cycloalkylalkyl$, $-O(aryl)$, $-O(aryl)-(C_0-C_3)alkyl$, $-O(heteroaryl)$, $-O(heteroaryl)-(C_0-C_3)alkyl$, $-N((C_0-C_6)alkyl)_2$, $-N((C_0-C_6)alkyl)((C_3-C_7)cycloalkyl)$ or $-N((C_0-C_6)alkyl)(aryl)$ substituents;

E is selected from the group consisting of hydrogen, halogen, $-OCF_3$, $-CF_3$, $-(C_1-C_6)alkyl$, $-(C_3-C_6)cycloalkyl$, $-(C_0-C_6)alkylOR^{12}$, $-(C_0-C_6)alkylNR^{13}R^{14}$, $-(C_0-C_6)alkylS(=O)R^{13}$, $-(C_0-C_6)alkylS(=O)_2R^{13}$, $-(C_0-C_6)alkylC(=O)R^{13}$, $-(C_0-C_6)alkylC(=O)OR^{13}$, $-(C_0-C_6)alkylC(=O)NR^{13}R^{14}$; wherein optionally two substituents are combined to the intervening atoms to form a bicyclic heterocycloalkyl, aryl or heteroaryl ring; wherein each ring is optionally further substituted with 1 to 5 independent halogen, $-CN$, an optionally substituted $-(C_1-C_6)alkyl$, $-O(C_0-C_6)alkyl$, $-O(C_3-C_7)cycloalkyl$, $-O(aryl)$, $-O(aryl)-(C_0-C_3)alkyl$, $-O(heteroaryl)$, $-O(heteroaryl)-(C_0-C_3)alkyl$, $-O(C_1-C_3)alkylaryl$, $-O(C_1-C_3)alkylheteroaryl$, $-N((C_0-C_6)alkyl)_2$, $-N((C_0-C_6)alkyl)((C_0-C_3)alkylaryl)$ or $-N((C_0-C_6)alkyl)((C_0-C_3)alkylheteroaryl)$ groups;

R^{12} at each occurrence is selected from hydrogen, $-(C_1-C_6)alkyl$, $-(C_2-C_6)alkenyl$, $-(C_3-C_7)cycloalkyl$, $-C(=O)R^{16}$; R^{16} are selected from hydrogen, $-(C_1-C_6)alkyl$, $-(C_3-C_7)cycloalkyl$ and $-(C_6-C_{10})aryl$;

R^{13} and R^{14} each independently is hydrogen, $-(C_1-C_6)alkyl$, $-(C_3-C_6)cycloalkyl$, $-(C_2-C_6)alkenyl$, $-(C_2-C_6)alkynyl$, $-(C_1-C_6)alkylhalo$, heterocycloalkyl, heteroaryl, heteroarylalkyl, alkylheteroaryl, arylalkyl, alkylaryl or aryl; any of which is optionally substituted with 1 to 5 independent halogen, $-CN$, $-(C_1-C_6)alkyl$, $-O(C_0-C_6)alkyl$, $-O(C_3-C_7)cycloalkylalkyl$, $-O(aryl)$, $-O(aryl)-(C_0-C_3)alkyl$, $-O(heteroaryl)$, $-O(heteroaryl)-(C_0-C_3)alkyl$, $-N((C_0-C_6)alkyl)_2$, $-N((C_0-C_6)alkyl)((C_3-C_7)cycloalkyl)$ or $-N((C_0-C_6)alkyl)(aryl)$ substituents;

When E is $-H$ the compounds for which X_3 is not $CH=$ or $-CF=$ are excluded from the invention;

L is independently selected from a $-(C_1-C_5)alkyl-Q_m$ - substituted with 1-6 T; m is 0 or 1;

T is 1 to 6 substituents independently selected from the group consisting of hydrogen, $-(C_1-C_6)alkyl$;

Q is $-C(=O)-$, $-C(=O)O-$, $-C(=O)NR^{22}-$, $-OC(=O)-$, $-OC(=O)NR^{22}-$, $-NR^{22}-$, $-NR^{22}C(=O)-$, $-NR^{22}C(=O)_2-$, $-NR^{22}C(=S)-$, $-S-$, $-S(=O)-$, or $-S(=O)_2-$; R^{22} is selected from $-(C_1-C_6)alkyl$, and $-(C_6-C_{10})aryl$; Any N may be an N-oxide.

4. A compound according to claims 1 to 3, which can exist as optical isomers, wherein said compound is either the racemic mixture or an individual optical isomer.

5. A compound according to claims 1 to 4, wherein said compound is selected from:

N-(3-(4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)methanesulfonamide, 4-(4-chlorobenzyl)-2-(2-methoxyphenyl)-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-cyanobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(naphthalen-2-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(benzo[c][1,2,5]oxadiazol-5-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-benzyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(3,5-dioxo-4-(4-(trifluoromethyl)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, methyl 4-((2-(3-acetamidophenyl)-3,5-dioxo-2,3-dihydro-1,2,4-triazin-4(5H)-yl)methyl)benzoate, N-(3-(4-(naphthalen-1-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-bromobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(3-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-isopropylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(3,4-dimethylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(6-chloropyridin-3-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chloro-2-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-benzyl-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chloro-3-fluorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(6-fluoropyridin-3-yl)methyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(6-isopropylpyridin-3-yl)methyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-

fluorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(2-fluoro-5-(4-(4-fluorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-methoxybenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-isopropylbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-(4-chlorobenzyl)-6-methyl-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(6-methyl-4-(naphthalen-2-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((6-cyclopentylpyridin-3-yl)methyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(3,4-difluorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(3,4-dimethoxybenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(2,4-difluorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-isopropoxybenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methyl-4-(naphthalen-1-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methyl-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(2-fluoro-5-(6-methyl-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(2-fluoro-5-(4-(3-methoxybenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 3-(4-(4-fluorobenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-N-methylbenzamide, N-(3-(4-(4-fluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-3,5-dioxo-4-(4-(trifluoromethyl)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chloro-3-fluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-(4-chloro-3-fluorobenzyl)-6-methoxy-2-(2-methoxyphenyl)-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)methanesulfonamide, 4-(4-chlorobenzyl)-6-methoxy-2-(2-methoxyphenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 1-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-3-methylurea, N-(3-(4-((6-chloropyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(2-fluoro-5-(6-methoxy-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)pyridin-3-yl)acetamide, N-(3-(6-methoxy-4-((6-methylpyridin-3-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-(4-chloro-3-fluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)pyridin-3-yl)acetamide, 4-(4-chlorobenzyl)-2-(4-fluoro-2-methoxyphenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(2-(methoxymethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(3-(methoxymethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(3-(2-methoxyethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(2-(trifluoromethoxy)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(2-(morpholinomethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(2-(2-methoxyethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(3-((2-methyl-1,3-dioxolan-2-yl)methyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(3-(2-oxopropyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-fluorobenzyl)-2-(2-(dimethylamino)methyl)phenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-fluorobenzyl)-6-methoxy-2-(2-(pyrrolidin-1-ylmethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, N-(2-fluoro-5-(6-methoxy-4-(4-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-(4-fluorobenzyl)-2-(4-fluorophenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-fluorobenzyl)-6-methoxy-2-(2-(piperidin-1-ylmethyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(2-(((2-methoxyethyl)(methyl)amino)methyl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, N-(2-fluoro-5-(4-(3-fluoro-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 2-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile, 4-(4-chlorobenzyl)-2-(4-fluoro-2-(hydroxymethyl)phenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione, 4-(2,4-dichlorobenzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-benzyl-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-(4-isopropylbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(biphenyl-3-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-3,5-dioxo-4-(4-(trifluoromethoxy)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-(dimethylamino)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(biphenyl-4-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-benzyl-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-4-(naphthalen-2-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((6-isopropylpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-fluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-4-(4-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-3,5-dioxo-4-(quinolin-6-ylmethyl)-4,5-dihydro-1,2,4-triazin-

2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-isopropoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(3,4-dimethoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((6-cyclopentylpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-(4-chlorobenzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-((6-fluoropyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((6-chloro-5-fluoropyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((6-dimethylamino)pyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(3,4-difluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(2,4-difluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-4-((6-methoxypyridin-3-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-3,5-dioxo-4-(pyridin-3-ylmethyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-(1H-pyrazol-1-yl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(2,6-difluoro-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-3,5-dioxo-4-(quinolin-3-ylmethyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(3-fluoro-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(3,5-difluoro-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(2,3-difluoro-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-4-((1-methyl-1H-indol-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(2-fluoro-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-(3,4-difluorobenzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-(2-chloro-4-fluorobenzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chloro-2-fluorobenzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(6-methoxy-4-(3-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-((2,3-dihydrobenzo[b][1,4]dioxin-2-yl)methyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 6-methoxy-2-phenyl-4-((2-phenylthiazol-4-yl)methyl)-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-(4-chloro-2-fluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 6-methoxy-2-phenyl-4-(3-(trifluoromethoxy)benzyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(2,5-difluorobenzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chloro-3-(trifluoromethoxy)benzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-(2,4-difluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((6-fluoro-2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-(4-fluoro-3-(trifluoromethyl)benzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-benzoylbenzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(4-((6-methoxy-3,5-dioxo-2-phenyl-2,3-dihydro-1,2,4-triazin-4(5H)-yl)methyl)thiazol-2-yl)acetamide, N-(3-(4-(furo[2,3-b]pyridin-5-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(benzo[d]oxazol-6-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-fluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 3-((6-methoxy-3,5-dioxo-2-phenyl-2,3-dihydro-1,2,4-triazin-4(5H)-yl)methyl)benzonitrile, 4-((6-methoxy-3,5-dioxo-2-phenyl-2,3-dihydro-1,2,4-triazin-4(5H)-yl)methyl)benzonitrile, N-(3-(4-((7-fluoro-2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-(4-(hydroxymethyl)benzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-((6-chloro-4H-benzo[d][1,3]dioxin-8-yl)methyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 6-methoxy-2-phenyl-4-((5-(trifluoromethyl)furan-2-yl)methyl)-1,2,4-triazine-3,5(2H,4H)-dione, 6-methoxy-4-((1-methyl-1H-imidazol-2-yl)methyl)-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-(2,4-dichlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((6-(diethylamino)pyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 4-((5-chlorothiophen-2-yl)methyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-((5-chlorobenzo[b]thiophen-3-yl)methyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-(2,4-difluorobenzyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-((2,3-dihydrofuro[2,3-b]pyridin-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((4-fluoro-2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-fluoro-3-(trifluoromethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((2-ethylbenzo[d]oxazol-6-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-cyanobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-4-((2-methylbenzo[d]oxazol-6-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((6-(dimethylamino)-5-fluoropyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-(dimethylamino)-3-fluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-methoxy-4-((1-methylindolin-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(2-

fluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((4-fluoro-2,3-dihydrobenzofuran-7-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((5-chloropyridin-2-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-((5-isopropylpyridin-2-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-(3,5-difluoro-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(2-fluoro-5-(4-(4-fluoro-3-(trifluoromethoxy)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(2-fluoro-5-(4-(3-fluoro-4-(trifluoromethoxy)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide, N-(3-(6-methoxy-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide, N-(5-(4-((2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(3-(6-methoxy-4-(4-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide, N-(3-(4-((2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide, N-(3-(4-(4-fluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide, N-(3-(4-(2,4-difluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide, 6-methoxy-4-phenethyl-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-2-methoxyacetamide, N-(3-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-3-methoxypropanamide, 4-(4-chlorobenzyl)-2-(2-hydroxyphenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(3-(thiazol-2-ylamino)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(3-(2-oxopyrrolidin-1-yl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-methoxy-2-(3-(5-methyl-1,2,4-oxadiazol-3-yl)phenyl)-1,2,4-triazine-3,5(2H,4H)-dione, 6-methoxy-4-(4-methoxyphenethyl)-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-fluorophenethyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-(3,4-dimethoxyphenethyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 6-methoxy-4-(3-methoxyphenethyl)-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 6-methoxy-4-(4-methoxyphenethyl)-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione, 4-(4-fluorophenethyl)-6-methoxy-2-phenyl-1,2,4-triazine-3,5(2H,4H)-dione N-(3-(6-methoxy-4-(3-methoxyphenethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-benzyl-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(2-fluoro-5-(4-(4-fluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-(2,4-difluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(5-(4-(4,5-difluoro-2-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide N-(2-fluoro-5-(6-methoxy-3,5-dioxo-4-phenethyl-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-((6-chloro-5-fluoropyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(2-fluoro-5-(6-methoxy-4-(3-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(2,4-difluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-4-fluorophenyl)acetamide, N-(2-fluoro-5-(6-methoxy-4-((1-methylindolin-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(2-fluoro-5-(4-((6-fluoro-2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-(4-(dimethylamino)-3-fluorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(5-(4-(3,4-difluoro-2-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, 6-methoxy-2-phenyl-4-(3-phenylpropyl)-1,2,4-triazine-3,5(2H,4H)-dione, 2-(2-(4-(4-chlorobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetoneitrile, N-(5-(4-(4-cyanobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, 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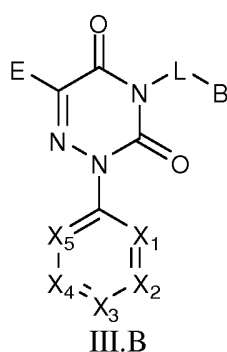
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N-(5-(4-(4-chloro-3-(pyrrolidin-1-yl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(2-fluoro-5-(4-(4-fluoro-3-(trifluoromethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-(2,4-difluoro-3-isopropoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(5-(4-((5,6-dichloropyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(5-(4-(4-cyano-2-fluoro-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, 4-((5,6-dichloropyridin-3-yl)methyl)-2-(4-fluorophenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione, N-(5-(4-(4-chloro-3-morpholinobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(5-(4-(3-(dimethylamino)-4-(trifluoromethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, 4-(4-chloro-3-methoxybenzyl)-2-(4-fluorophenyl)-6-methoxy-1,2,4-triazine-3,5(2H,4H)-dione, N-(2-fluoro-5-(4-(3-fluoro-4-(trifluoromethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamide, N-(5-(4-(4-chloro-3-(methoxymethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(2-fluoro-5-(4-(4-fluoro-3-morpholinobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(5-(4-((6-chloro-5-isopropylpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-isopropyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-ethyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-butyl-4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, 2-(6-bromo-4-(4-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile, N-(3-(4-(4-chlorobenzyl)-6-(2-(dimethylamino)ethoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-(2-methoxyethoxy)-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-(2-methoxyethoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-(methylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-ethoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-bromo-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-isopropoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-bromo-4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-(dimethylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-(hydroxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-((dimethylamino)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(6-((1H-pyrazol-1-yl)methyl)-4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-(methoxymethyl)-3,5-dioxo-4,5-dihydro-

1,2,4-triazin-2(3H)-yl)phenyl)acetamide, methyl 2-(3-acetamidophenyl)-4-(4-chlorobenzyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylate, isopropyl 2-(3-acetamidophenyl)-4-(4-chlorobenzyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazine-6-carboxylate, N-(3-(4-(4-chlorobenzyl)-6-(3-methyl-1,2,4-oxadiazol-5-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-(5-methyl-1,2,4-oxadiazol-5-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(3-(4-(4-chlorobenzyl)-6-(1,2,4-oxadiazol-5-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide, N-(2-fluoro-5-(4-(4-fluorobenzyl)-6-(2-methoxyethoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamide.

6. A pharmaceutical composition comprising a therapeutically effective amount of a compound according to claims 1 to 5 and a pharmaceutically acceptable carrier and/or excipient.
7. A compound/composition according to claims 1 to 6 for use in treating or preventing a condition in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of a GABA_B positive allosteric modulator (enhancer).
8. A compound/composition according to claims 1 to 6 for use in treating or preventing central nervous system disorders selected from the group consisting of anxiety disorders: Agoraphobia, Generalized Anxiety Disorder (GAD), Obsessive-Compulsive Disorder (OCD), Panic Disorder, Posttraumatic Stress Disorder (PTSD), Social Phobia, Other Phobias, Substance-Induced Anxiety Disorder.
9. A compound/composition according to claims 1 to 6 for use in treating or preventing central nervous system disorders selected from the group consisting of childhood disorders: Attention-Deficit/Hyperactivity Disorder).
10. A compound/composition according to claims 1 to 6 for use in treating or preventing central nervous system disorders selected from the group consisting of eating Disorders (Anorexia Nervosa, Bulimia Nervosa).
11. A compound/composition according to claims 1 to 6 for use in treating or preventing central nervous system disorders selected from the group consisting of mood disorders: Bipolar Disorders (I & II), Cyclothymic Disorder, Depression, Dysthymic Disorder, Major Depressive Disorder, Substance-Induced Mood Disorder.
12. A compound/composition according to claims 1 to 6 for use in treating or preventing central nervous system disorders selected from the group consisting of psychotic disorders: Schizophrenia, Delusional Disorder, Schizoaffective Disorder, Schizophreniform Disorder, Substance-Induced Psychotic Disorder.
13. A compound/composition according to claims 1 to 6 for use in treating or preventing central nervous system disorders selected from the group consisting of cognitive disorders: 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, Substance-Induced Persisting Dementia, Mild Cognitive Impairment.
14. A compound/composition according to claims 1 to 6 for use in treating or preventing central nervous system disorders selected from the group consisting of personality disorders: Obsessive-Compulsive Personality Disorder, Schizoid, Schizotypal disorder.
15. A compound/composition according to claims 1 to 6 for use in treating or preventing central nervous system disorders selected from the group consisting of substance-related disorders: Alcohol abuse, Alcohol dependence, Alcohol withdrawal, Alcohol withdrawal delirium, Alcohol-induced psychotic disorder, Amphetamine dependence, Amphetamine withdrawal, Cocaine dependence, Cocaine withdrawal, Nicotine dependence, Nicotine withdrawal, Opioid dependence, Opioid withdrawal.
16. A compound/composition according to claims 1 to 6 for use in treating or preventing inflammatory central nervous system disorders selected from multiple sclerosis form such as benign multiple sclerosis, amyotrophic lateral sclerosis, relapsing-remitting multiple sclerosis, secondary progressive multiple sclerosis, primary progressive multiple sclerosis, progressive-relapsing multiple sclerosis.
17. A compound/composition according to claims 1 to 6 for use in treating or preventing inflammatory or neuropathic pain, post-operative pain.
18. A compound/composition according to claims 1 to 6 for use in treating or preventing epilepsy and tremor.

19. A compound/composition according to claims 1 to 6 for use in treating or preventing urge urinary incontinence.
20. A compound/composition according to claims 1 to 6 for use in treating or preventing gastrointestinal disorders selected from the group of gastroesophageal reflux disease, transient lower oesophageal sphincter relaxations, dyspepsia, functional gastrointestinal disorders and irritable bowel syndrome.
21. A compound/composition according to claims 1 to 6 for use in treating or preventing spasticity and skeletal muscle rigidity, spinal cord injury, cerebral palsy.
22. Use of a compound according to claims 1 to 5 to prepare a tracer for imaging GABA_B receptors.
23. A compound/composition according to claims 1 to 6 in combination with an agonist of GABA_B receptor for use in treating or preventing a condition as defined in any one of claims 7 to 21, in a mammal, including a human, the treatment or prevention of which is affected or facilitated by the neuromodulatory effect of GABA_B positive allosteric modulators.
24. A process for producing a compound according to claim 3 having the formula III.B



and stereoisomeric forms, mixtures of stereoisomeric forms or pharmaceutically acceptable salts, hydrates or solvates of such compounds, wherein:

B is selected from C₆-C₁₀ aryl optionally substituted with one to 5 Y, 5 to 14 membered heteroaryl group optionally substituted by one to 8 Y, wherein said heteroaryl group comprises one, two or three heteroatoms selected from N, O, S;

Y is 1 to 5 substituents independently selected from the group consisting of hydrogen, halogen, -OCF₃, -NO₂, -CN, -CF₃, an optionally substituted-(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, halo-(C₁-C₆)alkyl, -(C₀-C₆)alkylOR¹, -(C₀-C₆)alkylNR²R³, -(C₀-C₆)alkylSR²;

R¹ at each occurrence is selected from hydrogen, -(C₁-C₆)alkyl;

R² and R³ each independently is -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl;

X₁, X₂, X₃, X₄ and X₅ are each independently selected from the group consisting of -CR⁶=, -N=;

R⁶ is 1 to 5 substituents independently selected from the group consisting of hydrogen, halogen, -OCF₃, -CN, -CF₃, -NO₂, an optionally substituted-(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₀-C₆)alkylOR⁷, -(C₀-C₆)alkylNR⁸R⁹, -(C₀-C₆)alkylSR⁸, -(C₀-C₆)alkylNR⁸C(=O)R⁹, -(C₀-C₆)alkylNR⁸C(=O)₂R⁹, -(C₀-C₆)alkylNR⁸S(=O)₂R⁹, -(C₀-C₆)alkylNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)alkylS(=O)₂R⁸, -(C₀-C₆)alkylS(=O)₂NR⁸R⁹, -(C₀-C₆)alkylC(=O)R⁸, -(C₀-C₆)alkylC(=O)OR⁷, -(C₁-C₆)alkylC(=O)NR⁸R⁹;

R⁷, R⁸ and R⁹ at each independently is selected from hydrogen, -(C₁-C₆)alkyl, -(C₃-C₇)cycloalkyl;

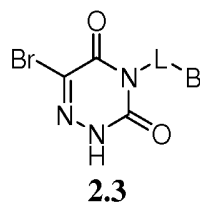
E is selected from the group consisting of -(C₁-C₆)alkyl, -(C₃-C₆)cycloalkyl, -(C₀-C₆)alkylOR¹², -(C₀-C₆)alkylNR¹³R¹⁴;

R¹², R¹³ and R¹⁴ each independently is selected from hydrogen, -(C₁-C₆)alkyl, -(C₂-C₆)alkenyl, -(C₃-C₇)cycloalkyl;

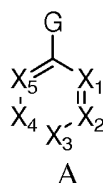
L is independently selected from a -(C₁-C₅)alkyl- substituted with 1-6 T;

T is 1 to 6 substituents independently selected from the group consisting of hydrogen, -(C₁-C₆)alkyl;

which comprises allowing a compound of formula 2.3



to react with the aryl or heteroaryl boronic acid or ester of formula A in a presence of catalytic copper systems and a base



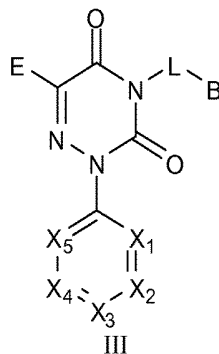
wherein:

G is a boronic acid, boronate (e.g. dimethyl boronate, pinacol boronate and the like)

And the resulting reaction product is then submitted to nucleophilic displacement of the bromine by various groups to provide the compound of formula III.B.

Patentansprüche

1. Verbindung der Formel III



und stereoisomere Formen, Gemische von stereoisomeren Formen oder pharmazeutisch akzeptable Salze, Hydrate oder Solvate solcher Verbindungen, wobei:

B ausgewählt ist aus C₆-C₁₀ Aryl, optional substituiert durch ein bis fünf Y, C₅-C₁₀ Cycloalkenyl, optional substituiert durch ein bis sieben Y, eine 5- bis 14-gliedrige Heteroarylgruppe, optional substituiert durch ein bis acht Y, wobei die genannte Heteroarylgruppe ein, zwei oder drei aus N, O, S ausgewählte Heteroatome umfasst; Y 1 bis 5 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -NO₂, -CN, -CF₃, einem optional substituierten - (C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, Halo-(C₁-C₆)Alkyl, -(C₀-C₆)AlkylOR¹, -(C₀-C₆)AlkylNR²R³, -(C₀-C₆)Alkyl- C(=NR⁴)NR²R³, -(C₀-C₆)AlkylSR², -(C₀-C₆)AlkylNR²C(=O)R³, -(C₀-C₆)AlkylNR²S(=O)₂R³, -(C₀-C₆)AlkylNR⁴C(=O)NR²R³, -(C₀-C₆)AlkylS(=O)R², -(C₀-C₆)AlkylS(=O)₂R², -(C₀-C₆)AlkylS(=O)₂NR²R³, -(C₀-C₆)AlkylC(=O)R², -(C₀-C₆)AlkylC(=O)OR¹, -(C₀-C₆)AlkylC(=O)NR²R³, -(C₀-C₆)AlkylC(=NR²)R³ oder -(C₀-C₆)AlkylC(=NOR¹)R³, Heteroaryl, Heteroarylalkyl, Arylalkyl, Arylalkylaryl, Alkylheteroaryl; wobei zwei Substituenten optional mit den intervenie-

renden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring ferner optional substituiert ist durch 1-5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R¹

bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, 3- bis 7- gliedrigem Heterocycloalkyl, -(C₆-C₁₀)Aryl, 5- oder 6- gliedrigem Heteroaryl, -(C₇-C₁₀)Arylalkyl, -C(=O)R⁵; R⁵ ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl und -(C₆-C₁₀)Aryl;

R², R³ und R⁴ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Arylalkyl oder Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Heteroaryl), -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇)cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten; wobei R², R³, R⁴ Substituenten optional mit den intervenierenden Atomen kombiniert sind, um ein bicyclisches Heterocycloalkyl zu bilden;

X₁, X₂, X₃, X₄ und X₅ jeweils unabhängig ausgewählt sind aus der Gruppe bestehend aus -CR⁶=, -N=;

R⁶ 1 bis 5 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -CN, -CF₃, -NO₂, optional substituiertem -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, Halo-(C₁-C₆)Alkyl, -(C₀-C₆)AlkylOR⁷, -(C₀-C₆)AlkylNR²R⁹, -(C₀-C₆)Alkyl-C(=NR¹⁰)NR⁸R⁹, -(C₀-C₆)AlkylSR², -(C₀-C₆)AlkylNR⁸C(=O)R⁹, -(C₀-C₆)AlkylNR²S(=O)₂R⁹, -(C₀-C₆)AlkylNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)AlkylS(=O)R⁸, -(C₀-C₆)AlkylS(=O)₂R⁸, -(C₀-C₆)AlkylS(=O)₂NR⁸R⁹, -(C₀-C₆)AlkylC(=O)R⁸, -(C₀-C₆)AlkylC(=O)OR⁷, -(C₁-C₆)AlkylC(=O)NR⁸R⁹ oder -(C₀-C₆)AlkylC(=NOR⁷)R⁹, Heteroaryl, Heteroarylalkyl, Arylalkyl, Aryl, einem 3 bis 6 Heterocycloalkyl; wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden, wobei jeder Ring optional ferner substituiert ist durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)Alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R⁷ bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, 3- bis 7- gliedrigem Heterocycloalkyl, -(C₆-C₁₀)Aryl, 5- oder 6- gliedrigem Heteroaryl, -(C₇-C₁₀)Arylalkyl, -C(=O)R¹¹; R¹¹ ausgewählt ist aus -(C₁-C₆)Alkyl, -(C₃-C₇)Cycloalkyl und -(C₆-C₁₀)Aryl;

R², R⁹ und R¹⁰ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Arylalkyl oder Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇)cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten;

E ausgewählt ist aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -NO₂, -CF₃, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, Halo-(C₁-C₆)Alkyl, -(C₀-C₆)AlkylOR¹², -(C₀-C₆)AlkylNR¹³R¹⁴, -(C₀-C₆)Alkyl-C(=NR¹⁵)NR¹³R¹⁴, -(C₀-C₆)AlkylSR¹², -(C₀-C₆)AlkylNR¹³C(=O)R¹⁴, -(C₀-C₆)AlkylNR¹³S(=O)₂R¹⁴, -(C₀-C₆)AlkylNR¹⁵C(=O)NR¹³R¹⁴, -(C₀-C₆)AlkylS(=O)R¹³, -(C₀-C₆)AlkylS(=O)₂R¹³, -(C₀-C₆)AlkylS(=O)₂NR¹³R¹⁴, -(C₀-C₆)AlkylC(=O)R¹³, -(C₀-C₆)AlkylC(=O)OR¹³, -(C₀-C₆)AlkylC(=O)NR¹³R¹⁴, -(C₀-C₆)AlkylC(=NR¹³)R¹⁴ oder -(C₀-C₆)AlkylC(=NOR¹³)R¹⁴; wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring ferner optional substituiert ist durch 1 bis 5 unabhängige Halogen-, -CN, optional substituierte -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)Alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)Alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R¹² bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, -C(=O)R¹⁶;

R¹⁶ ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₇)Cycloalkyl und -(C₆-C₁₀)Aryl;

R¹³, R¹⁴ und R¹⁵ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Alkylheteroaryl, Arylalkyl, Alkylaryl oder Aryl; von denen jedes optional substituiert ist durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇)cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten;

wenn E -H ist, dann sind die Verbindungen, für die X₃ nicht CH= oder -CF= ist, aus der Erfindung ausgeschlossen;

L unabhängig ausgewählt ist aus einem $-(C_1-C_5)Alkyl-Q_m)-$, substituiert durch 1-6 T;
m 0 oder 1 ist;

T 1 bis 6 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Fluor, $-OCF_3$, $-NO_2$, $-CN$, $-CF_3$, $-(C_1-C_6)Alkyl$, $-(C_3-C_6)Cycloalkyl$, $-(C_2-C_6)Alkenyl$, $-(C_2-C_6)Alkynyl$, Halo- $-(C_1-C_6)Alkyl$, $-(C_0-C_6)alkylOR^{17}$, $-(C_0-C_6)AlkylNR^{18}R^{19}$, $-(C_0-C_6)Alkyl-C(=NR^{20})NR^{18}R^{19}$, $-(C_0-C_6)AlkylSR^{10}$, $-(C_0-C_6)AlkylNR^{18}C(=O)R^{19}$, $-(C_0-C_6)AlkylNR^{18}C(=O)_2R^{19}$, $-(C_0-C_6)AlkylNR^{18}S(=O)_2R^{19}$, $-(C_0-C_6)AlkylNR^{20}C(=O)NR^{18}R^{19}$, $-(C_0-C_6)AlkylS(=O)R^{18}$, $-(C_0-C_6)AlkylS(=O)_2R^{18}$, $-(C_0-C_6)AlkylS(=O)_2NR^{18}R^{19}$, $-(C_0-C_6)AlkylC(=O)R^{18}$, $-(C_0-C_6)AlkylC(=O)OR^{17}$, $-(C_0-C_6)AlkylC(=O)NR^{18}R^{19}$, $-(C_0-C_6)AlkylC(=NR^{18})R^{19}$ oder $-(C_0-C_6)AlkylC(=NOR^{17})R^{19}$, Heteroaryl, Heteroarylalkyl, Arylalkyl, Alkylaryl, Alkylheteroaryl, Aryl; wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring optional ferner substituiert ist durch 1 bis 5 unabhängige Halogen-, $-CN$, $-(C_1-C_6)Alkyl$, $-O(C_0-C_6)Alkyl$, $-O(C_3-C_7)Cycloalkyl$, $-O(Aryl)$, $-O(Aryl)-(C_0-C_3)alkyl$, $-O(Heteroaryl)$, $-O(Heteroaryl)-(C_0-C_3)Alkyl$, $-O(C_1-C_3)Alkylaryl$, $-O(C_1-C_3)Alkylheteroaryl$, $-N((C_0-C_6)Alkyl)((C_0-C_3)alkylaryl)$ oder $-N((C_0-C_6)Alkyl)((C_0-C_3)alkylheteroaryl)$ Gruppen;

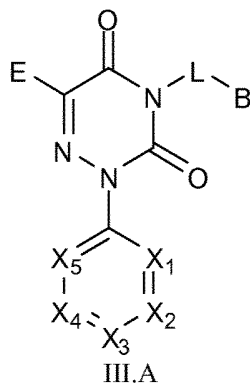
R^{17} bei jedem Vorkommen ausgewählt ist aus Wasserstoff, $-(C_1-C_6)Alkyl$, $-(C_2-C_6)Alkenyl$, $-(C_3-C_7)Cycloalkyl$, 3- bis 7- gliedrigem Heterocycloalkyl, $-(C_6-C_{10})Aryl$, 5- oder 6- gliedrigem Heteroaryl, $-(C_7-C_{10})Arylalkyl$, $-C(=O)R^{21}$; R^{21} ausgewählt ist aus Wasserstoff, $-(C_1-C_6)Alkyl$, $-(C_3-C_7)Cycloalkyl$ und $-(C_6-C_{10})Aryl$;

R^{18} , R^{19} und R^{20} jeweils unabhängig Folgendes sind: Wasserstoff, $-(C_1-C_6)Alkyl$, $-(C_3-C_6)Cycloalkyl$, $-(C_2-C_6)Alkenyl$, $-(C_2-C_6)Alkynyl$, $-(C_1-C_6)Alkylhalo$, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Arylalkyl, Alkylaryl, Alkylheteroaryl, Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, $-CN$, $-(C_1-C_6)Alkyl$, $-O(C_0-C_6)Alkyl$, $-O(C_3-C_7)Cycloalkylalkyl$, $-O(Aryl)$, $-O(Aryl)-(C_0-C_3)alkyl$, $-O(Heteroaryl)$, $-O(Heteroaryl)-(C_0-C_3)alkyl$, $-N((C_0-C_6)Alkyl)_2$, $-N((C_0-C_6)Alkyl)((C_3-C_7)cycloalkyl)$ oder $-N((C_0-C_6)Alkyl)(aryl)$ Substituenten;

$Q-C(=O)-$, $-C(0)O-$, $-C(=O)NR^{22}-$, $-OC(=O)-$, $-OC(=O)NR^{22}-$, $-NR^{22}-$, $-NR^{22}C(=O)-$, $-NR^{22}C(=O)_2-$, $-NR^{22}C(=S)-$, $-S-$, $-S(=O)-$ oder $-S(=O)_2-$ ist;

R^{22} ausgewählt ist aus $-(C_1-C_6)Alkyl$ und $-(C_6-C_{10})Aryl$;
jedes N ein N-Oxid sein kann.

2. Verbindung nach Anspruch 1 mit der Formel III.A



und stereoisomere Formen, Gemische von stereoisomeren Formen oder pharmazeutisch akzeptable Salze, Hydrate oder Solvate solcher Verbindungen,
wobei:

B ausgewählt ist aus C_6-C_{10} Aryl, optional substituiert durch ein bis fünf Y, C_5-C_{10} Cycloalkenyl, optional substituiert durch ein bis sieben Y, eine 5- bis 14-gliedrige Heteroarylgruppe, optional substituiert durch ein bis acht Y, wobei die genannte Heteroarylgruppe ein, zwei oder drei aus N, O, S ausgewählte Heteroatome umfasst;
Y 1 bis 5 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Halogen, $-OCF_3$, $-NO_2$, $-CN$, $-CF_3$, einem optional substituierten $-(C_1-C_6)Alkyl$, $-(C_3-C_6)Cycloalkyl$, $-(C_2-C_6)Alkenyl$, $-(C_2-C_6)Alkynyl$, Halo- $-(C_1-C_6)Alkyl$, $-(C_0-C_6)AlkylOR^1$, $-(C_0-C_6)AlkylNR^2R^3$, $-(C_0-C_6)Alkyl-C(=NR^4)NR^2R^3$, $-(C_0-C_6)AlkylSR^2$, $-(C_0-C_6)AlkylNR^2C(=O)R^3$, $-(C_0-C_6)AlkylNR^2S(=O)_2R^3$, $-(C_0-C_6)AlkylNR^4C(=O)NR^2R^3$, $-(C_0-C_6)AlkylS(=O)R^2$, $-(C_0-C_6)AlkylS(=O)_2R^2$, $-(C_0-C_6)AlkylS(=O)_2NR^2R^3$, $-(C_0-C_6)AlkylC(=O)R^2$, $-(C_0-C_6)AlkylC(=O)OR^1$, $-(C_0-C_6)AlkylC(=O)NR^2R^3$, $-(C_0-C_6)AlkylC(=NR^2)R^3$ oder $-(C_0-C_6)AlkylC(=NOR^1)R^3$, Heteroaryl, Heteroarylalkyl, Arylalkyl, Arylalkylaryl, Alkylheteroaryl; wobei optional zwei Substituenten mit den intervenie-

renden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring ferner optional substituiert ist durch 1-5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R¹ bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, 3- bis 7- gliedrigem Heterocycloalkyl, -(C₆-C₁₀)Aryl, 5- oder 6- gliedrigem Heteroaryl, -(C₇-C₁₀)Arylalkyl, -C(=O)R⁵; R⁵ ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl und -(C₆-C₁₀)Aryl;

R², R³ und R⁴ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆) Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Arylalkyl oder Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Heteroaryl), -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇) cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten; wobei optional R², R³, R⁴ Substituenten mit den intervenierenden Atomen kombiniert sind, um ein bicyclisches Heterocycloalkyl zu bilden;

X₁, X₂, X₃, X₄ und X₅ jeweils unabhängig ausgewählt sind aus der Gruppe bestehend aus -CR⁶=, -N=;

R⁶ 1 bis 5 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -CN, -CF₃, -NO₂, optional substituiertem -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, Halo-(C₁-C₆)Alkyl, -(C₀-C₆)AlkylOR⁷, -(C₀-C₆)AlkylNR²R⁹, -(C₀-C₆)Alkyl-C(=NR¹⁰)NR⁸R⁹, -(C₀-C₆)AlkylSR², -(C₀-C₆)AlkylNR⁸C(=O)R⁹, -(C₀-C₆)AlkylNR²S(=O)₂R⁹, -(C₀-C₆)AlkylNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)AlkylS(=O)R⁸, -(C₀-C₆)AlkylS(=O)₂R⁸, -(C₀-C₆)AlkylS(=O)₂NR⁸R⁹, -(C₀-C₆)AlkylC(=O)R⁸, -(C₀-C₆)AlkylC(=O)OR⁷, -(C₁-C₆)AlkylC(=O)NR⁸R⁹ oder -(C₀-C₆)AlkylC(=NOR⁷)R⁹, Heteroaryl, Heteroarylalkyl, Arylalkyl, Aryl, einem 3 bis 6 Heterocycloalkyl; wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring optional ferner substituiert ist durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)Alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R⁷ bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, 3- bis 7- gliedrigem Heterocycloalkyl, -(C₆-C₁₀)Aryl, 5- oder 6- gliedrigem Heteroaryl, -(C₇-C₁₀)Arylalkyl, -C(=O)R¹¹; R¹¹ ausgewählt ist aus -(C₁-C₆)Alkyl, -(C₃-C₇)Cycloalkyl und -(C₆-C₁₀)Aryl;

R⁸, R⁹ und R¹⁰ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Arylalkyl oder Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇) cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten;

E ausgewählt ist aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -NO₂, -CF₃, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, Halo-(C₁-C₆)Alkyl, -(C₀-C₆)AlkylOR¹², -(C₀-C₆)AlkylNR¹³R¹⁴, -(C₀-C₆)AlkylSR¹², -(C₀-C₆)AlkylNR¹³C(=O)R¹⁴, -(C₀-C₆)AlkylNR¹³S(=O)₂R¹⁴, -(C₀-C₆)AlkylNR¹⁵C(=O)NR¹³R¹⁴, -(C₀-C₆)AlkylS(=O)R¹³, -(C₀-C₆)AlkylS(=O)₂R¹³, -(C₀-C₆)AlkylS(=O)₂NR¹³R¹⁴, -(C₀-C₆)AlkylC(=O)R¹³, -(C₀-C₆)AlkylC(=O)OR¹³, -(C₀-C₆)AlkylC(=O)NR¹³R¹⁴, wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring ferner optional substituiert ist durch 1 bis 5 unabhängige Halogen-, -CN, optional substituierte -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)Alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)Alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R¹² bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, -C(=O)R¹⁶; R¹⁶ ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₇)Cycloalkyl und -(C₆-C₁₀)Aryl;

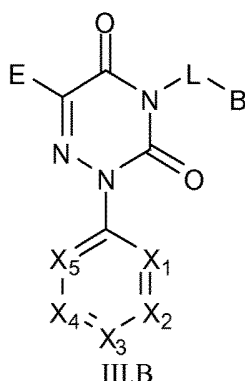
R¹³, R¹⁴ und R¹⁵ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Alkylheteroaryl, Arylalkyl, Alkylaryl oder Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇)cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten;

wenn E -H ist, dann sind die Verbindungen, für die X₃ nicht CH= oder -CF= sind, aus der Erfindung ausgeschlossen;

L unabhängig ausgewählt ist aus einem -(C₁-C₅)Alkyl-Q_m-, substituiert durch 1-6 T;
m 0 oder 1 ist;

T 1 bis 6 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Fluor, $-\text{OCF}_3$, $-\text{NO}_2$, $-\text{CN}$, $-\text{CF}_3$, $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-(\text{C}_3\text{-C}_6)\text{Cycloalkyl}$, $-(\text{C}_2\text{-C}_6)\text{Alkenyl}$, $-(\text{C}_2\text{-C}_6)\text{Alkynyl}$, Halo- $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-(\text{C}_0\text{-C}_6)\text{alkylOR}^{17}$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^{18}\text{R}^{19}$, $-(\text{C}_0\text{-C}_6)\text{Alkyl-C(=NR}^{20})\text{NR}^{18}\text{R}^{19}$, $-(\text{C}_0\text{-C}_6)\text{AlkylSR}^{18}$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^{18}\text{C(=O)R}^{19}$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^{18}\text{C(=O)}_2\text{R}^{19}$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^{18}\text{S(=O)}_2\text{R}^{19}$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^{20}\text{C(=O)NR}^{18}\text{R}^{19}$, $-(\text{C}_0\text{-C}_6)\text{AlkylS(=O)R}^{18}$, $-(\text{C}_0\text{-C}_6)\text{AlkylS(=O)}_2\text{R}^{18}$, $-(\text{C}_0\text{-C}_6)\text{AlkylS(=O)}_2\text{NR}^{18}\text{R}^{19}$, $-(\text{C}_0\text{-C}_6)\text{AlkylC(=O)R}^{18}$, $-(\text{C}_0\text{-C}_6)\text{AlkylC(=O)OR}^{17}$, $-(\text{C}_0\text{-C}_6)\text{AlkylC(=O)NR}^{18}\text{R}^{19}$, $-(\text{C}_0\text{-C}_6)\text{AlkylC(=NR}^{18})\text{R}^{19}$ oder $-(\text{C}_0\text{-C}_6)\text{AlkylC(=NOR}^{17})\text{R}^{19}$, Heteroaryl, Heteroarylalkyl, Arylalkyl, Alkylaryl, Alkylheteroaryl, Aryl; wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring optional ferner substituiert ist durch 1 bis 5 unabhängige Halogen-, $-\text{CN}$, $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-\text{O}(\text{C}_0\text{-C}_6)\text{Alkyl}$, $-\text{O}(\text{C}_3\text{-C}_7)\text{Cycloalkyl}$, $-\text{O}(\text{Aryl})$, $-\text{O}(\text{Aryl})-(\text{C}_0\text{-C}_3)\text{alkyl}$, $-\text{O}(\text{Heteroaryl})$, $-\text{O}(\text{Heteroaryl})-(\text{C}_0\text{-C}_3)\text{Alkyl}$, $-\text{O}(\text{C}_1\text{-C}_3)\text{Alkylaryl}$, $-\text{O}(\text{C}_1\text{-C}_3)\text{Alkylheteroaryl}$, $-\text{N}((\text{C}_0\text{-C}_6)\text{Alkyl})((\text{C}_0\text{-C}_3)\text{alkylaryl})$ oder $-\text{N}((\text{C}_0\text{-C}_6)\text{Alkyl})((\text{C}_0\text{-C}_3)\text{alkylheteroaryl})$ Gruppen; R^{17} bei jedem Vorkommen ausgewählt ist aus Wasserstoff, $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-(\text{C}_2\text{-C}_6)\text{Alkenyl}$, $-(\text{C}_3\text{-C}_7)\text{Cycloalkyl}$, 3- bis 7- gliedrigem Heterocycloalkyl, $-(\text{C}_6\text{-C}_{10})\text{Aryl}$, 5- oder 6- gliedrigem Heteroaryl, $-(\text{C}_7\text{-C}_{10})\text{Arylalkyl}$, $-\text{C(=O)R}^{21}$; R^{21} ausgewählt ist aus Wasserstoff, $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-(\text{C}_3\text{-C}_7)\text{Cycloalkyl}$ und $-(\text{C}_6\text{-C}_{10})\text{Aryl}$; R^{18} , R^{19} und R^{20} jeweils unabhängig Folgendes sind: Wasserstoff-, $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-(\text{C}_3\text{-C}_6)\text{Cycloalkyl}$, $-(\text{C}_2\text{-C}_6)\text{Alkenyl}$, $-(\text{C}_2\text{-C}_6)\text{Alkynyl}$, $-(\text{C}_1\text{-C}_6)\text{Alkylhalo}$, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Arylalkyl, Alkylaryl, Alkylheteroaryl, Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, $-\text{CN}$, $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-\text{O}(\text{C}_0\text{-C}_6)\text{Alkyl}$, $-\text{O}(\text{C}_3\text{-C}_7)\text{Cycloalkylalkyl}$, $-\text{O}(\text{Aryl})$, $-\text{O}(\text{Aryl})-(\text{C}_0\text{-C}_3)\text{alkyl}$, $-\text{O}(\text{Heteroaryl})$, $-\text{O}(\text{Heteroaryl})-(\text{C}_0\text{-C}_3)\text{alkyl}$, $-\text{N}((\text{C}_0\text{-C}_6)\text{Alkyl})_2$, $-\text{N}((\text{C}_0\text{-C}_6)\text{Alkyl})((\text{C}_3\text{-C}_7)\text{cycloalkyl})$ oder $-\text{N}((\text{C}_0\text{-C}_6)\text{Alkyl})(\text{aryl})$ Substituenten; $\text{Q} -\text{C(=O)-}$, $-\text{C(=O)O-}$, $-\text{C(=O)NR}^{22}$, $-\text{OC(=O)-}$, $-\text{OC(=O)NR}^{22}$, $-\text{NR}^{22}$, $-\text{NR}^{22}\text{C(=O)-}$, $-\text{NR}^{22}\text{C(=O)}_2$, $-\text{NR}^{22}\text{C(=S)-}$, $-\text{S-}$, $-\text{S(=O)-}$ oder $-\text{S(=O)}_2$ ist; R^{22} ausgewählt ist aus $-(\text{C}_1\text{-C}_6)\text{Alkyl}$ und $-(\text{C}_6\text{-C}_{10})\text{Aryl}$; jedes N ein N-Oxid sein kann.

3. Verbindung nach Anspruch 2 mit der Formel III.B



und stereoisomere Formen, Gemische von stereoisomeren Formen oder pharmazeutisch akzeptable Salze, Hydrate oder Solvate solcher Verbindungen, wobei:

B ausgewählt ist aus $\text{C}_6\text{-C}_{10}$ Aryl, optional substituiert durch ein bis fünf Y, $\text{C}_5\text{-C}_{10}$ Cycloalkenyl, optional substituiert durch ein bis sieben Y, eine 5- bis 14-gliedrige Heteroarylgruppe, optional substituiert durch ein bis acht Y, wobei die genannte Heteroarylgruppe ein, zwei oder drei aus N, O, S ausgewählte Heteroatome umfasst; Y 1 bis 5 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Halogen, $-\text{OCF}_3$, $-\text{NO}_2$, $-\text{CN}$, $-\text{CF}_3$, einem optional substituierten $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-(\text{C}_3\text{-C}_6)\text{Cycloalkyl}$, $-(\text{C}_2\text{-C}_6)\text{Alkenyl}$, $-(\text{C}_2\text{-C}_6)\text{Alkynyl}$, Halo- $-(\text{C}_1\text{-C}_6)\text{Alkyl}$, $-(\text{C}_0\text{-C}_6)\text{AlkylOR}^1$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^2\text{R}^3$, $-(\text{C}_0\text{-C}_6)\text{Alkyl-C(=NR}^4)\text{NR}^2\text{R}^3$, $-(\text{C}_0\text{-C}_6)\text{AlkylSR}^2$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^2\text{C(=O)R}^3$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^2\text{S(=O)}_2\text{R}^3$, $-(\text{C}_0\text{-C}_6)\text{AlkylNR}^2\text{C(=O)NR}^2\text{R}^3$, $-(\text{C}_0\text{-C}_6)\text{AlkylS(=O)R}^2$, $-(\text{C}_0\text{-C}_6)\text{AlkylS(=O)}_2\text{R}^2$, $-(\text{C}_0\text{-C}_6)\text{AlkylS(=O)}_2\text{NR}^2\text{R}^3$, $-(\text{C}_0\text{-C}_6)\text{AlkylC(=O)R}^2$, $-(\text{C}_0\text{-C}_6)\text{AlkylC(=O)OR}^1$, $-(\text{C}_0\text{-C}_6)\text{AlkylC(=O)NR}^2\text{R}^3$, $-(\text{C}_0\text{-C}_6)\text{AlkylC(=NR}^2)\text{R}^3$ oder $-(\text{C}_0\text{-C}_6)\text{AlkylC(=NOR}^1)\text{R}^3$, Heteroaryl, Heteroarylalkyl, Arylalkyl, Arylalkylaryl, Alkylheteroaryl; wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring ferner optional substituiert ist durch 1-5 unabhängige Halogen-, $-\text{CN}$, $-(\text{C}_1\text{-C}_6)\text{Alkyl}$,

-O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R¹ bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, 3- bis 7- gliedrigem Heterocycloalkyl, -(C₆-C₁₀)Aryl, 5- oder 6- gliedrigem Heteroaryl, -(C₇-C₁₀)Arylalkyl, -C(=O)R⁵; R⁵ ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl und -(C₆-C₁₀)Aryl;

R², R³ und R⁴ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Arylalkyl oder Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Heteroaryl), -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇)cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten; wobei optional R², R³, R⁴ Substituenten mit den intervenierenden Atomen kombiniert sind, um ein bicyclisches Heterocycloalkyl zu bilden;

X₁, X₂, X₃, X₄ und X₅ jeweils unabhängig ausgewählt sind aus der Gruppe bestehend aus -CR⁶=, -N=;

R⁶ 1 bis 5 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -CN, -CF₃, -NO₂, optional substituiertem -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, Halo-(C₁-C₆)Alkyl, -(C₀-C₆)AlkylOR⁷, -(C₀-C₆)AlkylNR²R⁹, -(C₀-C₆)AlkylSR⁸, -(C₀-C₆)AlkylNR⁸C(=O)⁹, -(C₀-C₆)AlkylNR⁸S(=O)₂R⁹, -(C₀-C₆)AlkylNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)AlkylS(=O)R², -(C₀-C₆)AlkylS(=O)₂R⁸, -(C₀-C₆)AlkylS(=O)₂NR²R⁹, -(C₀-C₆)AlkylC(=O)R⁸, -(C₀-C₆)AlkylC(=O)OR⁷, -(C₁-C₆)AlkylC(=O)NR⁸R⁹, Heteroaryl, Heteroarylalkyl, Arylalkyl, Aryl, einem 3 bis 6 Heterocycloalkyl; wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring optional ferner substituiert ist durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R⁷ bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, 3- bis 7- gliedrigem Heterocycloalkyl, -(C₆-C₁₀)Aryl, 5- oder 6- gliedrigem Heteroaryl, -(C₇-C₁₀)Arylalkyl, -C(=O)R¹¹; R¹¹ ausgewählt ist aus -(C₁-C₆)Alkyl, -(C₃-C₇)Cycloalkyl und -(C₆-C₁₀)Aryl;

R², R⁹ und R¹⁰ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Arylalkyl oder Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇)cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten;

E ausgewählt ist aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -CF₃, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₀-C₆)AlkylOR¹², -(C₀-C₆)AlkylNR¹³R¹⁴, -(C₀-C₆)AlkylS(=O)R¹³, -(C₀-C₆)AlkylS(=O)₂R¹³, -(C₀-C₆)AlkylC(=O)R¹³, -(C₀-C₆)AlkylC(=O)OR¹³, -(C₀-C₆)AlkylC(=O)NR¹³R¹⁴; wobei optional zwei Substituenten mit den intervenierenden Atomen kombiniert sind, um einen bicyclischen Heterocycloalkyl-, Aryl- oder Heteroarylring zu bilden; wobei jeder Ring optional ferner substituiert ist durch 1 bis 5 unabhängige Halogen-, -CN, optional substituierte -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -O(C₁-C₃)Alkylaryl, -O(C₁-C₃)Alkylheteroaryl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₀-C₃)Alkylaryl) oder -N((C₀-C₆)Alkyl)((C₀-C₃)alkylheteroaryl)) Gruppen;

R¹² bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl, -C(=O)R¹⁶; R¹⁶ ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₇)Cycloalkyl und -(C₆-C₁₀)Aryl;

R¹³ und R¹⁴ jeweils unabhängig Folgendes sind: Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₂-C₆)Alkenyl, -(C₂-C₆)Alkynyl, -(C₁-C₆)Alkylhalo, Heterocycloalkyl, Heteroaryl, Heteroarylalkyl, Alkylheteroaryl, Arylalkyl, Alkylaryl oder Aryl; von denen beliebige optional substituiert sind durch 1 bis 5 unabhängige Halogen-, -CN, -(C₁-C₆)Alkyl, -O(C₀-C₆)Alkyl, -O(C₃-C₇)Cycloalkylalkyl, -O(Aryl), -O(Aryl)-(C₀-C₃)alkyl, -O(Heteroaryl), -O(Heteroaryl)-(C₀-C₃)alkyl, -N((C₀-C₆)Alkyl)₂, -N((C₀-C₆)Alkyl)((C₃-C₇)cycloalkyl) oder -N((C₀-C₆)Alkyl)(aryl) Substituenten;

wenn E -H ist, dann sind die Verbindungen, für die X₃ nicht CH= oder -CF= sind, aus der Erfindung ausgeschlossen;

L unabhängig ausgewählt ist aus einem -(C₁-C₅)Alkyl-Q_m-, substituiert durch 1-6 T; m 0 oder 1 ist;

T 1 bis 6 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, -(C₁-C₆)Alkyl; Q-C(=O)-, -C(=O)O-, -C(=O)NR²²-, -OC(=O)-, -OC(=O)NR²²-, -NR²²-, -NR²²C(=O)-, -NR²²C(=O)₂-, -NR²²C(=S)-, -S-, -S(=O)- oder -S(=O)₂- ist; R²² ausgewählt ist aus -(C₁-C₆)Alkyl und -(C₆-C₁₀)Aryl; jedes N ein N-Oxid sein kann.

4. Verbindung nach den Ansprüchen 1 bis 3, die als optische Isomere vorliegen können, wobei die genannte Verbindung entweder das racemische Gemisch oder ein individuelles optisches Isomer ist.
5. Verbindung nach den Ansprüchen 1 bis 4, wobei die genannte Verbindung ausgewählt ist aus:

N-(3-(4-(4-Chlorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)methansulfonamid, 4-(4-chlorbenzyl)-2-(2-methoxyphenyl)-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-Nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Cyanobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(Naphthalen-2-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(Benzo[c][1,2,5]oxadiazol-5-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Fluorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-Benzyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(3,5-Dioxo-4-(4-(trifluormethyl)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, Methyl 4-((2-(3-acetamidophenyl)-3,5-dioxo-2,3-dihydro-1,2,4-triazin-4(5H)-yl)methyl)benzoat, N-(3-(4-(naphthalen-1-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Brombenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(3-Chlorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Isopropylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(3,4-Dimethylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Chlorpyridin-3-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlor-2-fluorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-Benzyl-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlor-3-fluorbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Fluorpyridin-3-yl)methyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Isopropylpyridin-3-yl)methyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Fluorbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(4-(4-fluorbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Methoxybenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Isopropylbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-(4-Chlorbenzyl)-6-methyl-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(6-Methyl-4-(naphthalen-2-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Cyclopentylpyridin-3-yl)methyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(3,4-Difluorbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(3,4-Dimethoxybenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(2,4-Difluorbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Isopropoxybenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methyl-4-(naphthalen-1-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methyl-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(6-methyl-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(4-(3-methoxybenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 3-(4-(4-Fluorbenzyl)-6-methyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-N-methylbenzamid, N-(3-(4-(4-Fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-3,5-dioxo-4-(4-(trifluormethyl)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlor-3-fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-(4-Chlor-3-fluorbenzyl)-6-methoxy-2-(2-methoxyphenyl)-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-Chlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)methansulfonamid, 4-(4-Chlorbenzyl)-6-methoxy-2-(2-methoxyphenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 1-(3-(4-(4-Chlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-3-methylharnstoff, N-(3-(4-((6-Chlorpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(6-methoxy-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(4-Chlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)pyridin-3-yl)acetamid, N-(3-(6-Methoxy-4-((6-methylpyridin-3-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(4-Chlor-3-fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)pyridin-3-yl)acetamid, 4-(4-Chlorbenzyl)-2-(4-fluor-2-methoxyphenyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(2-(methoxymethyl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(3-(2-methoxyethyl)phenyl)-1,2,4-

triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(2-(trifluormethoxy)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-2-(2,3-dimethoxyphenyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(2-(morpholinomethyl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(2-(2-methoxyethyl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(3-((2-methyl-1,3-dioxolan-2-yl)methyl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(3-(2-oxopropyl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Fluorbenzyl)-2-(2-((dimethylamino)methyl)phenyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Fluorbenzyl)-6-methoxy-2-(2-(pyrrolidin-1-ylmethyl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, N-(2-Fluor-5-(6-methoxy-4-(4-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-(4-Fluorbenzyl)-2-(4-fluorphenyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Fluorbenzyl)-6-methoxy-2-(2-(piperidin-1-ylmethyl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(2-(((2-methoxyethyl)(methyl)amino)methyl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, N-(2-Fluor-5-(4-(3-fluor-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 2-(4-(4-Chlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitril, 4-(4-Chlorbenzyl)-2-(4-fluor-2-(hydroxymethyl)phenyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, 4-(2,4-Dichlorbenzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-Benzyl-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-Isopropylbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(Biphenyl-3-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-3,5-dioxo-4-(4-(trifluormethoxy)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-(Dimethylamino)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(Biphenyl-4-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-Benzyl-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-(naphthalen-2-ylmethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Isopropylpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-(4-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-(4-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Isopropoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(3,4-Dimethoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Cyclopentylpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-(4-Chlorbenzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-((6-Fluorpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Chlor-5-fluorpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Dimethylamino)pyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((2,3-Dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(3,4-Difluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(2,4-Difluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-((6-methoxypyridin-3-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-3,5-dioxo-4-(pyridin-3-ylmethyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-(1H-Pyrazol-1-yl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(2,6-Difluor-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-3,5-dioxo-4-(chinolin-3-ylmethyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(3-Fluor-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(3,5-Difluor-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(2,3-Difluor-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-((1-methyl-1H-indol-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(2-Fluor-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-(3,4-Difluorbenzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-(2-Chlor-4-fluorbenzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlor-2-fluorbenzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(6-Methoxy-4-(3-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-((2,3-Dihydrobenzo[b][1,4]dioxin-2-yl)methyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 6-Methoxy-2-phenyl-4-((2-phenylthiazol-4-yl)methyl)-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-Chlor-2-fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 6-Methoxy-2-phenyl-4-(3-(trifluormethoxy)benzyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(2,5-Difluorbenzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlor-3-(trifluormethoxy)benzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(2,4-Difluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-Fluor-2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-(4-Fluor-3-(trifluormethyl)benzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Benzoylbenzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(4-((6-Methoxy-3,5-dioxo-2-phenyl-2,3-dihydro-1,2,4-triazin-

4(5H)-yl)methylthiazol-2-yl)acetamid, N-(3-(4-(Furo[2,3-b]pyridin-5-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(Benzo[d]oxazol-6-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Fluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 3-((6-Methoxy-3,5-dioxo-2-phenyl-2,3-dihydro-1,2,4-triazin-4(5H)-yl)methyl)benzonitril, 4-((6-Methoxy-3,5-dioxo-2-phenyl-2,3-dihydro-1,2,4-triazin-4(5H)-yl)methyl)benzonitril, N-(3-(4-((7-Fluor-2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-(4-(Hydroxymethyl)benzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-((6-Chlor-4H-benzo[d][1,3]dioxin-8-yl)methyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 6-Methoxy-2-phenyl-4-((5-(trifluormethyl)furan-2-yl)methyl)-1,2,4-triazin-3,5(2H,4H)-dion, 6-Methoxy-4-((1-methyl-1H-imidazol-2-yl)methyl)-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(2,4-Dichlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-(Diethylamino)pyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 4-((5-Chlorothiophen-2-yl)methyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-((5-Chlorbenzo[b]thiophen-3-yl)methyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-(2,4-Difluorbenzyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-((2,3-Dihydrofuro[2,3-b]pyridin-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((4-Fluor-2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Fluor-3-(trifluormethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((2-Ethylbenzo[d]oxazol-6-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Cyanobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-((2-methylbenzo[d]oxazol-6-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((6-(Dimethylamino)-5-fluorpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-(Dimethylamino)-3-fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-((1-methylindolin-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(2-Fluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((4-Fluor-2,3-dihydrobenzofuran-7-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((2,2-Difluorbenzo[d][1,3]dioxol-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((5-Chlorpyridin-2-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-((5-Isopropylpyridin-2-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(3,5-Difluor-4-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(4-(4-fluor-3-(trifluormethoxy)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(4-(3-fluor-4-(trifluormethoxy)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamid, N-(3-(6-Methoxy-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamid, N-(5-(4-((2,3-Dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(3-(6-Methoxy-4-(4-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamid, N-(3-(4-((2,3-Dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamid, N-(3-(4-(4-Fluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamid, N-(3-(4-(2,4-Difluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-N-methylacetamid, 6-Methoxy-4-phenethyl-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-Chlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-2-methoxyacetamid, N-(3-(4-(4-Chlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)-3-methoxypropanamid, 4-(4-Chlorbenzyl)-2-(2-hydroxyphenyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(3-(thiazol-2-ylamino)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(3-(2-oxopyrrolidin-1-yl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Chlorbenzyl)-6-methoxy-2-(3-(5-methyl-1,2,4-oxadiazol-3-yl)phenyl)-1,2,4-triazin-3,5(2H,4H)-dion, 6-Methoxy-4-(4-methoxyphenethyl)-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Fluorphenethyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-(3,4-Dimethoxyphenethyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 6-Methoxy-4-(3-methoxyphenethyl)-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 6-Methoxy-4-(4-methoxyphenethyl)-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-Fluorphenethyl)-6-methoxy-2-phenyl-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(6-Methoxy-4-(3-methoxyphenethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-Benzyl-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(4-(4-fluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(2,4-Difluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-(4,5-Difluor-2-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(6-methoxy-3,5-dioxo-4-phenethyl-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-((6-Chlor-5-fluorpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(6-

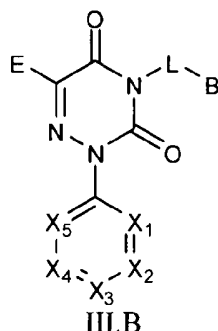
methoxy-4-(3-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(2,4-difluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-4-fluorphenyl)acetamid, N-(2-Fluor-5-(6-methoxy-4-((1-methylindolin-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(4-((6-fluor-2,3-dihydrobenzofuran-5-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(4-(Dimethylamino)-3-fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-(3,4-Difluor-2-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, 6-Methoxy-2-phenyl-4-(3-phenylpropyl)-1,2,4-triazin-3,5(2H,4H)-dion, 2-(2-(4-(4-Chlorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetanitril, N-(5-(4-(4-Cyanobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(6-methoxy-4-(2-methoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(4-Cyano-3-fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-((6-Cyanopyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(4-((6-isopropylpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(6-methoxy-4-((6-methoxypyridin-3-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(6-methoxy-3,5-dioxo-4-(chinolin-6-ylmethyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(Benzofuran-5-ylmethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(4-(4-fluorphenethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 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N-(3-(6-Methoxy-4-((1-methyl-1H-indazol-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-4-((2-methyl-2H-indazol-5-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-((2,3-Dihydrobenzofuran-6-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-((2,3-Dihydrobenzofuran-4-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(3-(6-Methoxy-4-((5-methoxypyridin-3-yl)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Cyano-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Methoxy-3,5-dioxo-4-phenethyl-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(4-(4-fluor-2-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 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1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-(4-Chlor-2-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-(4-Chlor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-(2,4-Difluorphenethyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(3-(4-((5-Fluor-6-methoxypyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Chlor-5-(4-(4-chlor-2-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Chlor-5-(4-(4-chlor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(3,4-Difluor-5-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Chlor-5-(4-(4-chlor-2-fluorbenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 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N-(5-(4-(4-Brom-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(4-(4-isopropyl-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 2-(4-Fluorphenyl)-4-(4-isopropyl-3-methoxybenzyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, N-(5-(4-(4-Chlor-3-(pyrrolidin-1-yl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(4-(4-fluor-3-(trifluormethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-(2,4-Difluor-3-isopropoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-((5,6-Dichlorpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-(4-Cyano-2-fluor-3-methoxybenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, 4-((5,6-Dichlorpyridin-3-yl)methyl)-2-(4-fluorphenyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, N-(5-(4-(4-Chlor-3-morpholinobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(5-(4-(3-(Dimethylamino)-4-(trifluormethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, 4-(4-Chlor-3-methoxybenzyl)-2-(4-fluorphenyl)-6-methoxy-1,2,4-triazin-3,5(2H,4H)-dion, N-(2-Fluor-5-(4-(3-fluor-4-(trifluormethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-

2(3H)-yl)phenyl)-N-methylacetamid, N-(5-(4-(4-Chlor-3-(methoxymethyl)benzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(2-Fluor-5-(4-(4-fluor-3-morpholinobenzyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(5-(4-((6-Chlor-5-isopropylpyridin-3-yl)methyl)-6-methoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorphenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-isopropyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-ethyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Butyl-4-(4-chlorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, 2-(6-Brom-4-(4-fluorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitril, N-(3-(4-(4-Chlorbenzyl)-6-(2-(dimethylamino)ethoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-(2-Methoxyethoxy)-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-(2-methoxyethoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-(methylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-ethoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Brom-4-(4-methylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-isopropoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-Brom-4-(4-chlorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-(dimethylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-(hydroxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-((dimethylamino)methyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(6-((1H-Pyrazol-1-yl)methyl)-4-(4-chlorbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-(methoxymethyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, Methyl 2-(3-acetamidophenyl)-4-(4-chlorbenzyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazin-6-carboxylat, Isopropyl 2-(3-acetamidophenyl)-4-(4-chlorbenzyl)-3,5-dioxo-2,3,4,5-tetrahydro-1,2,4-triazin-6-carboxylat, N-(3-(4-(4-Chlorbenzyl)-6-(3-methyl-1,2,4-oxadiazol-5-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-(5-methyl-1,2,4-oxadiazol-3-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(3-(4-(4-Chlorbenzyl)-6-(1,2,4-oxadiazol-5-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid, N-(2-Fluor-5-(4-(4-fluorbenzyl)-6-(2-methoxyethoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phenyl)acetamid.

6. Pharmazeutische Zusammensetzung, die eine therapeutisch wirksame Menge einer Verbindung nach den Ansprüchen 1 bis 5 und einen pharmazeutisch akzeptablen Träger und/oder Hilfsstoff beinhaltet.
7. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung eines Zustands in einem Säugetier, einschließlich einem Menschen, dessen Behandlung oder Verhütung durch die neuromodulatorische Wirkung eines GABA_B-positiven allosterischen Modulators (Enhancers) beeinflusst oder erleichtert wird.
8. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Störungen des zentralen Nervensystems, ausgewählt aus der Gruppe bestehend aus Angststörungen: Agoraphobie, generalisierte Angststörung (GAD), Zwangsstörung (OCD), Panikstörung, posttraumatische Belastungsstörung (PTSD), soziale Phobie, andere Phobien, substanzinduzierte Angststörung.
9. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Störungen des zentralen Nervensystems, ausgewählt aus der Gruppe bestehend aus Kindheitsstörungen: Aufmerksamkeitsdefizit/Hyperaktivitäts-Störung.
10. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Störungen des zentralen Nervensystems, ausgewählt aus der Gruppe bestehend aus Essstörungen (Anorexia nervosa, Bulimia nervosa).
11. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Störungen des zentralen Nervensystems, ausgewählt aus der Gruppe bestehend aus Gemütsstörungen: bipolare Störungen (I & II), zyklotyme Störung, Depression, dysthyme Störung, Major Depressive Disorder (MDD), substanzinduzierte Gemütsstörung.
12. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Störungen des zentralen Nervensystems, ausgewählt aus der Gruppe bestehend aus psychotischen Störungen: Schizophrenie, Wahnstörungen, schizoaffektive Störung, schizophreniforme Störung, substanzinduzierte psychotische Störung.

13. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Störungen des zentralen Nervensystems, ausgewählt aus der Gruppe bestehend aus kognitiven Störungen: Delirium, substanzinduziertes fortdauerndes Delirium, Demenz, Demenz aufgrund einer HIV-Erkrankung, Demenz aufgrund der Huntington-Krankheit, Demenz aufgrund der Parkinson-Krankheit, Demenz des Alzheimer-Typs, substanzinduzierte fortdauernde Demenz, milde kognitive Beeinträchtigung.
14. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Störungen des zentralen Nervensystems, ausgewählt aus der Gruppe bestehend aus Persönlichkeitsstörungen: zwanghafte Persönlichkeitsstörung, schizoide, schizotype Störung.
15. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Störungen des zentralen Nervensystems, ausgewählt aus der Gruppe bestehend aus substanzbezogenen Störungen: Alkoholmissbrauch, Alkoholabhängigkeit, Alkoholentzug, Alkoholentzugsdelirium, alkoholinduzierte psychotische Störung, Amphetaminabhängigkeit, Amphetaminentzug, Kokainabhängigkeit, Kokainentzug, Nikotinabhängigkeit, Nikotinentzug, Opioidabhängigkeit, Opioidentzug.
16. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von entzündlichen Störungen des zentralen Nervensystems, ausgewählt aus Formen von multipler Sklerose wie benigne multiple Sklerose, amyotrophe Lateralsklerose, schubförmige multiple Sklerose, sekundäre progressive multiple Sklerose, primäre progressive multiple Sklerose, primäre progressive multiple Sklerose, progressive wiederkehrende multiple Sklerose.
17. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von entzündlichen oder neuropathischen Schmerzen, postoperativen Schmerzen.
18. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Epilepsie und Tremor.
19. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Harndranginkontinenz.
20. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von gastrointestinalen Störungen, ausgewählt aus der Gruppe bestehend aus Refluxösophagitis, transienten unteren ösophagialen Sphinkterrelaxationen, Dyspepsie, funktionellen gastrointestinalen Störungen und Reizdarmsyndrom.
21. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 zur Verwendung bei der Behandlung oder Verhütung von Spastizität und Skelettmuskelrigidität, Rückenmarksverletzung, Zerebralparese.
22. Verwendung einer Verbindung nach den Ansprüchen 1 bis 5 zur Herstellung eines Tracers zum Abbilden von GABA_B-Rezeptoren.
23. Verbindung/Zusammensetzung nach den Ansprüchen 1 bis 6 in Kombination mit einem GABA_B-Rezeptor-Agonisten zur Verwendung bei der Behandlung oder Verhütung eines Zustands wie in einem der Ansprüche 7 bis 21 definiert in einem Säugetier, einschließlich einem Menschen, dessen Behandlung oder Verhütung durch die neuromodulatorische Wirkung von GABA_B-positiven allosterischen Modulatoren beeinflusst oder erleichtert wird.
24. Verfahren zur Herstellung einer Verbindung nach Anspruch 3 mit der Formel III.B



und stereoisomere Formen, Gemische von stereoisomeren Formen oder pharmazeutisch akzeptable Salze, Hydrate oder Solvate solcher Verbindungen, wobei:

B ausgewählt ist aus C₆-C₁₀ Aryl, optional substituiert durch ein bis fünf Y, eine 5- bis 14-gliedrige Heteroarylgruppe, optional substituiert durch ein bis acht Y, wobei die genannte Heteroarylgruppe ein, zwei oder drei aus N, O, S ausgewählte Heteroatome umfasst;

Y 1 bis 5 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -NO₂, -CN, -CF₃, einem optional substituierten -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, Halo-(C₁-C₆)Alkyl, -(C₀-C₆)AlkylOR¹, -(C₀-C₆)AlkylNR²R³, -(C₀-C₆)AlkylSR²;

R¹ bei jedem Vorkommen ausgewählt ist aus Wasserstoff, -(C₁-C₆)Alkyl;

R² und R³ jeweils unabhängig -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl sind;

X₁, X₂, X₃, X₄ und X₅ jeweils unabhängig ausgewählt sind aus der Gruppe bestehend aus -CR⁶=, -N=;

R⁶ 1 bis 5 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, Halogen, -OCF₃, -CN, -CF₃, -NO₂, optional substituiertem -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₀-C₆)AlkylOR⁷, -(C₀-C₆)AlkylNR⁸R⁹, -(C₀-C₆)AlkylSR⁸, -(C₀-C₆)AlkylNR⁸C(=O)R⁹, -(C₀-C₆)AlkylNR⁸C(=O)₂R⁹, -(C₀-C₆)AlkylNR⁸S(=O)₂R⁹, -(C₀-C₆)AlkylNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)AlkylS(=O)₂R⁸, -(C₀-C₆)AlkylS(=O)₂NR⁸R⁹, -(C₀-C₆)AlkylC(=O)R⁸, -(C₀-C₆)AlkylC(=O)OR⁷, -(C₁-C₆)AlkylC(=O)NR⁸R⁹;

R⁷, R⁸ und R⁹ jeweils unabhängig ausgewählt sind aus Wasserstoff, -(C₁-C₆)Alkyl, -(C₃-C₇)Cycloalkyl;

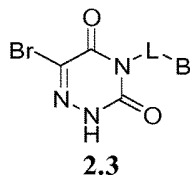
E ausgewählt ist aus der Gruppe bestehend aus -(C₁-C₆)Alkyl, -(C₃-C₆)Cycloalkyl, -(C₀-C₆)AlkylOR¹², -(C₀-C₆)AlkylNR¹³R¹⁴;

R¹², R¹³ und R¹⁴ jeweils unabhängig ausgewählt sind aus: Wasserstoff, -(C₁-C₆)Alkyl, -(C₂-C₆)Alkenyl, -(C₃-C₇)Cycloalkyl;

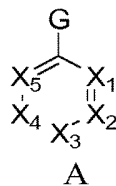
L unabhängig ausgewählt ist aus einem -(C₁-C₅)Alkyl-, substituiert durch 1-6 T;

T 1 bis 6 Substituenten ist, unabhängig ausgewählt aus der Gruppe bestehend aus Wasserstoff, -(C₁-C₆)Alkyl;

das das Reagierenlassen einer Verbindung von Formel 2.3



mit der/dem Aryl- oder Heteroaryl-Borsäure oder -ester von Formel A in Anwesenheit von katalytischen Kupfersystemen und einer Base

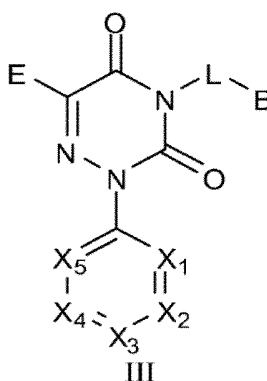


beinhaltet, wobei:

G eine Borsäure, Boronat (z.B. Dimethylboronat, Pinacolboronat und dergleichen) ist, und das resultierende Reaktionsprodukt dann einer nukleophilen Verdrängung des Broms durch verschiedene Gruppen unterzogen wird, um die Verbindung von Formel III.B zu erhalten.

Revendications

1. Composé de formule III



et formes stéréoisomères, mélanges de formes stéréoisomères ou sels, hydrates ou solvates pharmaceutiquement acceptables de tels composés, dans lequel :

B est sélectionné parmi un aryle en C₆-C₁₀ en option substitué par un à 5 Y, un cycloalkényle en C₅-C₁₀ en option substitué par un à 7 Y, un groupement hétéroaryle à 5 à 14 chaînons en option substitué par un à 8 Y, ledit groupement hétéroaryle comprenant un, deux ou trois hétéroatomes sélectionnés parmi un N, un O et un S ; Y représente 1 à 5 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, halogène, -OCF₃, -NO₂, -CN, -CF₃, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR¹, -(alkyle en C₀-C₆)NR²R³, -(alkyle en C₀-C₆)C(=NR⁴)NR²R³, -(alkyle en C₀-C₆)SR², -(alkyle en C₀-C₆)NR²C(=O)R³, -(alkyle en C₀-C₆)NR²S(=O)₂R³, -(alkyle en C₀-C₆)NR⁴C(=O)NR²R³, -(alkyle en C₀-C₆)S(=O)R², -(alkyle en C₀-C₆)S(=O)₂R², -(alkyle en C₀-C₆)S(=O)₂NR²R³, -(alkyle en C₀-C₆)C(=O)R², -(alkyle en C₀-C₆)C(=O)OR¹, -(alkyle en C₀-C₆)C(=O)NR²R³, -(alkyle en C₀-C₆)C(=NR²)R³, -(alkyle en C₀-C₆)C(=NOR¹)R³, hétéroaryle, hétéroarylalkyle, arylalkyle, arylalkylaryle ou alkylhétéroaryle en option substitué ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) ; Chaque R¹ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇, hétérocycloalkyle à 3 à 7 chaînons, aryle en C₆-C₁₀, hétéroaryle à 5 ou 6 chaînons, -(aryle en C₇-C₁₀)alkyle et -C(=O)R⁵ ; R⁵ est sélectionné parmi un hydrogène, un alkyle en C₁-C₆ et un aryle en C₆-C₁₀ ; R², R³ et R⁴ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle,

arylalkyle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(hétéroaryle), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ; les substituants R², R³, R⁴ étant en option combinés aux atomes impliqués pour former un hétérocycloalkyle bicyclique ;

X₁, X₂, X₃, X₄ et X₅ sont chacun indépendamment sélectionnés dans le groupe consistant en un -CR⁶= et un -N= ; R⁶ représente 1 à 5 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, halogène, -OCF₃, -CN, -CF₃, -NO₂, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR⁷, -(alkyle en C₀-C₆)NR⁸R⁹, -(alkyle en C₀-C₆)C(=NR¹⁰)NR⁸R⁹, -(alkyle en C₀-C₆)SR⁸, -(alkyle en C₀-C₆)NR⁸C(=O)R⁹, -(alkyle en C₀-C₆)NR⁸S(=O)₂R⁹, -(alkyle en C₀-C₆)NR¹⁰C(=O)NR⁸R⁹, -(alkyle en C₀-C₆)S(=O)R⁸, -(alkyle en C₀-C₆)S(=O)₂R⁸, -(alkyle en C₀-C₆)S(=O)₂NR⁸R⁹, -(alkyle en C₀-C₆)C(=O)R⁸, -(alkyle en C₀-C₆)C(=O)OR⁷, -(alkyle en C₁-C₆)C(=O)NR⁸R⁹, -(alkyle en C₀-C₆)C(=NOR⁷)R⁹, hétéroaryle, hétéroarylalkyle, arylalkyle, aryle ou hétérocycloalkyle à 3 à 6 chaînons en option substitué ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) ;

Chaque R⁷ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇, hétérocycloalkyle à 3 à 7 chaînons, aryle en C₆-C₁₀, hétéroaryle à 5 ou 6 chaînons, -(aryle en C₇-C₁₀)alkyle et -C(=O)R¹¹ ; R¹¹ est sélectionné parmi un alkyle en C₁-C₆, un cycloalkyle en C₃-C₇ et un aryle en C₆-C₁₀ ;

R⁸, R⁹ et R¹⁰ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, arylalkyle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ;

E est sélectionné dans le groupe consistant en un hydrogène, halogène, -OCF₃, -NO₂, -CF₃, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR¹², -(alkyle en C₀-C₆)NR¹³R¹⁴, -(alkyle en C₀-C₆)C(=NR¹⁵)NR¹³R¹⁴, -(alkyle en C₀-C₆)SR¹², -(alkyle en C₀-C₆)NR¹³C(=O)R¹⁴, -(alkyle en C₀-C₆)NR¹³S(=O)₂R¹⁴, -(alkyle en C₀-C₆)NR¹⁵C(=O)NR¹³R¹⁴, -(alkyle en C₀-C₆)S(=O)₂NR¹³R¹⁴, -(alkyle en C₀-C₆)C(=O)R¹³, -(alkyle en C₀-C₆)C(=O)OR¹³, -(alkyle en C₀-C₆)C(=O)NR¹³R¹⁴, -(alkyle en C₀-C₆)C(=NR¹³)R¹⁴ ou -(alkyle en C₀-C₆)C(=NOR¹³)R¹⁴ ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) en option substitué ;

Chaque R¹² est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇ et -C(=O)R¹⁶ ; R¹⁹ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₇ et aryle en C₆-C₁₀ ; R¹³R¹⁴ et R¹⁵ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, alkylhétéroaryle, arylalkyle, alkylaryle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ;

Quand E représente un -H, les composés dans lesquels X₃ ne représente pas un CH= ou un -CF= sont exclus de l'invention ;

L est indépendamment sélectionné parmi un -(alkyle en C₁-C₅)Q_m- substitué par 1-6T ;

m peut prendre la valeur 0 ou 1 ;

T représente 1 à 6 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, fluor, -OCF₃, -NO₂, -CN, -CF₃, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR¹⁷, -(alkyle en C₀-C₆)NR¹⁸R¹⁹, -(alkyle en C₀-C₆)C(=NR²⁰)NR¹⁸R¹⁹,

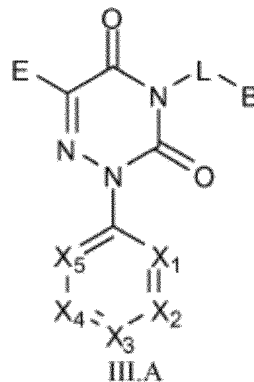
-(alkyle en C₀-C₆)SR¹⁸, -(alkyle en C₀-C₆)NR¹⁸C(=O)R¹⁹, -(alkyle en C₀-C₆)NR¹⁸C(=O)₂R¹⁹, -(alkyle en C₀-C₆)NR¹⁸S(=O)₂R¹⁹, -(alkyle en C₀-C₆)NR²⁰C(=O)NR¹⁸R¹⁹, -(alkyle en C₀-C₆)S(=O)R¹⁸, -(alkyle en C₀-C₆)S(=O)₂R¹⁸, -(alkyle en C₀-C₆)S(=O)₂NR¹⁸R¹⁹, -(alkyle en C₀-C₆)C(=O)R¹⁸, -(alkyle en C₀-C₆)C(=O)OR¹⁷, -(alkyle en C₀-C₆)C(=O)NR¹⁸R¹⁹, -(alkyle en C₀-C₆)C(=NR¹⁸)R¹⁹, -(alkyle en C₀-C₆)C(=NOR¹⁷)R¹⁹, hétéroaryle, hétéroarylalkyle, arylalkyle, alkylaryle, alkylhétéroaryle ou aryle ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)-(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)-(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) ;

Chaque R¹⁷ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇, hétérocycloalkyle à 3 à 7 chaînons, aryle en C₆-C₁₀, hétéroaryle à 5 ou 6 chaînons, -(aryle en C₇-C₁₀)alkyle et -C(=O)R²¹ ; R²¹ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₇ et aryle en C₆-C₁₀ ; R¹⁸, R¹⁹ et R²⁰ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, arylalkyle, alkylaryle, alkylhétéroaryle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ;

Q représente un -C(=O)-, -C(=O)O-, -C(=O)NR²²-, -OC(=O)-, -OC(=O)NR²²-, -NR²²-, -NR²²C(=O)-, -NR²²C(=O)₂-, -NR²²C(=S)-, -S-, -S(=O)- ou -S(=O)₂- ; R²² est sélectionné parmi un alkyle en C₁-C₆ et un aryle en C₆-C₁₀ ;

Tout N peut représenter un N-oxyde.

2. Composé selon la revendication 1 répondant à la formule III.A



et formes stéréoisomères, mélanges de formes stéréoisomères ou sels, hydrates ou solvates pharmaceutiquement acceptables de tels composés, dans lequel :

B est sélectionné parmi un aryle en C₆-C₁₀ en option substitué par un à 5 Y, un cycloalkényle en C₅-C₁₀ en option substitué par un à 7 Y, un groupement hétéroaryle à 5 à 14 chaînons en option substitué par un à 8 Y, ledit groupement hétéroaryle comprenant un, deux ou trois hétéroatomes sélectionnés parmi un N, un O et un S ; Y représente 1 à 5 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, halogène, -OCF₃, -NO₂, -CN, -CF₃, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR¹, -(alkyle en C₀-C₆)NR²R³, -(alkyle en C₀-C₆)C(=NR⁴)NR²R³, -(alkyle en C₀-C₆)SR², -(alkyle en C₀-C₆)NR²C(=O)R³, -(alkyle en C₀-C₆)NR²S(=O)₂R³, -(alkyle en C₀-C₆)NR⁴C(=O)NR²R³, -(alkyle en C₀-C₆)S(=O)R², -(alkyle en C₀-C₆)S(=O)₂R², -(alkyle en C₀-C₆)S(=O)₂NR²R³, -(alkyle en C₀-C₆)C(=O)R², -(alkyle en C₀-C₆)C(=O)OR¹, -(alkyle en C₀-C₆)C(=O)NR²R³, -(alkyle en C₀-C₆)C(=NR²)R³, -(alkyle en C₀-C₆)C(=NOR¹)R³, hétéroaryle, hétéroarylalkyle, arylalkyle, arylalkylaryle ou alkylhétéroaryle en option substitué ; deux substituants étant en option combinés aux atomes impli-

qués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) ;

Chaque R¹ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇, hétérocycloalkyle à 3 à 7 chaînons, aryle en C₆-C₁₀, hétéroaryle à 5 ou 6 chaînons, -(aryle en C₇-C₁₀)alkyle et -C(=O)R⁵ ; R⁵ est sélectionné parmi un hydrogène, un alkyle en C₁-C₆ et un aryle en C₆-C₁₀ ;

R², R³ et R⁴ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, arylalkyle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(hétéroaryle), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ; les substituants R², R³, R⁴ étant en option combinés aux atomes impliqués pour former un hétérocycloalkyle bicyclique ;

X₁, X₂, X₃, X₄ et X₅ sont chacun indépendamment sélectionnés dans le groupe consistant en un -CR⁶= et un -N= ; R⁶ représente 1 à 5 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, halogène, -OCF₃, -CN, -CF₃, -NO₂, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR⁷, -(alkyle en C₀-C₆)NR⁸R⁹, -(alkyle en C₀-C₆)C(=NR¹⁰)NR⁸R⁹, -(alkyle en C₀-C₆)SR⁸, -(alkyle en C₀-C₆)NR⁸C(=O)R⁹, -(alkyle en C₀-C₆)NR⁸S(=O)₂R⁹, -(alkyle en C₀-C₆)NR¹⁰C(=O)NR⁸R⁹, -(alkyle en C₀-C₆)S(=O)R⁸, -(alkyle en C₀-C₆)S(=O)₂R⁸, -(alkyle en C₀-C₆)S(=O)₂NR⁸R⁹, -(alkyle en C₀-C₆)C(=O)R⁸, -(alkyle en C₀-C₆)C(=O)OR⁷, -(alkyle en C₁-C₆)C(=O)NR⁸R⁹, -(alkyle en C₀-C₆)C(=NOR⁷)R⁹, hétéroaryle, hétéroarylalkyle, arylalkyle, aryle ou hétérocycloalkyle à 3 à 6 chaînons en option substitué ; deux substituants étant en option combinés aux atomes impliqués pour former

un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) ;

Chaque R⁷ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇, hétérocycloalkyle à 3 à 7 chaînons, aryle en C₆-C₁₀, hétéroaryle à 5 ou 6 chaînons, -(aryle en C₇-C₁₀)alkyle et -C(=O)R¹¹ ; R¹¹ est sélectionné parmi un alkyle en C₁-C₆, un cycloalkyle en C₃-C₇ et un aryle en C₆-C₁₀ ;

R⁸, R⁹ et R¹⁰ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, arylalkyle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ;

E est sélectionné dans le groupe consistant en un hydrogène, halogène, -OCF₃, -NO₂, -CF₃, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR¹², -(alkyle en C₀-C₆)NR¹³R¹⁴, -(alkyle en C₀-C₆)SR¹², -(alkyle en C₀-C₆)NR¹³C(=O)R¹⁴, -(alkyle en C₀-C₆)NR¹³S(=O)₂R¹⁴, -(alkyle en C₀-C₆)NR¹⁵C(=O)NR¹³R¹⁴, -(alkyle en C₀-C₆)S(=O)R¹³, -(alkyle en C₀-C₆)S(=O)₂R¹³, -(alkyle en C₀-C₆)S(=O)₂NR¹³R¹⁴, -(alkyle en C₀-C₆)C(=O)R¹³, -(alkyle en C₀-C₆)C(=O)OR¹³ ou -(alkyle en C₀-C₆)C(=O)NR¹³R¹⁴ ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) en option substitué ;

Chaque R¹² est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇ et -C(=O)R¹⁶ ;

R¹⁹ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₇ et aryle en C₆-C₁₀ ;

R¹³, R¹⁴ et R¹⁵ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, alkyhétéroaryle, arylalkyle, alkylaryle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en

C_0-C_6 (aryle) ;

Quand E représente un H, les composés dans lesquels X_3 ne représente pas un CH= ou un -CF= sont exclus de l'invention ;

L est indépendamment sélectionné parmi un -(alkyle en C_1-C_5) Q_m - substitué par 1-6T ;
m peut prendre la valeur 0 ou 1 ;

T représente 1 à 6 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, fluor, -OCF₃, -NO₂, -CN, -CF₃, alkyle en C_1-C_6 , cycloalkyle en C_3-C_6 , alkényle en C_2-C_6 , alkynyle en C_2-C_6 , halo(alkyle en C_1-C_6), -(alkyle en C_0-C_6)OR¹⁷, -(alkyle en C_0-C_6)NR¹⁸R¹⁹, -(alkyle en C_0-C_6)C(=NR²⁰)NR¹⁸R¹⁹, -(alkyle en C_0-C_6)SR¹⁸, -(alkyle en C_0-C_6)NR¹⁸C(=O)R¹⁹, -(alkyle en C_0-C_6)NR¹⁸C(=O)₂R¹⁹, -(alkyle en C_0-C_6)NR¹⁸S(=O)₂R¹⁹, -(alkyle en C_0-C_6)NR²⁰C(=O)NR¹⁸R¹⁹, -(alkyle en C_0-C_6)S(=O)R¹⁸, -(alkyle en C_0-C_6)S(=O)₂R¹⁸, -(alkyle en C_0-C_6)S(=O)₂NR¹⁸R¹⁹, -(alkyle en C_0-C_6)C(=O)R¹⁸, -(alkyle en C_0-C_6)C(=O)OR¹⁷, -(alkyle en C_0-C_6)C(=O)NR¹⁸R¹⁹, -(alkyle en C_0-C_6)C(=NR¹⁸)R¹⁹, -(alkyle en C_0-C_6)C(=NOR¹⁷)R¹⁹, hétéroaryle, hétéroarylalkyle, arylalkyle, alkylaryle, alkylhétéroaryle ou aryle ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C_1-C_6 , -O(alkyle en C_0-C_6), -O(cycloalkyle en C_3-C_7), -O(aryle), -O(aryle)-(alkyle en C_0-C_3), -O(hétéroaryle), -O(hétéroaryle)-(alkyle en C_0-C_3), -O(alkyle en C_1-C_3)aryle, -O(alkyle en C_1-C_3)hétéroaryle, -N((alkyle en C_0-C_6)(alkyle en C_0-C_3)aryle) ou -N((alkyle en C_0-C_6)(alkyle en C_0-C_3)hétéroaryle) ;

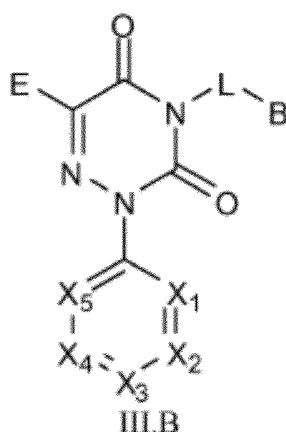
Chaque R¹⁷ est sélectionné parmi un hydrogène, alkyle en C_1-C_6 , alkényle en C_2-C_6 , cycloalkyle en C_3-C_7 , hétérocycloalkyle à 3 à 7 chaînons, aryle en C_6-C_{10} , hétéroaryle à 5 ou 6 chaînons, -(aryle en C_7-C_{10})alkyle et -C(=O)R²¹ ; R²¹ est sélectionné parmi un hydrogène, alkyle en C_1-C_6 , cycloalkyle en C_3-C_7 et aryle en C_6-C_{10} ; R¹⁸, R¹⁹ et R²⁰ représentent chacun indépendamment un hydrogène, alkyle en C_1-C_6 , cycloalkyle en C_3-C_6 , alkényle en C_2-C_6 , alkynyle en C_2-C_6 , -(alkyle en C_1-C_6)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, arylalkyle, alkylaryle, alkylhétéroaryle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C_1-C_6 , -O(alkyle en C_0-C_6), -O(cycloalkyle en C_3-C_7)alkyle, -O(aryle), -O(aryle)(alkyle en C_0-C_3), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C_0-C_3), -N(alkyle en C_0-C_6), -N(alkyle en C_0-C_6)(cycloalkyle en C_3-C_7) ou -N(alkyle en C_0-C_6)(aryle) ;

Q représente un -C(=O)-, -C(=O)O-, -C(=O)NR²²-, -OC(=O)-, -OC(=O)NR²²-, -NR²²-, -NR²²C(=O)-, -NR²²C(=O)₂-, -NR²²C(=S)-, -S-, -S(=O)- ou -S(=O)₂- ;

R²² est sélectionné parmi un alkyle en C_1-C_6 et un aryle en C_6-C_{10} ;

Tout N peut représenter un N-oxyde.

3. Composé selon la revendication 2 qui répond à la formule III.B



et formes stéréoisomères, mélanges de formes stéréoisomères ou sels, hydrates ou solvates pharmaceutiquement acceptables de tels composés, dans lequel :

B est sélectionné parmi un aryle en C₆-C₁₀ en option substitué par un à 5 Y, un cycloalkényle en C₅-C₁₀ en option substitué par un à 7 Y, un groupement hétéroaryle à 5 à 14 chaînons en option substitué par un à 8 Y, ledit groupement hétéroaryle comprenant un, deux ou trois hétéroatomes sélectionnés parmi un N, un O et un S ; Y représente 1 à 5 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, halogène, -OCF₃, -NO₂, -CN, -CF₃, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR¹, -(alkyle en C₀-C₆)NR²R³, -(alkyle en C₀-C₆)C(=NR⁴)NR²R³, -(alkyle en C₀-C₆)SR², -(alkyle en C₀-C₆)NR²C(=O)R³, -(alkyle en C₀-C₆)NR²S(=O)₂R³, -(alkyle en C₀-C₆)NR⁴C(=O)NR²R³, -(alkyle en C₀-C₆)S(=O)R², -(alkyle en C₀-C₆)S(=O)₂R², -(alkyle en C₀-C₆)S(=O)₂NR²R³, -(alkyle en C₀-C₆)C(=O)R², -(alkyle en C₀-C₆)C(=O)OR¹, -(alkyle en C₀-C₆)C(=O)NR²R³, -(alkyle en C₀-C₆)C(=NR²)R³, -(alkyle en C₀-C₆)C(=NOR¹)R³ hétéroaryle, hétéroarylalkyle, arylalkyle, arylalkylaryle ou alkylhétéroaryle en option substitué ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) ;

Chaque R¹ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇, hétérocycloalkyle à 3 à 7 chaînons, aryle en C₆-C₁₀, hétéroaryle à 5 ou 6 chaînons, -(aryle en C₇-C₁₀)alkyle et -C(=O)R⁵ ; R⁵ est sélectionné parmi un hydrogène, un alkyle en C₁-C₆ et un aryle en C₆-C₁₀ ;

R², R³ et R⁴ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, arylalkyle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(hétéroaryle), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ; les substituants R², R³, R⁴ étant en option combinés aux atomes impliqués pour former un hétérocycloalkyle bicyclique ;

X₁, X₂, X₃, X₄ et X₅ sont chacun indépendamment sélectionnés dans le groupe consistant en un -CR⁶= et un -N= ; R⁶ représente 1 à 5 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, halogène, -OCF₃, -CN, -CF₃, -NO₂, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, halo(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR⁷, -(alkyle en C₀-C₆)NR⁸R⁹, -(alkyle en C₀-C₆)SR⁸, -(alkyle en C₀-C₆)NR⁸C(=O)R⁹, -(alkyle en C₀-C₆)NR⁸S(=O)₂R⁹, -(alkyle en C₀-C₆)NR¹⁰C(=O)NR⁸R⁹, -(alkyle en C₀-C₆)S(=O)R⁸, -(alkyle en C₀-C₆)S(=O)₂R⁸, -(alkyle en C₀-C₆)S(=O)₂NR⁸R⁹, -(alkyle en C₀-C₆)C(=O)R⁸, -(alkyle en C₀-C₆)C(=O)OR⁷, -(alkyle en C₁-C₆)C(=O)NR⁸R⁹, hétéroaryle, hétéroarylalkyle, arylalkyle, aryle ou hétérocycloalkyle à 3 à 6 chaînons en option substitué ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) ;

Chaque R⁷ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇, hétérocycloalkyle à 3 à 7 chaînons, aryle en C₆-C₁₀, hétéroaryle à 5 ou 6 chaînons, -(aryle en C₇-C₁₀)alkyle et -C(=O)R¹¹ ; R¹¹ est sélectionné parmi un alkyle en C₁-C₆, un cycloalkyle en C₃-C₇ et un aryle en C₆-C₁₀ ;

R⁸, R⁹ et R¹⁰ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, arylalkyle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ;

E est sélectionné dans le groupe consistant en un hydrogène, halogène, -OCF₃, -CF₃, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, -(alkyle en C₀-C₆)OR¹², -(alkyle en C₀-C₆)NR¹³R¹⁴, -(alkyle en C₀-C₆)S(=O)R¹³, -(alkyle en C₀-C₆)S(=O)₂R¹³, -(alkyle en C₀-C₆)C(=O)R¹³, -(alkyle en C₀-C₆)C(=O)OR¹³ ou -(alkyle en C₀-C₆)C(=O)NR¹³R¹⁴ ; deux substituants étant en option combinés aux atomes impliqués pour former un noyau hétérocycloalkyle, aryle ou hétéroaryle bicyclique ; chaque noyau étant de plus en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇), -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -O(alkyle en C₁-C₃)aryle, -O(alkyle en C₁-C₃)hétéroaryle, -N(alkyle en C₀-C₆)₂, -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)aryle) ou -N((alkyle en C₀-C₆)(alkyle en C₀-C₃)hétéroaryle) en option substitué ;

Chaque R¹² est sélectionné parmi un hydrogène, alkyle en C₁-C₆, alkényle en C₂-C₆, cycloalkyle en C₃-C₇ et

-C(=O)R¹⁶ ; R¹⁶ est sélectionné parmi un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₇ et aryle en C₆-C₁₀ ; R¹³ et R¹⁴ représentent chacun indépendamment un hydrogène, alkyle en C₁-C₆, cycloalkyle en C₃-C₆, alkényle en C₂-C₆, alkynyle en C₂-C₆, -(alkyle en C₁-C₆)halo, hétérocycloalkyle, hétéroaryle, hétéroarylalkyle, alkylhétéroaryle, arylalkyle, alkylaryle ou aryle ; l'un quelconque de ces groupements étant en option substitué par 1 à 5 substituants indépendamment sélectionnés parmi un halogène, -CN, alkyle en C₁-C₆, -O(alkyle en C₀-C₆), -O(cycloalkyle en C₃-C₇)alkyle, -O(aryle), -O(aryle)(alkyle en C₀-C₃), -O(hétéroaryle), -O(hétéroaryle)(alkyle en C₀-C₃), -N(alkyle en C₀-C₆)₂, -N(alkyle en C₀-C₆)(cycloalkyle en C₃-C₇) ou -N(alkyle en C₀-C₆)(aryle) ;

Quand E représente un -H, les composés dans lesquels X₃ ne représente pas un CH= ou un -CF= sont exclus de l'invention ;

L est indépendamment sélectionné parmi un -(alkyle en C₁-C₅)Q_m- substitué par 1-6T ;

m peut prendre la valeur 0 ou 1 ;

T représente 1 à 6 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène et un alkyle en C₁-C₆ ;

Q représente un C(=O)-, -C(=O)-, -C(=O)NR²²-, -OC(=O)-, -OC(=O)NR²²-, -NR²²-, -NR²²C(=O)-, -NR²²C(=O)₂-, -NR²²C(=S)-, -S-, -S(=O)- ou -S(=O)₂- ; R²² est sélectionné parmi un alkyle en C₁-C₆ et un aryle en C₆-C₁₀ ; Tout N peut représenter un N-oxyde.

4. Composé selon les revendications 1 à 3 qui peut exister sous la forme d'isomères optiques, ledit composé étant soit le mélange racémique, soit un isomère optique individuel.

5. Composé selon les revendications 1 à 4, ledit composé étant sélectionné parmi les suivants :

N-(3-(4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)méthanesulfonamide, 4-(4-chlorobenzyl)-2-(méthoxyphényl)-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-cyanobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(naphtalén-2-ylméthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(benzo[c][1,2,5]oxadiazol-5-ylméthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-benzyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(3,5-dioxo-4-(4-(trifluorométhyl)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-((2-(3-acétamidophényl)-3,5-dioxo-2,3-dihydro-1,2,4-triazin-4(5H)-yl)méthyl)benzoate de méthyle, N-(3-(4-(naphtalén-1-ylméthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-méthoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-bromobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-méthylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-isopropylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3,4-diméthylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((6-chloropyridin-3-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chloro-2-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-benzyl-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chloro-3-fluorobenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((6-fluoropyridin-3-yl)méthyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((6-isopropylpyridin-3-yl)méthyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-fluorobenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(4-fluorobenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-méthoxybenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-isopropylbenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-(4-chlorobenzyl)-6-méthyl-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(6-méthyl-4-(naphtalén-2-ylméthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((6-cyclopentylpyridin-3-yl)méthyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3,4-difluorobenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3,4-diméthoxybenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(2,4-difluorobenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-isopropoxybenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthyl-4-(naphtalén-1-ylméthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthyl-4-(4-méthylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-

2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(6-méthyl-4-(4-méthylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(3-méthoxybenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 3-(4-(4-fluorobenzyl)-6-méthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-N-méthylbenzamide, N-(3-(4-(4-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-(4-méthylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-3,5-dioxo-4-(4-(trifluorométhyl)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chloro-3-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-(4-chloro-3-fluorobenzyl)-6-méthoxy-2-(2-méthoxyphényl)-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)méthanesulfonamide, 4-(4-chlorobenzyl)-6-méthoxy-2-(2-méthoxyphényl)-1,2,4-triazin-3,5(2H,4H)-dione, 1-(3-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-3-méthylurée, N-(3-(4-(6-chloropyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(6-méthoxy-4-(4-méthylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)pyridin-3-yl)acétamide, N-(3-(6-méthoxy-4-(6-méthylpyridin-3-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chloro-3-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)pyridin-3-yl)acétamide, 4-(4-chlorobenzyl)-2-(4-fluoro-2-méthoxyphényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(2-(méthoxyméthyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(3-(méthoxyméthyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(2-(trifluorométhoxy)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-2-(2,3-diméthoxyphényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(2-(morpholinométhyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(2-(2-méthoxyéthyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(3-((2-méthyl-1,3-dioxolan-2-yl)méthyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(3-(2-oxopropyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-fluorobenzyl)-2-(2-(diméthylamino)méthyl)phényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-fluorobenzyl)-6-méthoxy-2-(2-(pyrrolidin-1-yl)méthyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, N-(2-fluoro-5-(6-méthoxy-4-(4-méthoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-(4-fluorobenzyl)-2-(4-fluorophényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-fluorobenzyl)-6-méthoxy-2-(2-(piperidin-1-yl)méthyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(2-(((2-méthoxyéthyl)(méthyl)amino)méthyl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, N-(2-fluoro-5-(4-(3-fluoro-4-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 2-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile, 4-(4-chlorobenzyl)-2-(4-fluoro-2-(hydroxyméthyl)phényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, 4-(2,4-dichlorobenzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-benzyl-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-(4-isopropylbenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(biphényl-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-3,5-dioxo-4-(4-(trifluorométhoxy)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-(diméthylamino)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(biphényl-4-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-benzyl-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-(naphtalén-2-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(6-isopropylpyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-(4-méthoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-3,5-dioxo-4-(quinoléin-6-yl)méthyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-isopropoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3,4-diméthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(6-cyclopentylpyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-(4-chlorobenzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-(6-fluoropyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(6-chloro-5-fluoropyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(6-(diméthylamino)pyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3,4-difluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(2,4-difluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-(6-méthoxypyridin-3-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-3,5-dioxo-4-(pyridin-3-yl)méthyl)-4,5-dihydro-

1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-(1H-pyrazol-1-yl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(2,6-difluoro-4-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-3,5-dioxo-4-(quinoléin-3-ylméthyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3-fluoro-4-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3,5-difluoro-4-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(2,3-difluoro-4-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-((1-méthyl-1H-indol-5-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(2-fluoro-4-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-(3,4-difluorobenzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-(2-chloro-4-fluorobenzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chloro-2-fluorobenzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(6-méthoxy-4-(3-méthoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-((2,3-dihydrobenzo[b][1,4]dioxin-2-yl)méthyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 6-méthoxy-2-phényl-4-((2-phénylthiazol-4-yl)méthyl)-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-(4-chloro-2-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 6-méthoxy-2-phényl-4-(3-(trifluorométhoxybenzyl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(2,5-difluorobenzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chloro-3-(trifluorométhoxy)benzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-(2,4-difluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((6-fluoro-2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-(4-fluoro-3-(trifluorométhyl)benzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-benzoylbenzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(4-((6-méthoxy-3,5-dioxo-2-phényl-2,3-dihydro-1,2,4-triazin-4(5H)-yl)méthyl)thiazol-2-yl)acétamide, N-(3-(4-(furo[2,3-b]pyridin-5-ylméthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(benzo[d]oxazol-6-ylméthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-fluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 3-((6-méthoxy-3,5-dioxo-2-phényl-2,3-dihydro-1,2,4-triazin-4(5H)-yl)méthyl)benzonitrile, 4-((6-méthoxy-3,5-dioxo-2-phényl-2,3-dihydro-1,2,4-triazin-4(5H)-yl)méthyl)benzonitrile, N-(3-(4-((7-fluoro-2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-(4-(hydroxyméthyl)benzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-((6-chloro-4H-benzo[d][1,3]dioxin-8-yl)méthyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 6-méthoxy-2-phényl-4-((5-(trifluorométhyl)furan-2-yl)méthyl)-1,2,4-triazin-3,5(2H,4H)-dione, 6-méthoxy-4-((1-méthyl-1H-imidazol-2-yl)méthyl)-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-(2,4-dichlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((6-(diéthylamino)pyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-((5-chlorothiophén-2-yl)méthyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-((5-chlorobenzo[b]thiophén-3-yl)méthyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-(2,4-difluorobenzyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-((2,3-dihydrofuro[2,3-b]pyridin-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((4-fluoro-2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-fluoro-3-(trifluorométhyl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((2-éthylbenzo[d]oxazol-6-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-cyanobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-((2-méthylbenzo[d]oxazol-6-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((6-(diméthylamino)-5-fluoropyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-(diméthylamino)-3-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-((1-méthylindolin-5-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(2-fluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((4-fluoro-2,3-dihydrobenzofuran-7-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((2,2-difluorobenzo[d][1,3]dioxol-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((5-chloropyridin-2-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-((5-isopropylpyridin-2-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(3,5-difluoro-4-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(4-fluoro-3-(trifluorométhoxy)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(3-fluoro-4-(trifluorométhoxy)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-N-méthylacétamide, N-(3-(6-méthoxy-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-N-méthylacétamide, N-(5-(4-((2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-

2-fluorophényl)acétamide, N-(3-(6-méthoxy-4-(4-méthoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-N-méthylacétamide, N-(3-(4-((2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-N-méthylacétamide, N-(3-(4-(4-fluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-N-méthylacétamide, N-(3-(4-(2,4-difluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-N-méthylacétamide, 6-méthoxy-4-phenéthyl-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-2-méthoxyacétamide, N-(3-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-3-méthoxypropanamide, 4-(4-chlorobenzyl)-2-(2-hydroxyphényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(3-(thiazo-1-2-ylamino)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(3-(2-oxopyrrolidin-1-yl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-chlorobenzyl)-6-méthoxy-2-(3-(5-méthyl-1,2,4-oxadiazol-3-yl)phényl)-1,2,4-triazin-3,5(2H,4H)-dione, 6-méthoxy-4-(4-méthoxyphenéthyl)-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-(4-fluorophenéthyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 4-(3,4-diméthoxyphenéthyl)-6-méthoxy-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 6-méthoxy-4-(3-méthoxyphenéthyl)-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, 6-méthoxy-4-(4-méthoxyphenéthyl)-2-phényl-1,2,4-triazin-3,5(2H,4H)-dione, N-(3-(6-méthoxy-4-(3-méthoxyphenéthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-benzyl-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(4-fluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(2,4-difluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(4,5-difluoro-2-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(6-méthoxy-3,5-dioxo-4-phenéthyl-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-((6-chloro-5-fluoropyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(6-méthoxy-4-(3-méthoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(2,4-difluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-4-fluorophényl)acétamide, N-(2-fluoro-5-(6-méthoxy-4-((1-méthylindolin-5-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-((6-fluoro-2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-(diméthylamino)-3-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(3,4-difluoro-2-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, 6-méthoxy-2-phényl-4-(3-phénylpropyl)-1,2,4-triazin-3,5(2H,4H)-dione, 2-(2-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acetonitrile, N-(5-(4-(4-cyanobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(6-méthoxy-4-(2-méthoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-cyano-3-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-((6-cyanopyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-((6-isopropylpyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(6-méthoxy-4-((6-méthoxypyridin-3-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(6-méthoxy-3,5-dioxo-4-(quinoléin-6-yl)méthyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(benzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(4-fluorophenéthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(3-chloro-4-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(3-fluoro-4-(pyrrolidin-1-yl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(3-fluoro-4-morpholinobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(2,6-difluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(6-méthoxy-3,5-dioxo-4-(4-(trifluorométhyl)benzyl)-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(6-méthoxy-4-(3-(méthoxyméthyl)benzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(furo[2,3-b]pyridin-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(2-fluoro-5-(méthoxyméthyl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(4-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(6-méthoxy-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-(1H-pyrazol-1-yl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(3-fluoro-5-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(3-(diméthylamino)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(3-(6-méthoxy-4-((1-méthyl-1H-indazol-6-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-((4-méthyl-3,4-dihydro-2H-benzo[b][1,4]oxazin-7-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-((2,3-dihydroben-

zo[b][1,4]dioxin-6-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(3-(6-méthoxy-4-((1-méthyl-1H-indazol-5-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-4-((2-méthyl-2H-indazol-5-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-((2,3-dihydrobenzofuran-6-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-((2,3-dihydrobenzofuran-4-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(3-(6-méthoxy-4-((5-méthoxypyridin-3-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-cyano-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-méthoxy-3,5-dioxo-4-phenéthyl-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(4-fluoro-2-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-(1H-pyrazol-1-yl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(3,4-difluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(6-méthoxy-4-((1-méthyl-1H-indazol-6-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(3,4-dichlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(3-(diméthylamino)-4-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-chloro-5-(4-(4-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-((2,2-diméthyl-2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, 4-(4-chlorobenzyl)-2-(4-fluorophényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, N-(5-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(3-(4-(benzo[d]isoxazol-6-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chloro-3-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-chloro-5-(4-(4-chloro-3-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-(4-fluoro-2-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(2,4-difluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(4-chloro-2-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(4-chloro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(2,4-difluorophenéthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(3-(4-((5-fluoro-6-méthoxypyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-(4-chloro-2-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-(4-chloro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(3,4-difluoro-5-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-chloro-5-(4-(4-chloro-2-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-(2,4-difluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-(4-chlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chloro-2-(diméthylamino)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-chloro-5-(4-(4-chloro-2-(diméthylamino)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-(3-(diméthylamino)-4-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-benzyl-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-chlorophényl)acétamide, N-(5-(4-(2-(diméthylamino)-4-fluorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-chloro-5-(4-(4-fluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(3-fluoro-4-(trifluorométhyl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chloro-3-(diméthylamino)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-(4-chloro-3-(diméthylamino)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(3,4-dichlorobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-chloro-5-(4-((6-(diméthylamino)pyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chloro-3-(trifluorométhoxy)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-chloro-5-(6-méthoxy-4-(3-méthoxybenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-((2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-chloro-5-(4-(3,4-difluoro-2-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 4-(4-chlorobenzyl)-2-(4-fluorophényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, N-(2-chloro-5-(4-(4-chloro-3-(trifluorométhyl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chloro-3-isopropoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-chloro-5-(6-méthoxy-4-((1-méthyl-1H-indazol-6-yl)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chloro-3-isopropylbenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophé-

nyl)acétamide, N-(2-fluoro-5-(4-(3-fluoro-4-morpholinobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(4-chloro-3-hydroxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-((3,3-diméthyl-2,3-dihydrobenzofuran-5-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-((3,3-diméthyl-2,3-dihydrobenzofuran-6-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(3-fluoro-4-isopropylbenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chloro-3-(diméthylamino)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-4-fluorophényl)acétamide, N-(5-(4-(4-cyano-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, 4-(3,4-dichlorobenzyl)-2-(4-fluorophényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, N-(5-(4-(4-chloro-2-fluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, 4-(4-chloro-2-fluoro-3-méthoxybenzyl)-2-(4-fluorophényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, N-(5-(4-(4-bromo-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(4-isopropyl-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 2-(4-fluorophényl)-4-(4-isopropyl-3-méthoxybenzyl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, N-(5-(4-(4-chloro-3-(pyrrolidin-1-yl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(4-fluoro-3-(trifluorométhyl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-(2,4-difluoro-3-isopropoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-((5,6-dichloropyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(4-cyano-2-fluoro-3-méthoxybenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, 4-((5,6-dichloropyridin-3-yl)méthyl)-2-(4-fluorophényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, N-(5-(4-(4-chloro-3-morpholinobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(5-(4-(3-(diméthylamino)-4-(trifluorométhyl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, 4-(4-chloro-3-méthoxybenzyl)-2-(4-fluorophényl)-6-méthoxy-1,2,4-triazin-3,5(2H,4H)-dione, N-(2-fluoro-5-(4-(3-fluoro-4-(trifluorométhyl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)-N-méthylacétamide, N-(5-(4-(4-chloro-3-(méthoxyméthyl)benzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(2-fluoro-5-(4-(4-fluoro-3-morpholinobenzyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(5-(4-((6-chloro-5-isopropylpyridin-3-yl)méthyl)-6-méthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)-2-fluorophényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-isopropyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-éthyl-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-butyl-4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 2-(6-bromo-4-(4-fluorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)benzonitrile, N-(3-(4-(4-chlorobenzyl)-6-(2-(diméthylamino)éthoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-(2-méthoxyéthoxy)-4-(4-nitrobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-(2-méthoxyéthoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-(méthylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-éthoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-bromo-4-(4-méthylbenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-isopropoxy-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-bromo-4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-(diméthylamino)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-(hydroxyméthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-((diméthylamino)méthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(6-((1H-pyrazol-1-yl)méthyl)-4-(4-chlorobenzyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-(méthoxyméthyl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, 2-(3-acétamidophényl)-4-(4-chlorobenzyl)-3,5-dioxo-2,3,4,5-tétrahydro-1,2,4-triazin-6-carboxylate de méthyle, 2-(3-acétamidophényl)-4-(4-chlorobenzyl)-3,5-dioxo-2,3,4,5-tétrahydro-1,2,4-triazin-6-carboxylate d'isopropyle, N-(3-(4-(4-chlorobenzyl)-6-(3-méthyl-1,2,4-oxadiazol-5-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-(5-méthyl-1,2,4-oxadiazol-3-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(3-(4-(4-chlorobenzyl)-6-(1,2,4-oxadiazol-5-yl)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide, N-(2-fluoro-5-(4-(4-fluorobenzyl)-6-(2-méthoxyéthoxy)-3,5-dioxo-4,5-dihydro-1,2,4-triazin-2(3H)-yl)phényl)acétamide.

6. Composition pharmaceutique comprenant une quantité thérapeutiquement efficace d'un composé selon les revendications 1 à 5 et un véhicule et/ou un excipient pharmaceutiquement acceptable.

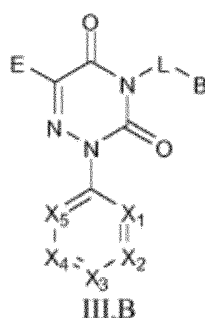
7. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention chez un

mammifère, y compris un être humain, d'une maladie dont le traitement ou la prévention est influencé ou facilité par l'effet neuromodulateur de modulateurs allostériques positifs (stimulateurs) du récepteur GABA_B.

- 5 8. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies du système nerveux central sélectionnées dans le groupe consistant en des troubles anxieux : agoraphobie, trouble anxieux généralisé (TAG), trouble obsessionnel compulsif (TOC), trouble panique, syndrome de stress post-traumatique (SSPT), phobie sociale, autres phobies, trouble anxieux induit par une substance.
- 10 9. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies du système nerveux central sélectionnées dans le groupe consistant en des maladies infantiles : trouble de déficit de l'attention/hyperactivité.
- 15 10. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies du système nerveux central sélectionnées dans le groupe consistant en des troubles de la conduite alimentaire (anorexie mentale, boulimie).
- 20 11. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies du système nerveux central sélectionnées dans le groupe consistant en des troubles de l'humeur : maladies à forme bipolaire (I & II), trouble cyclothymique, dépression, trouble dysthymique, trouble dépressif majeur, trouble de l'humeur induit par une substance.
- 25 12. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies du système nerveux central sélectionnées dans le groupe consistant en des maladies psychotiques : schizophrénie, trouble délirant, trouble schizoaffectif, trouble schizophréniforme, trouble psychotique induit par une substance.
- 30 13. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies du système nerveux central sélectionnées dans le groupe consistant en des troubles cognitifs : délire, délire persistant induit par une substance, démence, complexe démentiel associé au VIH, démence de la maladie de Huntington, démence de la maladie de Parkinson, démence de type Alzheimer, démence persistante induite par une substance, déclin cognitif léger.
- 35 14. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies du système nerveux central sélectionnées dans le groupe consistant en des troubles de la personnalité : trouble de la personnalité obsessionnelle-compulsive, schizoïdie, trouble schizotypique.
- 40 15. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies du système nerveux central sélectionnées dans le groupe consistant en des troubles liés à l'usage d'une substance : usage abusif d'alcool, alcool-dépendance, sevrage éthylique, délire du syndrome de sevrage éthylique, trouble psychotique d'origine éthylique, dépendance aux amphétamines, sevrage des amphétamines, dépendance à la cocaïne, sevrage de la cocaïne, dépendance à la nicotine, sevrage de la nicotine, dépendance aux opioïdes, sevrage des opioïdes.
- 45 16. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de maladies inflammatoires du système nerveux central sélectionnées dans le groupe consistant en une forme de sclérose en plaques comme la forme bénigne de sclérose en plaques, la forme récurrente-rémittente de sclérose en plaques, la forme progressive secondaire de sclérose en plaques, la forme progressive primaire de sclérose en plaques et la forme progressive récurrente de sclérose en plaques.
- 50 17. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de la douleur inflammatoire ou neuropathique et de la douleur postopératoire.
18. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de l'épilepsie et des tremblements.
- 55 19. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de l'incontinence urinaire d'urgence mictionnelle.
20. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de per-

turbations gastro-intestinales sélectionnées dans le groupe consistant en le reflux gastro-oesophagien, le relâchement transitoire du sphincter oesophagien inférieur, la dyspepsie, les troubles gastro-intestinaux fonctionnels et le syndrome du côlon irritable.

21. Composé/composition selon les revendications 1 à 6 pour l'utilisation dans le traitement ou la prévention de la spasticité et de la rigidité des muscles squelettique, d'une atteinte rachidienne et de l'infirmité motrice cérébrale.
22. Utilisation d'un composé selon les revendications 1 à 5 pour la préparation de traceurs pour l'imagerie des récepteurs au GABA_B.
23. Composé/composition selon les revendications 1 à 6 combiné à un agoniste du récepteur GABA_B pour l'utilisation dans le traitement ou la prévention chez un mammifère, y compris un être humain, d'une maladie telle que définie à l'une quelconque des revendications 7 à 11 dont le traitement ou la prévention est influencé ou facilité par l'effet neuromodulateur de modulateurs positifs allostériques du récepteur GABA_B.
24. Procédé de production d'un composé selon la revendication 3 répondant à la formule III.B



et formes stéréoisomères, mélanges de formes stéréoisomères ou sels, hydrates ou solvates pharmaceutiquement acceptables de tels composés, dans lequel :

B est sélectionné parmi un aryle en C₆-C₁₀ en option substitué par un à 5 Y ou un groupement hétéroaryle à 5 à 14 chaînons en option substitué par un à 8 Y, ledit groupement hétéroaryle comprenant un, deux ou trois hétéroatomes sélectionnés parmi un N, un O et un S ;

Y représente 1 à 5 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, halogène, -OCF₃, -NO₂, -CN, -CF₃ et alkyle en C₁-C₆, cycloalkyle en C₃-C₆, halo-(alkyle en C₁-C₆), -(alkyle en C₀-C₆)OR¹, -(alkyle en C₀-C₆)NR²R³ ou -(alkyle en C₀-C₆)SR² en option substitué ;

Chaque R¹ est sélectionné parmi un hydrogène et un alkyle en C₁-C₆ ;

R² et R³ représentent chacun indépendamment un alkyle en C₁-C₆ ou un cycloalkyle en C₃-C₆ ;

X₁, X₂, X₃, X₄ et X₅ sont chacun indépendamment sélectionnés dans le groupe consistant en un -CR⁶= et un -N= ;

R⁶ représente 1 à 5 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène, halogène, -OCF₃, -CN, -CF₃, -NO₂ et alkyle en C₁-C₆, cycloalkyle en C₃-C₆, -(alkyle en C₀-C₆)OR⁷, -(alkyle en C₀-C₆)NR⁸R⁹, -(alkyle en C₀-C₆)SR⁸, -(alkyle en C₀-C₆)NR⁸C(O)R⁹, -(alkyle en C₀-C₆)NR⁸C(=O)₂R⁹, -(alkyle en C₀-C₆)NR⁸S(=O)₂R⁹, -(alkyle en C₀-C₆)NR¹⁰C(=O)NR⁸R⁹, -(alkyle en C₀-C₆)S(=O)₂R⁸, -(alkyle en C₀-C₆)S(=O)₂NR⁸R⁹, -(alkyle en C₀-C₆)C(=O)R⁸, -(alkyle en C₀-C₆)C(=O)OR⁷ ou -(alkyle en C₁-C₆)C(=O)NR⁸R⁹ en option substitué ;

R⁷, R⁸ et R⁹ sont chacun indépendamment sélectionnés parmi un hydrogène, un alkyle en C₁-C₆ et un cycloalkyle en C₃-C₇ ;

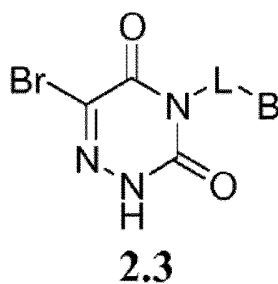
E est sélectionné dans le groupe consistant en un alkyle en C₁-C₆, cycloalkyle en C₃-C₆, -(alkyle en C₀-C₆)OR¹² et -(alkyle en C₀-C₆)NR¹³R¹⁴ ;

R¹², R¹³ et R¹⁴ sont chacun indépendamment sélectionnés parmi un hydrogène, alkyle en C₁-C₆, un alkényle en C₂-C₆ et un cycloalkyle en C₃-C₇ ;

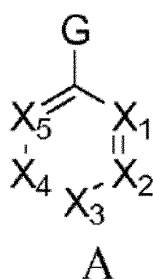
L est indépendamment sélectionné parmi un -(alkyle en C₁-C₅) substitué par 1-6 T ;

T représente 1 à 6 substituants indépendamment sélectionnés dans le groupe consistant en un hydrogène et un alkyle en C₁-C₆ ;

qui comprend la mise en réaction d'un composé de formule 2.3



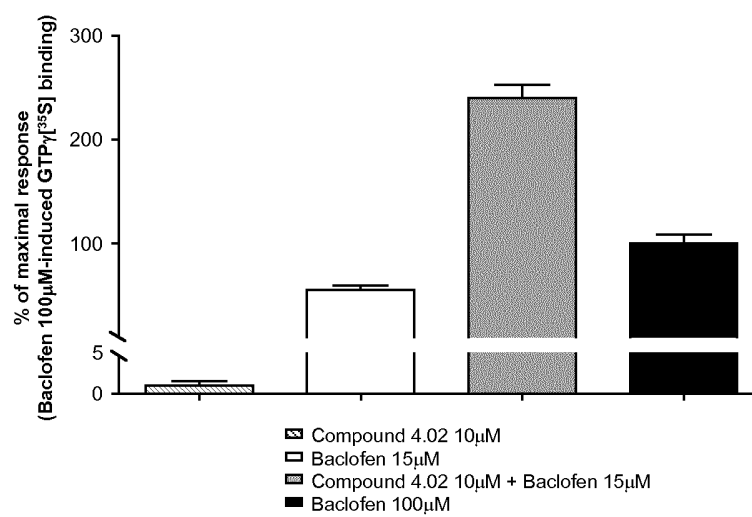
avec l'acide ou l'ester aryl- ou hétéroarylboronique de formula A en présence de systèmes catalytiques au cuivre et d'une base



dans lequel :

G est un acide boronique, un boronate (p. ex. boronate diméthylque, boronate pinacolique et autres composés semblables),

Le produit de réaction obtenu étant ensuite soumis à un déplacement nucléophile du brome par divers groupements pour produire le composé de formule III.B.

FIGURE**Figure A**

REFERENCES CITED IN THE DESCRIPTION

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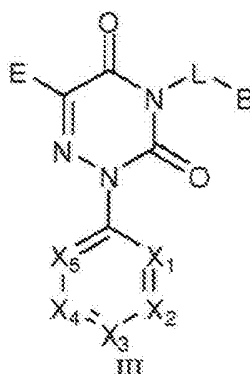
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Szabadalmi igénypontok

1. III képletű vegyületek



és az ilyen vegyületek sztereoiszomer alakjai, sztereoiszomer alakjainak keverékei vagy gyógyszerészetileg elfogadható sói, hidrátjai vagy szolvátjai, ahol:

B jelentése adott esetben 1-5 Y csoporttal helyettesített C_5 - C_{10} aril, adott esetben 1-7 Y csoporttal helyettesített C_5 - C_{10} cikloalkenil, adott esetben 1-8 Y csoporttal helyettesített 5-14 tagú heteroarilcsoport, ahol a heteroarilcsoport egy, kettő vagy három a N, O, S közül választott, heteroatomot tartalmaz;

Y jelentése az alábbiakból álló csoportból egymástól függetlenül választott 1-5 szubsztituens: hidrogén, halogén, $-OCF_3$, $-NO_2$, $-CN$, $-CF_3$, adott esetben szubsztituált- $-(C_1-C_6)$ alkil, $-(C_3-C_6)$ cikloalkil, $-(C_2-C_6)$ alkenil, $-(C_2-C_6)$ alkinil, halogén- $-(C_1-C_6)$ alkil, $-(C_0-C_6)$ alkil OR^1 , $-(C_0-C_6)$ alkil NR^2R^3 , $-(C_0-C_6)$ alkil $-C(=NR^4)NR^2R^3$, $-(C_0-C_6)$ alkil SR^2 , $-(C_0-C_6)$ alkil $NR^1C(=O)R^3$, $-(C_0-C_6)$ alkil $NR^2S(=O)_2R^3$, $-(C_0-C_6)$ alkil $NR^1C(=O)NR^2R^3$, $-(C_0-C_6)$ alkil $S(=O)R^2$, $-(C_0-C_6)$ alkil $S(=O)_2R^2$, $-(C_0-C_6)$ alkil $S(=O)_2NR^2R^3$, $-(C_0-C_6)$ alkil $-C(=O)R^2$, $-(C_0-C_6)$ alkil $C(=O)OR^1$, $-(C_0-C_6)$ alkil $C(=O)NR^2R^3$, $-(C_0-C_6)$ alkil $C(=NR^2)R^3$, vagy $-(C_0-C_6)$ alkil $C(=NOR^3)R^3$, heteroaril, heteroarilalkil, arilalkil, arilalkilaril, alkilheteroaril; ahol adott esetben két szubsztituens a közbúlsó atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol minden gyűrű adott esetben tovább szubsztituált 1-5 független halogén, $-CN$, $-(C_1-C_6)$ alkil, $-O(C_0-C_6)$ alkil, $-O(C_3-C_7)$ cikloalkil, $-O(aril)$, $-O(aril)-(C_0-C_3)$ alkil, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)$ alkil, $-O(C_1-C_3)$ alkilaril, $-O(C_1-C_3)$ alkilheteroaril, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)-((C_0-C_3)alkilaril)$ vagy $-N((C_0-C_6)alkil)((C_0-C_3)alkilheteroaril)$ csoporttal;

R^1 jelentését minden esetben az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)$ alkil, $-(C_2-C_6)$ alkenil, $-(C_3-C_7)$ cikloalkil, 3-7 tagú heterocikloalkil, $-(C_5-C_{10})$ aril, 5 vagy 6 tagú heteroaril, $-(C_7-C_{10})$ arilalkil, $-C(=O)R^5$;

R^5 jelentését az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)$ alkil, és $-(C_5-C_{10})$ aril;

R^2 , R^3 és R^4 mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)$ alkil, $-(C_3-C_6)$ cikloalkil, $-(C_2-C_6)$ alkenil, $-(C_2-C_6)$ alkinil, $-(C_1-C_6)$ alkilhalogén, heterocikloalkil,

heteroaril, heteroarilalkil, arilalkil vagy aril; amelyek közül bármelyik adott esetben szubsztituált 1-5 egymástól független halogén, -CN, -(C₁-C₆)alkil, -O(C₀-C₆)alkil, -O(C₃-C₇)cikloalkilalkil, -O(aril), -O(heteroaril), -N((C₀-C₆)alkil)₂, -N((C₀-C₆)alkil)((C₃-C₇)cikloalkil) vagy -N((C₀-C₆)alkil)(aril) szubsztituenssel; ahol adott esetben R², R³, R⁴ szubsztituensek a közbülső atomokkal együttesen biciklusos heterocikloalkil csoportot alkotnak;

X₁, X₂, X₃, X₄ és X₅ mindegyikének jelentését egymástól függetlenül az alábbiakból álló csoportból választjuk: -CR⁶=, -N=;

R⁶ jelentése az alábbiakból álló csoportból egymástól függetlenül választott 1-5 szubsztituens: hidrogén, halogén, -OCF₃, -CN, -CF₃, -NO₂, adott esetben szubsztituált-(C₁-C₆)alkil, -(C₃-C₆)cikloalkil, -(C₂-C₆)alkenil, -(C₂-C₆)alkinil, halogén-(C₁-C₆)alkil, -(C₀-C₆)alkilOR⁷, -(C₀-C₆)alkilNR⁸R⁹, -(C₀-C₆)alkil-C(=NR¹⁰)NR⁸R⁹, -(C₀-C₆)alkilSR⁸, -(C₀-C₆)alkilNR⁸C(=O)R⁹, -(C₀-C₆)alkilNR⁸S(=O)₂R⁹, -(C₀-C₆)alkilNR¹⁰C(=O)NR⁸R⁹, -(C₀-C₆)alkilS(=O)R⁸, -(C₀-C₆)alkilS(=O)₂R⁸, -(C₀-C₆)alkilS(=O)₂NR⁸R⁹, -(C₀-C₆)alkil-C(=O)R⁸, -(C₀-C₆)alkilC(=O)OR⁷, -(C₁-C₆)alkilC(=O)NR⁸R⁹, vagy -(C₀-C₆)alkil-C(=NOR⁷)R⁹, heteroaril, heteroarilalkil, arilalkil, aril, 3-6 heterocikloalkil; ahol adott esetben két szubsztituens a közbülső atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol mindegyik gyűrű adott esetben tovább szubsztituált 1-5 egymástól független halogén, -CN, -(C₁-C₆)alkil, -O(C₀-C₆)alkil, -O(C₃-C₇)cikloalkil, -O(aril), -O(aril)-(C₀-C₃)alkil, -O(heteroaril), -O(heteroaril)-(C₀-C₃)alkil, -O(C₁-C₃)alkilaril, -O(C₁-C₃)alkilheteroaril, -N((C₀-C₆)alkil)₂, -N((C₀-C₆)alkil)((C₀-C₃)alkilaril) vagy -N((C₀-C₆)alkil)((C₀-C₃)alkilheteroaril) csoporttal;

R⁷ jelentését minden esetben az alábbiak közül választjuk hidrogén, -(C₁-C₆)alkil, -(C₂-C₆)alkenil, -(C₃-C₇)cikloalkil, 3-7 tagú heterocikloalkil, -(C₆-C₁₀)aril, 5 vagy 6 tagú heteroaril, -(C₇-C₁₀)arilalkil, -C(=O)R¹¹;

R¹¹ jelentését az alábbiak közül választjuk: -(C₁-C₆)alkil, -(C₃-C₇)cikloalkil és -(C₆-C₁₀)aril;

R⁸, R⁹ és R¹⁰ mindegyikének jelentése egymástól függetlenül hidrogén, -(C₁-C₆)alkil, -(C₃-C₆)cikloalkil, -(C₂-C₆)alkenil, -(C₂-C₆)alkinil, -(C₁-C₆)alkilhalogén, heterocikloalkil, heteroaril, heteroarilalkil, arilalkil vagy aril; amelyek bármelyike adott esetben szubsztituált 1-5 egymástól független halogén, -CN, -(C₁-C₆)alkil, -O(C₀-C₆)alkil, -O(C₃-C₇)cikloalkilalkil, -O(aril), -O(aril)-(C₀-C₃)alkil, -O(heteroaril), -O(heteroaril)-(C₀-C₃)alkil, -N((C₀-C₆)alkil)₂, -N((C₀-C₆)alkil)((C₃-C₇)cikloalkil) vagy -N((C₀-C₆)alkil)(aril) szubsztituenssel;

E jelentését az alábbiakból álló csoportból választjuk: hidrogén, halogén, -OCF₃, -NO₂, -CF₃, -(C₁-C₆)alkil, -(C₃-C₆)cikloalkil, -(C₂-C₆)alkenil, -(C₂-C₆)alkinil, halogén-(C₁-C₆)alkil, -(C₀-C₆)alkilOR¹², -(C₀-C₆)alkilNR¹³R¹⁴, -(C₀-C₆)alkil-C(=NR¹⁵)NR¹³R¹⁴, -(C₀-C₆)alkilSR¹², -(C₀-C₆)alkilNR¹³C(=O)R¹⁴, -(C₀-C₆)alkilNR¹³S(=O)₂R¹⁴, -(C₀-C₆)alkilNR¹⁵C(=O)NR¹³R¹⁴,

$-(C_0-C_6)\text{alkilS}(=O)R^{13}$, $-(C_0-C_6)\text{alkilS}(=O)_2R^{13}$, $-(C_0-C_6)\text{alkilS}(=O)_2NR^{13}R^{14}$, $-(C_0-C_6)\text{alkilC}(=O)R^{13}$, $-(C_0-C_6)\text{alkilC}(=O)OR^{13}$, $-(C_0-C_6)\text{alkilC}(=O)NR^{13}R^{14}$, $-(C_0-C_6)\text{alkilC}(=NR^{13})R^{14}$, vagy $-(C_0-C_6)\text{alkilC}(=NOR^{13})R^{14}$; ahol adott esetben két szubsztituens a közbülső atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol minden gyűrű adott esetben tovább szubsztituált 1-5 egymástól független halogén, -CN, adott esetben szubsztituált $-(C_1-C_6)\text{alkil}$, $-O(C_0-C_6)\text{alkil}$, $-O(C_3-C_7)\text{cikloalkil}$, $-O(\text{aryl})$, $-O(\text{aryl})-(C_0-C_3)\text{alkil}$, $-O(\text{heteroaril})$, $-O(\text{heteroaril})-(C_0-C_3)\text{alkil}$, $-O(C_1-C_3)\text{alkilaril}$, $-O(C_1-C_3)\text{alkilheteroaril}$, $-N((C_0-C_6)\text{alkil})_2$, $-N((C_0-C_6)\text{alkil})((C_0-C_3)\text{alkilaril})$ vagy $-N((C_0-C_6)\text{alkil})-((C_0-C_3)\text{alkilheteroaril})$ csoporttal;

R^{12} jelentését minden esetben az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_2-C_6)\text{alkenil}$, $-(C_3-C_7)\text{cikloalkil}$, $-C(=O)R^{16}$;

R^{15} jelentését az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_3-C_7)\text{cikloalkil}$ és $-(C_6-C_{10})\text{aryl}$;

R^{13} , R^{14} és R^{15} mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_3-C_6)\text{cikloalkil}$, $-(C_2-C_6)\text{alkenil}$, $-(C_2-C_6)\text{alkinil}$, $-(C_1-C_6)\text{alkilhalogén}$, heterocikloalkil, heteroaril, heteroarilalkil, alkilheteroaril, arilalkil, alkilaril vagy aril; amelyek közül bármelyik adott esetben szubsztituált 1-5 egymástól független halogén, -CN, $-(C_1-C_6)\text{alkil}$, $-O(C_0-C_6)\text{alkil}$, $-O(C_3-C_7)\text{cikloalkilalkil}$, $-O(\text{aryl})$, $-O(\text{aryl})-(C_0-C_3)\text{alkil}$, $-O(\text{heteroaril})$, $-O(\text{heteroaril})-(C_0-C_3)\text{alkil}$, $-N((C_0-C_6)\text{alkil})_2$, $-N((C_0-C_6)\text{alkil})((C_3-C_7)\text{cikloalkil})$ vagy $-N((C_0-C_6)\text{alkil})(\text{aryl})$ szubsztituenssel;

Amikor E jelentése -H, akkor azok a vegyületek, amelyeknél X_3 jelentése nem $CH=$ vagy $-CF=$, nem képezik a találmány tárgyát;

L jelentését egymástól függetlenül 1-6 T szubsztituált $-((C_1-C_5)\text{alkil}-Q_m)-$ csoportból választjuk;

m értéke 0 vagy 1;

T jelentése 1-6 szubsztituens, amelyet egymástól függetlenül az alábbiakból álló csoportból választunk: hidrogén, fluor, $-OCF_3$, $-NO_2$, -CN, $-CF_3$, $-(C_1-C_6)\text{alkil}$, $-(C_3-C_6)\text{cikloalkil}$, $-(C_2-C_6)\text{alkenil}$, $-(C_2-C_6)\text{alkinil}$, halogén- $-(C_1-C_6)\text{alkil}$, $-(C_0-C_6)\text{alkilOR}^{17}$, $-(C_0-C_6)\text{alkilNR}^{18}R^{19}$, $-(C_0-C_6)\text{alkilC}(=NR^{20})NR^{18}R^{19}$, $-(C_0-C_6)\text{alkilSR}^{16}$, $-(C_0-C_6)\text{alkilNR}^{18}C(=O)R^{19}$, $-(C_0-C_6)\text{alkilNR}^{18}C(=O)_2R^{19}$, $-(C_0-C_6)\text{alkilNR}^{18}S(=O)_2R^{19}$, $-(C_0-C_6)\text{alkilNR}^{20}C(=O)NR^{18}R^{19}$, $-(C_0-C_6)\text{alkilS}(=O)R^{18}$, $-(C_0-C_6)\text{alkilS}(=O)_2R^{18}$, $-(C_0-C_6)\text{alkilS}(=O)_2NR^{18}R^{19}$, $-(C_0-C_6)\text{alkilC}(=O)R^{18}$, $-(C_0-C_6)\text{alkilC}(=O)OR^{17}$, $-(C_0-C_6)\text{alkilC}(=O)NR^{18}R^{19}$, $-(C_0-C_6)\text{alkilC}(=NR^{18})R^{19}$, vagy $-(C_0-C_6)\text{alkilC}(=NOR^{17})R^{19}$, heteroaril, heteroarilalkil, arilalkil, alkilaril, alkilheteroaril, aril; ahol adott esetben két szubsztituens a közbülső atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol mindegyik gyűrű adott esetben tovább szubsztituált 1-5 egymástól független halogén, -CN, $-(C_1-C_6)\text{alkil}$, $-O(C_0-C_6)\text{alkil}$, $-O(C_3-C_7)\text{cikloalkil}$, $-O(\text{aryl})$, $-O(\text{aryl})-(C_0-C_3)\text{alkil}$, $-O(\text{heteroaril})$,

-O(heteroaril)-(C₀-C₃)alkil, -O(C₁-C₃)alkilaril, -O(C₁-C₃)alkilheteroaril, -N((C₀-C₆)alkil)((C₀-C₃)alkilaril)) vagy -N((C₀-C₆)alkil)((C₀-C₃)alkilheteroaril)) csoporttal;

R¹⁷ jelentését minden esetben az alábbiak közül választjuk: hidrogén, -(C₁-C₆)alkil, -(C₂-C₆)alkenil, -(C₃-C₇)cikloalkil, 3-7 tagú heterocikloalkil, -(C₆-C₁₀)aril, 5-6 tagú heteroaril, -(C₇-C₁₀)arilalkil, -C(=O)R²¹;

R²¹ jelentése hidrogén, -(C₁-C₆)alkil, -(C₃-C₇)cikloalkil, vagy -(C₆-C₁₀)aril;

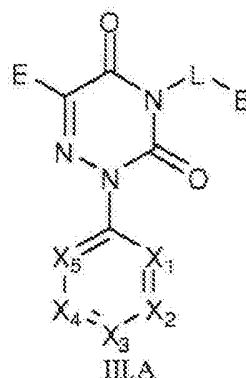
R¹⁸, R¹⁹ és R²⁰ mindegyikének jelentése egymástól függetlenül hidrogén, -(C₁-C₆)alkil, -(C₃-C₆)cikloalkil, -(C₂-C₆)alkenil, -(C₂-C₆)alkinil, -(C₁-C₆)alkilhalogén, heterocikloalkil, heteroaril, heteroarilalkil, arilalkil alkilaril, alkilheteroaril, aril; amelyek közül bármelyik adott esetben szubsztituált 1-5 egymástól független halogén, -CN, -(C₁-C₆)alkil, -O(C₀-C₆)alkil, -O(C₃-C₇)cikloalkilalkil, -O(aril), -O(aril)-(C₀-C₃)alkil, -O(heteroaril), -O(heteroaril)-(C₀-C₃)alkil, -N((C₀-C₆)alkil)₂, -N((C₀-C₆)alkil)((C₃-C₇)cikloalkil) vagy -N((C₀-C₆)alkil)(aril) szubsztituenssel;

Q jelentése -C(=O)-, -C(=O)O-, -C(=O)NR²²-, -OC(=O)-, -OC(=O)NR²²-, -NR²²-, -NR²²C(=O)-, -NR²²C(=O)₂-, -NR²²C(=S)-, -S-, -S(=O)-, vagy -S(=O)₂-;

R²² jelentése -(C₁-C₆)alkil, vagy -(C₆-C₁₀)aril;

Bármely N N-oxidot képezhet.

2. Az 1. igénypont szerinti III.A képletű vegyületek



és az ilyen vegyületek sztereoizomer alakjai, sztereoizomer alakjainak keverékei vagy gyógyszerészetileg elfogadható sói, hidrátjai vagy szolvátjai,

ahol:

B jelentése adott esetben 1-5 Y csoporttal helyettesített C₆-C₁₀ aril, adott esetben 1-7 Y csoporttal helyettesített C₅-C₁₀ cikloalkenil, adott esetben 1-8 Y csoporttal helyettesített 5-14 tagú heteroarilcsoport, ahol a heteroarilcsoport egy, kettő vagy három a N, O, S közül választott, heteroatomot tartalmaz;

Y jelentése az alábbiakból álló csoportból egymástól függetlenül választott 1-5 szubsztituens: hidrogén, halogén, -OCF₃, -NO₂, -CN, -CF₃, adott esetben szubsztituált-(C₁-C₆)alkil, -(C₃-C₆)cikloalkil, -(C₂-C₆)alkenil, -(C₂-C₆)alkinil, halogén-(C₁-C₆)alkil, -(C₀-C₆)alkilOR¹, -(C₀-C₆)alkilNR²R³, -(C₀-C₆)alkil-C(=NR⁴)NR²R³, -(C₀-C₆)alkilSR²,

$-(C_0-C_6)alkilNR^2C(=O)R^3$, $-(C_0-C_6)alkilNR^2S(=O)_2R^3$, $-(C_0-C_6)alkilNR^4C(=O)NR^2R^3$,
 $-(C_0-C_6)alkilS(=O)R^2$, $-(C_0-C_6)alkilS(=O)_2R^2$, $-(C_0-C_6)alkilS(=O)_2NR^2R^3$, $-(C_0-C_6)alkil-$
 $C(=O)R^2$, $-(C_0-C_6)alkilC(=O)OR^1$, $-(C_0-C_6)alkilC(=O)NR^2R^3$, $-(C_0-C_6)alkilC(=NR^3)R^3$,
 vagy $-(C_0-C_6)alkilC(=NOR^1)R^3$, heteroaril, heteroarilalkil, arilalkil, arilalkilaril,
 alkilheteroaril; ahol adott esetben két szubsztituens a közbülső atomokkal együttesen
 biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol minden gyűrű adott
 esetben tovább szubsztituált 1-5 egymástól független halogén, -CN, $-(C_1-C_6)alkil$, $-O(C_0-$
 $C_6)alkil$, $-O(C_3-C_7)cikloalkil$, $-O(aril)$, $-O(aril)-(C_0-C_3)alkil$, $-O(heteroaril)$, $-O(heteroaril)-$
 $-(C_0-C_3)alkil$, $-O(C_1-C_3)alkilaril$, $-O(C_1-C_3)alkilheteroaril$, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)al-$
 $kil)-((C_0-C_3)alkilaril)$ vagy $-N((C_0-C_6)alkil)((C_0-C_3)alkilheteroaril)$ csoporttal;

R^1 jelentését minden esetben az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)alkil$, $-(C_2-$
 $C_6)alkenil$, $-(C_3-C_7)cikloalkil$, 3-7 tagú heterocikloalkil, $-(C_5-C_{10})aril$, 5 vagy 6 tagú
 heteroaril, $-(C_7-C_{10})arilalkil$, $-C(=O)R^5$;

R^5 jelentését az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)alkil$, és $-(C_6-C_{10})aril$;

R^2 , R^3 és R^4 mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)alkil$,
 $-(C_3-C_6)cikloalkil$, $-(C_2-C_6)alkenil$, $-(C_2-C_6)alkinil$, $-(C_1-C_6)alkilhalogén$, heterocikloalkil,
 heteroaril, heteroarilalkil, arilalkil vagy aril; amelyek közül bármelyik adott esetben
 szubsztituált 1-5 egymástól független halogén, -CN, $-(C_1-C_6)alkil$, $-O(C_0-C_6)alkil$, $-O(C_3-$
 $C_7)cikloalkilalkil$, $-O(aril)$, $-O(heteroaril)$, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)((C_3-C_7)-$
 $cikloalkil)$ vagy $-N((C_0-C_6)alkil)(aril)$ szubsztituenssel; ahol adott esetben R^2 , R^3 , R^4
 szubsztituensek a közbülső atomokkal együttesen biciklusos heterocikloalkil csoportot
 alkotnak;

X_1 , X_2 , X_3 , X_4 és X_5 mindegyikének jelentését egymástól függetlenül az alábbiakból álló
 csoportból választjuk: $-CR^6=$, $-N=$;

R^6 jelentése az alábbiakból álló csoportból egymástól függetlenül választott 1-5
 szubsztituens: hidrogén, halogén, $-OCF_3$, -CN, $-CF_3$, $-NO_2$, adott esetben szubsztituált-
 $-(C_1-C_6)alkil$, $-(C_3-C_6)cikloalkil$, $-(C_2-C_6)alkenil$, $-(C_2-C_6)alkinil$, halogén- $-(C_1-C_6)alkil$,
 $-(C_0-C_6)alkilOR^7$, $-(C_0-C_6)alkilNR^8R^9$, $-(C_0-C_6)alkil-C(=NR^{10})NR^8R^9$, $-(C_0-C_6)alkilSR^6$,
 $-(C_0-C_6)alkilNR^8C(=O)R^9$, $-(C_0-C_6)alkilNR^8S(=O)_2R^9$, $-(C_0-C_6)alkilNR^{10}C(=O)NR^8R^9$,
 $-(C_0-C_6)alkilS(=O)R^8$, $-(C_0-C_6)alkilS(=O)_2R^8$, $-(C_0-C_6)alkilS(=O)_2NR^8R^9$, $-(C_0-C_6)alkil-$
 $C(=O)R^8$, $-(C_0-C_6)alkilC(=O)OR^7$, $-(C_1-C_6)alkilC(=O)NR^8R^9$, vagy $-(C_0-C_6)alkil-$
 $C(=NOR^7)R^9$, heteroaril, heteroarilalkil, arilalkil, aril, 3-6 heterocikloalkil; ahol adott
 esetben két szubsztituens a közbülső atomokkal együttesen biciklusos heterocikloalkil-,
 aril- vagy heteroarilgyűrűt alkot; ahol mindegyik gyűrű adott esetben tovább
 szubsztituált 1-5 egymástól független halogén, -CN, $-(C_1-C_6)alkil$, $-O(C_0-C_6)alkil$, $-O(C_3-$
 $C_7)cikloalkil$, $-O(aril)$, $-O(aril)-(C_0-C_3)alkil$, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)alkil$,
 $-O(C_1-C_3)alkilaril$, $-O(C_1-C_3)alkilheteroaril$, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)((C_0-$
 $C_3)alkilaril)$ vagy $-N((C_0-C_6)alkil)((C_0-C_3)alkilheteroaril)$ csoporttal;

R^7 jelentését minden esetben az alábbiak közül választjuk hidrogén, $-(C_1-C_6)$ alkil, $-(C_2-C_6)$ alkenil, $-(C_3-C_7)$ cikloalkil, 3-7 tagú heterocikloalkil, $-(C_6-C_{10})$ aril, 5 vagy 6 tagú heteroaril, $-(C_7-C_{10})$ arilalkil, $-C(=O)R^{11}$;

R^{11} jelentését az alábbiak közül választjuk: $-(C_1-C_6)$ alkil, $-(C_3-C_7)$ cikloalkil és $-(C_6-C_{10})$ aril;

R^8 , R^9 és R^{10} mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)$ alkil, $-(C_3-C_6)$ cikloalkil, $-(C_2-C_6)$ alkenil, $-(C_2-C_6)$ alkinil, $-(C_1-C_6)$ alkilhalogén, heterocikloalkil, heteroaril, heteroarilalkil, arilalkil vagy aril; amelyek bármelyike adott esetben szubsztituált 1-5 egymástól független halogén, $-CN$, $-(C_1-C_6)$ alkil, $-O(C_0-C_6)$ alkil, $-O(C_3-C_7)$ cikloalkilalkil, $-O(aril)$, $-O(aril)-(C_0-C_3)$ alkil, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)$ alkil, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)((C_3-C_7)cikloalkil)$ vagy $-N((C_0-C_6)alkil)(aril)$ szubsztituenssel;

E jelentését az alábbiakból álló csoportból választjuk: hidrogén, halogén, $-OCF_3$, $-NO_2$, $-CF_3$, $-(C_1-C_6)$ alkil, $-(C_3-C_6)$ cikloalkil, halogén- $-(C_1-C_6)$ alkil, $-(C_0-C_6)alkilOR^{12}$, $-(C_0-C_6)alkilNR^{13}R^{14}$, $-(C_0-C_6)alkilSR^{12}$, $-(C_0-C_6)alkilNR^{13}C(=O)R^{14}$, $-(C_0-C_6)alkilNR^{13}S(=O)_2R^{14}$, $-(C_0-C_6)alkilNR^{15}C(=O)NR^{13}R^{14}$, $-(C_0-C_6)alkilS(=O)R^{13}$, $-(C_0-C_6)alkilS(=O)_2R^{13}$, $-(C_0-C_6)alkilS(=O)_2NR^{13}R^{14}$, $-(C_0-C_6)alkil-C(=O)R^{13}$, $-(C_0-C_6)alkilC(=O)OR^{13}$, $-(C_0-C_6)alkilC(=O)NR^{13}R^{14}$; ahol adott esetben két szubsztituens a közbűlő atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol minden gyűrű adott esetben tovább szubsztituált 1-5 egymástól független halogén, $-CN$, adott esetben szubsztituált $-(C_1-C_6)$ alkil, $-O(C_0-C_6)$ alkil, $-O(C_3-C_7)$ cikloalkil, $-O(aril)$, $-O(aril)-(C_0-C_3)$ alkil, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)$ alkil, $-O(C_1-C_3)alkilaril$, $-O(C_1-C_3)alkilheteroaril$, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)((C_0-C_3)alkilaril)$ vagy $-N((C_0-C_6)alkil)-((C_0-C_3)alkilheteroaril)$ csoporttal;

R^{12} jelentését minden esetben az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)$ alkil, $-(C_2-C_6)$ alkenil, $-(C_3-C_7)$ cikloalkil, $-C(=O)R^{16}$;

R^{16} jelentését az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)$ alkil, $-(C_3-C_7)$ cikloalkil és $-(C_6-C_{10})$ aril;

R^{13} , R^{14} és R^{15} mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)$ alkil, $-(C_3-C_6)$ cikloalkil, $-(C_2-C_6)$ alkenil, $-(C_2-C_6)$ alkinil, $-(C_1-C_6)$ alkilhalogén, heterocikloalkil, heteroaril, heteroarilalkil, alkilheteroaril, arilalkil, alkilaril vagy aril; amelyek közül bármelyik adott esetben szubsztituált 1-5 egymástól független halogén, $-CN$, $-(C_1-C_6)$ alkil, $-O(C_0-C_6)$ alkil, $-O(C_3-C_7)$ cikloalkilalkil, $-O(aril)$, $-O(aril)-(C_0-C_3)$ alkil, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)$ alkil, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)((C_3-C_7)cikloalkil)$ vagy $-N((C_0-C_6)alkil)(aril)$ szubsztituenssel;

Amikor E jelentése $-H$, akkor azok a vegyületek, amelyeknél X_2 jelentése nem $CH=$ vagy $-CF=$, nem képezik a találmány tárgyát;

L jelentését egymástól függetlenül 1-6 T szubsztituált $-(C_1-C_5)\text{alkil}-Q_m$ - csoportból választjuk;

m értéke 0 vagy 1;

T jelentése 1-6 szubsztituens, amelyet egymástól függetlenül az alábbiakból álló csoportból választunk: hidrogén, fluor, $-OCF_3$, $-NO_2$, $-CN$, $-CF_3$, $-(C_1-C_5)\text{alkil}$, $-(C_3-C_6)\text{cikloalkil}$, $-(C_2-C_5)\text{alkenil}$, $-(C_2-C_5)\text{alkinil}$, halogén- $-(C_1-C_6)\text{alkil}$, $-(C_0-C_6)\text{alkilOR}^{17}$, $-(C_0-C_6)\text{alkilNR}^{18}\text{R}^{19}$, $-(C_0-C_6)\text{alkil}-C(=NR^{20})NR^{18}\text{R}^{19}$, $-(C_0-C_6)\text{alkilSR}^{18}$, $-(C_0-C_6)\text{alkilNR}^{18}C(=O)R^{19}$, $-(C_0-C_6)\text{alkilNR}^{18}C(=O)_2R^{19}$, $-(C_0-C_6)\text{alkilNR}^{18}S(=O)_2R^{19}$, $-(C_0-C_6)\text{alkilNR}^{20}C(=O)NR^{18}\text{R}^{19}$, $-(C_0-C_6)\text{alkilS}(=O)R^{18}$, $-(C_0-C_6)\text{alkilS}(=O)_2R^{18}$, $-(C_0-C_6)\text{alkilS}(=O)_2NR^{18}\text{R}^{19}$, $-(C_0-C_6)\text{alkilC}(=O)R^{18}$, $-(C_0-C_6)\text{alkilC}(=O)OR^{17}$, $-(C_0-C_6)\text{alkilC}(=O)NR^{18}\text{R}^{19}$, $-(C_0-C_6)\text{alkil}-C(=NR^{18})R^{19}$, vagy $-(C_0-C_6)\text{alkilC}(=NOR^{17})R^{19}$, heteroaril, heteroarilalkil, arilalkil, alkilaril, alkilheteroaril, aril; ahol adott esetben két szubsztituens a közbülső atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol mindegyik gyűrű adott esetben tovább szubsztituált 1-5 egymástól független halogén, $-CN$, $-(C_1-C_6)\text{alkil}$, $-O(C_0-C_6)\text{alkil}$, $-O(C_3-C_7)\text{cikloalkil}$, $-O(\text{aril})$, $-O(\text{aril})-(C_0-C_3)\text{alkil}$, $-O(\text{heteroaril})$, $-O(\text{heteroaril})-(C_0-C_3)\text{alkil}$, $-O(C_1-C_3)\text{alkilaril}$, $-O(C_1-C_3)\text{alkilheteroaril}$, $-N((C_0-C_6)\text{alkil})((C_0-C_3)\text{alkilaril})$ vagy $-N((C_0-C_6)\text{alkil})((C_0-C_3)\text{alkilheteroaril})$ csoporttal;

R^{17} jelentését minden esetben az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_2-C_6)\text{alkenil}$, $-(C_3-C_7)\text{cikloalkil}$, 3-7 tagú heterocikloalkil, $-(C_5-C_{10})\text{aril}$, 5 vagy 6 tagú heteroaril, $-(C_7-C_{10})\text{arilalkil}$, $-C(=O)R^{21}$;

R^{21} jelentése hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_3-C_7)\text{cikloalkil}$, vagy $-(C_5-C_{10})\text{aril}$;

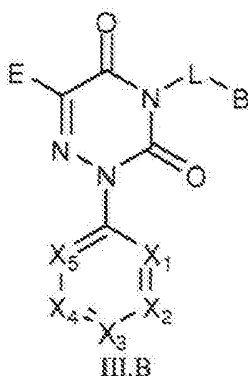
R^{18} , R^{19} és R^{20} mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_3-C_6)\text{cikloalkil}$, $-(C_2-C_6)\text{alkenil}$, $-(C_2-C_6)\text{alkinil}$, $-(C_1-C_6)\text{alkilhalogén}$, heterocikloalkil, heteroaril, heteroarilalkil, arilalkil alkilaril, alkilheteroaril, aril; amelyek közül bármelyik adott esetben szubsztituált 1-5 egymástól független halogén, $-CN$, $-(C_1-C_6)\text{alkil}$, $-O(C_0-C_6)\text{alkil}$, $-O(C_3-C_7)\text{cikloalkilalkil}$, $-O(\text{aril})$, $-O(\text{aril})-(C_0-C_3)\text{alkil}$, $-O(\text{heteroaril})$, $-O(\text{heteroaril})-(C_0-C_3)\text{alkil}$, $-N((C_0-C_6)\text{alkil})_2$, $-N((C_0-C_6)\text{alkil})((C_3-C_7)\text{cikloalkil})$ vagy $-N((C_0-C_6)\text{alkil})(\text{aril})$ szubsztituenssel;

Q jelentése $-C(=O)-$, $-C(=O)O-$, $-C(=O)NR^{22}-$, $-OC(=O)-$, $-OC(=O)NR^{22}-$, $-NR^{22}-$, $-NR^{22}C(=O)-$, $-NR^{22}C(=O)_2-$, $-NR^{22}C(=S)-$, $-S-$, $-S(=O)-$, vagy $-S(=O)_2-$;

R^{22} jelentése $-(C_1-C_6)\text{alkil}$, vagy $-(C_5-C_{10})\text{aril}$;

Bármely N N-oxidot képezhet.

3. A 2. igénypont szerinti III.B képletű vegyületek



és az ilyen vegyületek sztereoizomer alakjai, sztereoizomer alakjainak keverékei vagy gyógyszerészetileg elfogadható sói, hidrátjai vagy szolvátjai,

ahol:

B jelentése adott esetben 1-5 Y csoporttal helyettesített C_6 - C_{10} aril, adott esetben 1-7 Y csoporttal helyettesített C_5 - C_{10} cikloalkenil, adott esetben 1-8 Y csoporttal helyettesített 5-14 tagú heteroarilcsoport, ahol a heteroarilcsoport egy, kettő vagy három a N, O, S közül választott, heteroatomot tartalmaz;

Y jelentése az alábbiakból álló csoportból egymástól függetlenül választott 1-5 szubsztituens: hidrogén, halogén, $-OCF_3$, $-NO_2$, $-CN$, $-CF_3$, adott esetben szubsztituált- $-(C_1-C_6)$ alkil, $-(C_3-C_6)$ cikloalkil, $-(C_2-C_6)$ alkenil, $-(C_2-C_6)$ alkinil, halogén- $-(C_1-C_6)$ alkil, $-(C_0-C_6)$ alkil OR^1 , $-(C_0-C_6)$ alkil NR^2R^3 , $-(C_0-C_6)$ alkil- $C(=NR^4)NR^2R^3$, $-(C_0-C_6)$ alkil SR^2 , $-(C_0-C_6)$ alkil $NR^2C(=O)R^2$, $-(C_0-C_6)$ alkil $NR^2S(=O)_2R^3$, $-(C_0-C_6)$ alkil $NR^4C(=O)NR^2R^3$, $-(C_0-C_6)$ alkil $S(=O)R^2$, $-(C_0-C_6)$ alkil $S(=O)_2R^2$, $-(C_0-C_6)$ alkil $S(=O)_2NR^2R^3$, $-(C_0-C_6)$ alkil- $C(=O)R^2$, $-(C_0-C_6)$ alkil $C(=O)OR^1$, $-(C_0-C_6)$ alkil $C(=O)NR^2R^3$, $-(C_0-C_6)$ alkil $C(=NR^2)R^3$, vagy $-(C_0-C_6)$ alkil $C(=NOR^1)R^3$, heteroaril, heteroarilalkil, arilalkil, arilalkilaril, alkilheteroaril; ahol adott esetben két szubsztituens a közbülső atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol minden gyűrű adott esetben tovább szubsztituált 1-5 egymástól független halogén, $-CN$, $-(C_1-C_6)$ alkil, $-O(C_0-C_6)$ alkil, $-O(C_3-C_7)$ cikloalkil, $-O(aril)$, $-O(aril)-(C_0-C_3)$ alkil, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)$ -alkil, $-O(C_1-C_3)$ alkilaril, $-O(C_1-C_3)$ alkilheteroaril, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)-((C_0-C_3)alkilaril)$ vagy $-N((C_0-C_6)alkil)((C_0-C_3)alkilheteroaril)$ csoporttal;

R^1 jelentését minden esetben az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)$ alkil, $-(C_2-C_6)$ alkenil, $-(C_3-C_7)$ cikloalkil, 3-7 tagú heterocikloalkil, $-(C_6-C_{10})$ aril, 5 vagy 6 tagú heteroaril, $-(C_7-C_{10})$ arilalkil, $-C(=O)R^5$;

R^5 jelentését az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)$ alkil, és $-(C_6-C_{10})$ aril;

R^2 , R^3 és R^4 mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)$ alkil, $-(C_3-C_6)$ cikloalkil, $-(C_2-C_6)$ alkenil, $-(C_2-C_6)$ alkinil, $-(C_1-C_6)$ alkilhalogén, heterocikloalkil, heteroaril, heteroarilalkil, arilalkil vagy aril; amelyek közül bármelyik adott esetben szubsztituált 1-5 egymástól független halogén, $-CN$, $-(C_1-C_6)$ alkil, $-O(C_0-C_6)$ alkil, $-O(C_3-C_7)$ -cikloalkilalkil, $-O(aril)$, $-O(heteroaril)$, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)((C_3-C_7)-$

-cikloalkil) vagy $-N((C_0-C_6)alkil)(aril)$ szubsztituenssel; ahol adott esetben R^2 , R^3 , R^4 szubsztituensek a közbülső atomokkal együttesen biciklusos heterocikloalkil csoportot alkotnak;

X_1 , X_2 , X_3 , X_4 és X_5 mindegyikének jelentését egymástól függetlenül az alábbiakból álló csoportból választjuk: $-CR^6=$, $-N=$;

R^6 jelentése az alábbiakból álló csoportból egymástól függetlenül választott 1-5 szubsztituens: hidrogén, halogén, $-OCF_3$, $-CN$, $-CF_3$, $-NO_2$, adott esetben szubsztituált- $-(C_1-C_6)alkil$, $-(C_3-C_6)cikloalkil$, $-(C_2-C_6)alkenil$, $-(C_2-C_6)alkinil$, halogén- $-(C_1-C_6)alkil$, $-(C_0-C_6)alkilOR^7$, $-(C_0-C_6)alkilNR^8R^9$, $-(C_0-C_6)alkilSR^8$, $-(C_0-C_6)alkilNR^8C(=O)R^9$, $-(C_0-C_6)alkilNR^8S(=O)_2R^9$, $-(C_0-C_6)alkilNR^{10}C(=O)NR^8R^9$, $-(C_0-C_6)alkilS(=O)R^8$, $-(C_0-C_6)alkilS(=O)_2R^8$, $-(C_0-C_6)alkilS(=O)_2NR^8R^9$, $-(C_0-C_6)alkil-C(=O)R^8$, $-(C_0-C_6)alkilC(=O)OR^7$, $-(C_1-C_6)alkilC(=O)NR^8R^9$, heteroaril, heteroarilalkil, arilalkil, aril, 3-6 heterocikloalkil; ahol adott esetben két szubsztituens a közbülső atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol mindegyik gyűrű adott esetben tovább szubsztituált 1-5 egymástól független halogén, $-CN$, $-(C_1-C_6)alkil$, $-O(C_0-C_6)alkil$, $-O(C_3-C_7)cikloalkil$, $-O(aril)$, $-O(aril)-(C_0-C_3)alkil$, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)alkil$, $-O(C_1-C_3)alkilaril$, $-O(C_1-C_3)alkilheteroaril$, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)((C_0-C_3)alkilaril)$ vagy $-N((C_0-C_6)alkil)((C_0-C_3)alkilheteroaril)$ csoporttal;

R^7 jelentését minden esetben az alábbiak közül választjuk hidrogén, $-(C_1-C_6)alkil$, $-(C_2-C_6)alkenil$, $-(C_3-C_7)cikloalkil$, 3-7 tagú heterocikloalkil, $-(C_6-C_{10})aril$, 5 vagy 6 tagú heteroaril, $-(C_7-C_{10})arilalkil$, $-C(=O)R^{11}$;

R^{11} jelentését az alábbiak közül választjuk: $-(C_1-C_6)alkil$, $-(C_3-C_7)cikloalkil$ és $-(C_6-C_{10})aril$;

R^8 , R^9 és R^{10} mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)alkil$, $-(C_3-C_6)cikloalkil$, $-(C_2-C_6)alkenil$, $-(C_2-C_6)alkinil$, $-(C_1-C_6)alkilhalogén$, heterocikloalkil, heteroaril, heteroarilalkil, arilalkil vagy aril; amelyek bármelyike adott esetben szubsztituált 1-5 egymástól független halogén, $-CN$, $-(C_1-C_6)alkil$, $-O(C_0-C_6)alkil$, $-O(C_3-C_7)cikloalkilalkil$, $-O(aril)$, $-O(aril)-(C_0-C_3)alkil$, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)alkil$, $-N((C_0-C_6)alkil)_2$, $-N((C_0-C_6)alkil)((C_3-C_7)cikloalkil)$ vagy $-N((C_0-C_6)alkil)(aril)$ szubsztituenssel;

E jelentését az alábbiakból álló csoportból választjuk: hidrogén, halogén, $-OCF_3$, $-CF_3$, $-(C_1-C_6)alkil$, $-(C_3-C_6)cikloalkil$, $-(C_0-C_6)alkilOR^{12}$, $-(C_0-C_6)alkilNR^{13}R^{14}$, $-(C_0-C_6)alkilS(=O)R^{13}$, $-(C_0-C_6)alkilS(=O)_2R^{13}$, $-(C_0-C_6)alkil-C(=O)R^{13}$, $-(C_0-C_6)alkilC(=O)OR^{13}$, $-(C_0-C_6)alkilC(=O)NR^{13}R^{14}$; ahol adott esetben két szubsztituens a közbülső atomokkal együttesen biciklusos heterocikloalkil-, aril- vagy heteroarilgyűrűt alkot; ahol minden gyűrű adott esetben tovább szubsztituált 1-5 egymástól független halogén, $-CN$, adott esetben szubsztituált $-(C_1-C_6)alkil$, $-O(C_0-C_6)alkil$, $-O(C_3-C_7)cikloalkil$, $-O(aril)$, $-O(aril)-(C_0-C_3)alkil$, $-O(heteroaril)$, $-O(heteroaril)-(C_0-C_3)alkil$, $-O(C_1-C_3)alkilaril$, $-O(C_1-C_3)alkil-$

-heteroaril, $-N((C_0-C_6)\text{alkil})_2$, $-N((C_0-C_6)\text{alkil})((C_0-C_3)\text{alkilaril})$ vagy $-N((C_0-C_6)\text{alkil})((C_0-C_3)\text{alkilheteroaril})$ csoporttal;

R^{12} jelentését minden esetben az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_2-C_6)\text{alkenil}$, $-(C_3-C_7)\text{cikloalkil}$, $-C(=O)R^{15}$;

R^{15} jelentését az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_3-C_7)\text{cikloalkil}$ és $-(C_6-C_{10})\text{aril}$;

R^{13} és R^{14} mindegyikének jelentése egymástól függetlenül hidrogén, $-(C_1-C_6)\text{alkil}$, $-(C_3-C_6)\text{cikloalkil}$, $-(C_2-C_6)\text{alkenil}$, $-(C_2-C_6)\text{alkinil}$, $-(C_1-C_6)\text{alkilhalogén}$, heterocikloalkil, heteroaril, heteroarilalkil, alkilheteroaril, arilalkil, alkilaril vagy aril; amelyek közül bármelyik adott esetben szubsztituált 1-5 egymástól független halogén, $-\text{CN}$, $-(C_1-C_6)\text{alkil}$, $-\text{O}(C_0-C_6)\text{alkil}$, $-\text{O}(C_3-C_7)\text{cikloalkilalkil}$, $-\text{O}(\text{aril})$, $-\text{O}(\text{aril})-(C_0-C_3)\text{alkil}$, $-\text{O}(\text{heteroaril})$, $-\text{O}(\text{heteroaril})-(C_0-C_3)\text{alkil}$, $-N((C_0-C_6)\text{alkil})_2$, $-N((C_0-C_6)\text{alkil})((C_3-C_7)\text{cikloalkil})$ vagy $-N((C_0-C_6)\text{alkil})(\text{aril})$ szubsztituenssel;

Amikor E jelentése $-\text{H}$, akkor azok a vegyületek, amelyeknél X_3 jelentése nem CH= vagy $-\text{CF=}$, nem képezik a találmány tárgyát;

L jelentését egymástól függetlenül 1-6 T szubsztituált $-((C_1-C_5)\text{alkil}-Q_m)-$ csoportból választjuk;

m értéke 0 vagy 1;

T jelentése 1-6 szubsztituens, amelyet egymástól függetlenül az alábbiakból álló csoportból választunk: hidrogén, $-(C_1-C_6)\text{alkil}$,

Q jelentése $-C(=O)-$, $-C(=O)O-$, $-C(=O)NR^{22}-$, $-\text{OC}(=O)-$, $-\text{OC}(=O)NR^{22}-$, $-\text{NR}^{22}-$, $-\text{NR}^{22}C(=O)-$, $-\text{NR}^{22}C(=O)_2-$, $-\text{NR}^{22}C(=S)-$, $-\text{S}-$, $-\text{S}(=O)-$, vagy $-\text{S}(=O)_2-$;

R^{22} jelentése $-(C_1-C_6)\text{alkil}$, vagy $-(C_6-C_{10})\text{aril}$;

Bármely N N-oxidot képezhet.

4. Az 1-3. igénypontok bármelyike szerinti vegyület, amely optikai izomer alakban létezik, ahol a vegyület vagy racém keverék, vagy egy egyedi optikai izomer.

5. Az 1-4. igénypontok bármelyike szerinti vegyület, amely vegyületet az alábbiak közül választjuk:

$N\text{-}\{3\text{-}\{4\text{-}\{4\text{-klórbenzil}\}\text{-}3,5\text{-dioxo-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{acetamid}$, $N\text{-}\{3\text{-}\{4\text{-}\{4\text{-klórbenzil}\}\text{-}3,5\text{-dioxo-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{metánszulfonamid}$, $4\text{-}\{4\text{-klórbenzil}\}\text{-}2\text{-}\{2\text{-metoxifenil}\}\text{-}1,2,4\text{-triazin-}3,5(2H,4H)\text{-dion}$, $N\text{-}\{3\text{-}\{4\text{-}\{4\text{-nitrobenzil}\}\text{-}3,5\text{-dioxo-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{acetamid}$, $N\text{-}\{3\text{-}\{4\text{-}\{4\text{-cianobenzil}\}\text{-}3,5\text{-dioxo-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{acetamid}$, $N\text{-}\{3\text{-}\{4\text{-}\{4\text{-naftil-}2\text{-ilmetil}\}\text{-}3,5\text{-dioxo-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{acetamid}$, $N\text{-}\{3\text{-}\{4\text{-}\{benzo[c][1,2,5]oxadiazol-5\text{-ilmetil}\}\text{-}3,5\text{-dioxo-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{acetamid}$, $N\text{-}\{3\text{-}\{4\text{-}\{4\text{-fluorbenzil}\}\text{-}3,5\text{-dioxo-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{acetamid}$, $N\text{-}\{3\text{-}\{4\text{-benzil-}3,5\text{-dioxo-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{acetamid}$, $N\text{-}\{3\text{-}\{3,5\text{-dioxo-}4\text{-}\{4\text{-}\{trifluormetil}\}\text{benzil}\}\text{-}4,5\text{-dihidro-}1,2,4\text{-triazin-}2(3H)\text{-il}\}\text{fenil}\}\text{acetamid}$, metil-4-((2-(3-acetamidofenil)-3,5-

-dioxo-2,3-dihidro-1,2,4-triazin-4(5H)-il)metil)benzoát, N-(3-(4-(naftil-1-ilmetil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-metoxibenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-brómbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3-klórbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-metilbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-izopropilbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3,4-dimetilbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-klórpíridin-3-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klór-2-fluorbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-benzil-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klór-3-fluorbenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-fluorpiridin-3-il)metil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-izopropilpiridin-3-il)metil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-fluorbenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(4-fluorbenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-metoxibenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-izopropilbenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-(4-klórbenzil)-6-metil-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(6-metil-4-(naftil-2-ilmetil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-ciklopentilpiridin-3-il)metil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3,4-difluorbenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3,4-dimetoxibenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(2,4-difluorbenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-izopropoxibenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metil-4-(naftil-1-ilmetil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metil-4-(4-metilbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(6-metil-4-(4-metilbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(3-metoxibenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 3-(4-(4-fluorbenzil)-6-metil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-N-metilbenzamid, N-(3-(4-(4-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-(4-nitrobenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-(4-metilbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-3,5-dioxo-4-(4-(trifluor-metil)benzil)-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klór-3-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-(4-klór-3-

-fluorbenzil)-6-metoxi-2-(2-metoxifenil)-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)metánszulfonamid, 4-(4-klórbenzil)-6-metoxi-2-(2-metoxifenil)-1,2,4-triazin-3,5(2H,4H)-dion, 1-(3-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-3-metilkarbamid, N-(3-(4-(6-klórpíridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(6-metoxi-4-(4-metilbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)pyridin-3-il)acetamid, N-(3-(6-metoxi-4-(6-metilpiridin-3-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klór-3-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)piridin-3-il)acetamid, 4-(4-klórbenzil)-2-(4-fluor-2-metoxifenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(2-(metoximetil)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(3-(metoximetil)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(3-(2-metoxietil)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(2-(trifluor-metoxi)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-2-(2,3-dimetoxifenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(2-(morfolino-metil)-fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(2-(2-metoxietil)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(3-(2-metil-1,3-dioxolan-2-il)metil)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(3-(2-oxopropil)-fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-fluorbenzil)-2-(2-(dimetilamino)metil)fenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-fluorbenzil)-6-metoxi-2-(2-(pirrolidin-1-ilmetil)-fenil)-1,2,4-triazin-3,5(2H,4H)-dion, N-(2-fluor-5-(6-metoxi-4-(4-metoxibenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-(4-fluorbenzil)-2-(4-fluorfenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-fluorbenzil)-6-metoxi-2-(2-(piperidin-1-ilmetil)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(2-((2-metoxietil)(metil)amino)metil)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, N-(2-fluor-5-(4-(3-fluor-4-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 2-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)benzonitril, 4-(4-klórbenzil)-2-(4-fluor-2-(hidroximetil)fenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, 4-(2,4-diklórbenzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-benzil-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-izopropilbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(bifenil-3-ilmetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-3,5-dioxo-4-(4-(trifluor-metoxi)benzil)-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-(dimetil-amino)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(bifenil-4-ilmetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-benzil-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-(naftil-2-ilmetil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid,

N-(3-(4-((6-izopropilpiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-(4-metoxibenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-3,5-dioxo-4-(quinolin-6-ilmetil)-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-izopropoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3,4-dimetoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-ciklopentilpiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-(4-klórbenzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-((6-fluorpiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-klór-5-fluorpiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-(dimetilamino)piridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((2,3-dihidrobenezofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3,4-difluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(2,4-difluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-((6-metoxipiridin-3-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-3,5-dioxo-4-(piridin-3-ilmetil)-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-(1H-pirazol-1-il)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(2,6-difluor-4-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-3,5-dioxo-4-(kinolin-3-ilmetil)-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3-fluor-4-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3,5-difluor-4-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(2,3-difluor-4-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-((1-metil-1H-indol-5-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(2-fluor-4-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-(3,4-difluorbenzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-(2-klór-4-fluorbenzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klór-2-fluorbenzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(6-metoxi-4-(3-metoxibenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-((2,3-dihidrobenzo[b][1,4]dioxin-2-il)metil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 6-metoxi-2-fenil-4-((2-feniltiazol-4-il)metil)-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-klór-2-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 6-metoxi-2-fenil-4-(3-(trifluormetoxi)benzil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(2,5-difluorbenzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klór-3-(trifluormetoxi)benzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(2,4-difluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)a-

cetamid, N-(3-(4-((6-fluor-2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-(4-fluor-3-(trifluormetil)benzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-benzoilbenzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(4-((6-metoxi-3,5-dioxo-2-fenil-2,3-dihidro-1,2,4-triazin-4(5H)-il)metil)tiazol-2-il)acetamid, N-(3-(4-(furo[2,3-b]piridin-5-ilmetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(benzo[d]oxazol-6-ilmetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-fluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 3-((6-metoxi-3,5-dioxo-2-fenil-2,3-dihidro-1,2,4-triazin-4(5H)-il)metil)benzonitril, 4-((6-metoxi-3,5-dioxo-2-fenil-2,3-dihidro-1,2,4-triazin-4(5H)-il)metil)benzonitril, N-(3-(4-((7-fluor-2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-(4-(hidroximetil)benzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-((6-klór-4H-benzo[d][1,3]dioxin-8-il)metil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 6-metoxi-2-fenil-4-((5-(trifluormetil)furan-2-il)metil)-1,2,4-triazin-3,5(2H,4H)-dion, 6-metoxi-4-((1-metil-1H-imidazol-2-il)metil)-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(2,4-diklórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-(dietilamino)piridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-((5-klórtien-2-il)metil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-((5-klórbenzo[b]tien-3-il)metil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-(2,4-difluorbenzil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-((2,3-dihidrofuro[2,3-b]piridin-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((4-fluor-2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-fluor-3-(trifluormetil)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((2-etilbenzo[d]oxazol-6-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-cianobenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-((2-metilbenzo[d]oxazol-6-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((6-(dimetilamino)-5-fluorpiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-(dimetilamino)-3-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-((1-metilindolin-5-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(2-fluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((4-fluor-2,3-dihidrobenzofuran-7-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((2,2-difluorbenzo[d][1,3]dioxol-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((5-klórpíridin-2-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-((5-izopropilpiridin-2-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(3,5-difluor-4-

-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(4-fluor-3-(trifluormetoxi)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(3-fluor-4-(trifluormetoxi)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-N-metilacetamid, N-(3-(6-metoxi-4-(4-nitrobenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-N-metilacetamid, N-(5-(4-((2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(3-(6-metoxi-4-(4-metoxibenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-N-metilacetamid, N-(3-(4-((2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-N-metilacetamid, N-(3-(4-(4-fluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-N-metilacetamid, N-(3-(4-(2,4-difluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-N-metilacetamid, 6-metoxi-4-fenetil-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-2-metoxiacetamid, N-(3-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-3-metoxipropanamid, 4-(4-klórbenzil)-2-(2-hidroxifenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(3-(tiazol-2-ilamino)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(3-(2-oxopirrolidin-1-il)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-klórbenzil)-6-metoxi-2-(3-(5-metil-1,2,4-oxadiazol-3-il)fenil)-1,2,4-triazin-3,5(2H,4H)-dion, 6-metoxi-4-(4-metoxifenetil)-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-fluorfenetil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-(3,4-dimetoxifenetil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 6-metoxi-4-(3-metoxifenetil)-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 6-metoxi-4-(4-metoxifenetil)-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, 4-(4-fluorfenetil)-6-metoxi-2-fenil-1,2,4-triazin-3,5(2H,4H)-dion, N-(3-(6-metoxi-4-(3-metoxifenetil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-benzil-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(4-fluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(2,4-difluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(4,5-difluor-2-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(6-metoxi-3,5-dioxo-4-fenetil-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-((6-klór-5-fluorpiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(6-metoxi-4-(3-metoxibenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(2,4-difluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-4-fluorfenil)acetamid, N-(2-fluor-5-(6-metoxi-4-((1-metilindolin-5-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-((6-fluor-2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-(di-

metilamino)-3-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(3,4-difluor-2-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, 6-metoxi-2-fenil-4-(3-fenilpropil)-1,2,4-triazin-3,5(2H,4H)-dion, 2-(2-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-cianobenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(6-metoxi-4-(2-metoxibenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-ciano-3-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(6-cianopiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(6-izopropilpiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(6-metoxi-4-(6-metoxipiridin-3-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(6-metoxi-3,5-dioxo-4-(kinolin-6-ilmetil)-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(benzofuran-5-ilmetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(4-fluorfenetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(3-klór-4-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(3-fluor-4-(pirrolidin-1-il)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(3-fluor-4-morfolinobenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(2,6-difluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(6-metoxi-3,5-dioxo-4-(4-(trifluormetil)benzil)-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(6-metoxi-4-(3-(metoximetil)benzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(furo[2,3-b]piridin-5-ilmetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(2-fluor-5-(metoximetil)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(4-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(6-metoxi-4-(4-nitrobenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-(1H-pirazol-1-il)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(3-fluor-5-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(3-(dimetilamino)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(3-(6-metoxi-4-(1-metil-1H-indazol-6-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-(4-metil-3,4-dihidro-2H-benzo[b][1,4]oxazin-7-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(2,3-dihidrobenczo[b][1,4]dioxin-6-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(3-(6-metoxi-4-(1-metil-1H-indazol-5-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-4-(2-metil-2H-indazol-5-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-

-il)fenil)acetamid, N-(5-(4-((2,3-dihidrobenzofuran-6-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-((2,3-dihidrobenzofuran-4-il)-metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(3-(6-metoxi-4-((5-metoxipiridin-3-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-ciano-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-metoxi-3,5-dioxo-4-fenetil-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(4-fluor-2-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-(1H-pirazol-1-il)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(3,4-difluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(6-metoxi-4-((1-metil-1H-indazol-6-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(3,4-diklórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(3-(dimetilamino)-4-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-klor-5-(4-(4-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(2,2-dimetil-2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, 4-(4-klórbenzil)-2-(4-fluorfenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, N-(5-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(3-(4-(benzo[d]izoxazol-6-ilmetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klor-3-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-klor-5-(4-(4-klor-3-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-klor-5-(4-(4-fluor-2-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(2,4-difluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(4-klor-2-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(4-klor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(2,4-difluorfenetil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(3-(4-((5-fluor-6-metoxipiridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-klor-5-(4-(4-klor-2-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-klor-5-(4-(4-klor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(3,4-difluor-5-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-klor-5-(4-(4-klor-2-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-klor-5-(4-(2,4-difluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-klor-5-(4-(4-klórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klor-2-(dimetilamino)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)a-

cetamid, N-(2-klór-5-(4-(4-klór-2-(dimetilamino)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-klór-5-(4-(3-(dimetilamino)-4-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-benzil-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-klór)fenil)acetamid, N-(5-(4-(2-(dimetilamino)-4-fluorbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-klór-5-(4-(4-fluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(3-fluor-4-(trifluormetil)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klór-3-(dimetilamino)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-klór-5-(4-(4-klór-3-(dimetilamino)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(3,4-diklórbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-klór-5-(4-(6-(dimetilamino)-píridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klór-3-(trifluormetoxi)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-klór-5-(6-metoxi-4-(3-metoxibenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-klór-5-(4-(2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-klór-5-(4-(3,4-difluor-2-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 4-(4-klórbenzil)-2-(4-fluorfenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, N-(2-klór-5-(4-(4-klór-3-(trifluormetil)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klór-3-izopropoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-klór-5-(6-metoxi-4-(1-metil-1H-indazol-6-il)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klór-3-izopropilbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(3-fluor-4-morfolinobenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(4-klór-3-hidroxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(3,3-dimetil-2,3-dihidrobenzofuran-5-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(3,3-dimetil-2,3-dihidrobenzofuran-6-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(3-fluor-4-izopropilbenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klór-3-(dimetilamino)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-4-fluorfenil)acetamid, N-(5-(4-(4-ciano-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, 4-(3,4-diklórbenzil)-2-(4-fluorfenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, N-(5-(4-(4-klór-2-fluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, 4-(4-klór-2-fluor-3-metoxibenzil)-2-(4-fluorfenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, N-(5-(4-(4-bróm-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)aceta-

mid, N-(2-fluor-5-(4-(4-izopropil-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 2-(4-fluorfenil)-4-(4-izopropil-3-metoxibenzil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, N-(5-(4-(4-klór-3-(pirrolidin-1-il)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(4-fluor-3-(trifluormetil)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-(2,4-difluor-3-izopropoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-((5,6-diklórpíridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(4-ciano-2-fluor-3-metoxibenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, 4-((5,6-diklórpíridin-3-il)metil)-2-(4-fluorfenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, N-(5-(4-(4-klór-3-morfolinobenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(5-(4-(3-(dimetilamino)-4-(trifluormetil)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, 4-(4-klór-3-metoxibenzil)-2-(4-fluorfenil)-6-metoxi-1,2,4-triazin-3,5(2H,4H)-dion, N-(2-fluor-5-(4-(3-fluor-4-(trifluormetil)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)-N-metilacetamid, N-(5-(4-(4-klór-3-(metoximetil)benzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(2-fluor-5-(4-(4-fluor-3-morfolinobenzil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(5-(4-((6-klór-5-izopropilpíridin-3-il)metil)-6-metoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)-2-fluorfenil)acetamid, N-(3-(4-(4-klórbenzil)-6-izopropil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-etil-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-butil-4-(4-klórbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, 2-(6-bróm-4-(4-fluorbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)benzonitril, N-(3-(4-(4-klórbenzil)-6-(2-(dimetilamino)ethoxi)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-(2-metoxietoxi)-4-(4-nitrobenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-(2-metoxietoxi)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-(metilamino)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-(metilamino)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-etoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-bróm-4-(4-metilbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-izopropoxi-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-bróm-4-(4-klórbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-(dimetilamino)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-(hidroximetil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-((dimetilamino)metil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(6-((1H-pirazol-1-il)metil)-4-(4-klórbenzil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-(metoximetil)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, metil-2-(3-acetamido-

fenil)-4-(4-klórbenzil)-3,5-dioxo-2,3,4,5-tetrahidro-1,2,4-triazin-6-karboxilát, izopropil-2-(3-acetamidofenil)-4-(4-klórbenzil)-3,5-dioxo-2,3,4,5-tetrahidro-1,2,4-triazin-6-karboxilát, N-(3-(4-(4-klórbenzil)-6-(3-metil-1,2,4-oxadiazol-5-il)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-(5-metil-1,2,4-oxadiazol-3-il)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(3-(4-(4-klórbenzil)-6-(1,2,4-oxadiazol-5-il)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid, N-(2-fluor-5-(4-(4-fluorbenzil)-6-(2-metoxietoxi)-3,5-dioxo-4,5-dihidro-1,2,4-triazin-2(3H)-il)fenil)acetamid.

6. Az 1-5. igénypontok szerinti vegyületek terápiásan hatékony mennyiségét és gyógyszerészetileg elfogadható vivőanyagot és/vagy hordozót tartalmazó gyógyszerkészítmény.

7. Az 1-6. igénypontok szerinti vegyületek/készítmények egy emlősben, ideértve az embert, egy állapot kezelésében vagy megelőzésében történő alkalmazásra, ahol a kezelést vagy megelőzést befolyásolja vagy megkönnyíti egy GABA_B pozitív allosztérikus modulátor (fokozó).

8. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket a szorongásos betegségek csoportjából választunk: agorafóbia, generalizált szorongás szindróma (GAD), rögeszmés-kényszeres zavar (OCD), pánik betegség, poszt-traumatikus stressz szindróma (PTSD), szociális szorongásos zavar, egyéb fóbiák, anyagok fogyasztása okozta szorongásos zavar.

9. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket a gyermekkori betegségek csoportjából választunk (figyelemhiányos/hiperaktivitási zavar).

10. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket az étkezési betegségek csoportjából választunk (anorexia nervosa, bulimia nervosa).

11. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket a kedélybetegségek csoportjából választunk: bipoláris zavarok (I & II), ciklotimiás rendellenesség, depresszió, levertség zavar, major depressziós rendellenesség, anyagok fogyasztása okozta kedélybetegség.

12. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket a pszichotikus rendellenességek csoportjából választunk: skizofrénia, deluzív zavar, skizoaffektív zavar, skizofreniform zavar, anyagok fogyasztása okozta pszichotikus rendellenességek.

13. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket

a kognitív rendellenességek csoportjából választunk: delírium, anyagok fogyasztása okozta folytatódó delírium, demencia, HIV- okozta demencia, Huntington-kór okozta demencia, Parkinson-kór okozta demencia, Alzheimer-kór okozta demencia, anyagok fogyasztása okozta demencia, enyhe mentális hanyatlás.

14. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket a személyiség zavarok csoportjából választunk: rögeszmés-kényszeres személyiség zavar, skizoid, skizotíp személyiség zavar.

15. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket anyagok fogyasztása okozta rendellenességek csoportjából választunk: alkohol használat, alkohol függőség, alkohol megvonás, alkohol megvonási delírium, alkohol okozta pszichotikus rendellenesség, amfetamin függőség, amfetamin megvonás, kokain függőség, kokain megvonás, nikotin függőség, nikotin megvonás, opioid függőség, opioid megvonás.

16. Az 1-6. igénypontok szerinti vegyületek/készítmények a központi idegrendszeri gyulladásos rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket a szklerózis multiplex formák közül választunk: jóindulatú szklerózis multiplex, amiotrófiás laterális szklerózis, relapszáló-remittáló szklerózis multiplex, másodlagos progresszív szklerózis multiplex, elsődleges progresszív szklerózis multiplex, progresszív-relapszáló szklerózis multiplex.

17. Az 1-6. igénypontok szerinti vegyületek/készítmények a gyulladásos vagy neuropatikus fájdalom, operáció utáni fájdalom kezelésében vagy megelőzésében történő alkalmazásra.

18. Az 1-6. igénypontok szerinti vegyületek/készítmények epilepszia és remegés kezelésében vagy megelőzésében történő alkalmazásra.

19. Az 1-6. igénypontok szerinti vegyületek/készítmények készítményes vizelet inkontinencia kezelésében vagy megelőzésében történő alkalmazásra

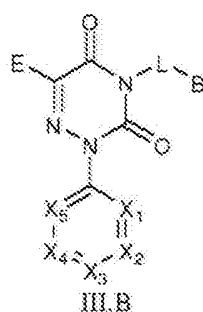
20. Az 1-6. igénypontok szerinti vegyületek/készítmények a gyomor-bél rendszeri rendellenességek kezelésében vagy megelőzésében történő alkalmazásra, amelyeket az alábbi csoportból választunk: gyomor-nyelőcső reflux betegség, nyelőcső alsó záróizmainak időszakos relaxációs zavara, emésztési zavar, funkcionális gyomor-bél rendszeri rendellenességek és gyulladásos bélbetegség.

21. Az 1-6. igénypontok szerinti vegyületek/készítmények görcs és váz-izom merevség, gerincvelő sérülés, agyi bénulás kezelésében vagy megelőzésében történő alkalmazásra.

22. Az 1-5. igénypontok szerinti vegyületek alkalmazása GABA_B receptorok leképezésére szolgáló nyomjelző készítésére.

23. Az 1-6. igénypontok szerinti vegyületek/készítmények GABA_B receptor agonistával kombinációban egy 7-21. igénypontok bármelyikében meghatározott állapot kezelésére vagy megelőzésére történő alkalmazásra egy emlősben, beleértve az embert, ahol a kezelést vagy megelőzést befolyásolja vagy megkönnyíti egy GABA_B pozitív allosztérikus modulátor neuromodulátor hatása.

24. Eljárás a 3. igénypont szerinti III.B képletű vegyületek



és az ilyen vegyületek sztereoizomer alakjai, sztereoizomer alakjainak keverékei vagy gyógyszerészetileg elfogadható sói, hidrátjai vagy szolvátjai előállítására, ahol:

B jelentése adott esetben 1-5 Y csoporttal helyettesített C₆-C₁₀ aril, adott esetben 1-8 Y csoporttal helyettesített 5-14 tagú heteroarilcsoport, ahol a heteroarilcsoport egy, kettő vagy három a N, O, S közül választott, heteroatomot tartalmaz;

Y jelentése az alábbiakból álló csoportból egymástól függetlenül választott 1-5 szubsztituens: hidrogén, halogén, -OCF₃, -NO₂, -CN, -CF₃, adott esetben szubsztituált-(C₁-C₆)alkil, -(C₃-C₆)cikloalkil, halogén-(C₁-C₆)alkil, -(C₀-C₆)alkilOR¹, -(C₀-C₆)alkilNR²R³, -(C₀-C₆)alkilSR²;

R¹ jelentését minden esetben az alábbiak közül választjuk: hidrogén, -(C₁-C₆)alkil;

R² és R³ mindegyikének jelentése egymástól függetlenül -(C₁-C₆)alkil, -(C₃-C₆)cikloalkil;

X₁, X₂, X₃, X₄ és X₅ mindegyikének jelentését egymástól függetlenül az alábbiakból álló csoportból választjuk: -CR⁶=, -N=;

R⁶ jelentése az alábbiakból álló csoportból egymástól függetlenül választott 1-5 szubsztituens: hidrogén, halogén, -OCF₃, -CN, -CF₃, -NO₂, adott esetben szubsztituált-(C₁-C₆)alkil, -(C₃-C₆)cikloalkil, -(C₀-C₆)alkilOR⁷, -(C₀-C₆)alkilNR⁸R⁹, -(C₀-C₆)alkilSR⁸, -(C₀-C₆)alkilNR⁸C(=O)R⁹, -(C₀-C₆)alkilNR⁸C(=O)₂R⁹, -(C₀-C₆)alkilNR⁸S(=O)₂R⁹, -(C₀-C₆)alkilNR¹⁰C(=O)NR⁹R⁹, -(C₀-C₆)alkilS(=O)₂R⁸, -(C₀-C₆)alkilS(=O)₂NR⁸R⁹, -(C₀-C₆)alkilC(=O)R⁸, -(C₀-C₆)alkilC(=O)OR⁷, -(C₁-C₆)alkilC(=O)NR⁸R⁹;

R⁷, R⁸ és R⁹ jelentését minden esetben az alábbiak közül választjuk hidrogén, -(C₁-C₆)alkil, -(C₃-C₇)cikloalkil;

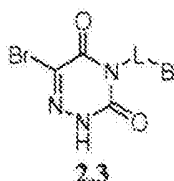
E jelentését az alábbiakból álló csoportból választjuk: $-(C_1-C_6)$ alkil, $-(C_3-C_6)$ cikloalkil, $-(C_6-C_6)$ alkilOR¹², $-(C_6-C_6)$ alkilNR¹³R¹⁴;

R¹², R¹³ és R¹⁴ mindegyikének jelentését egymástól függetlenül az alábbiak közül választjuk: hidrogén, $-(C_1-C_6)$ alkil, $-(C_2-C_6)$ alkenil, $-(C_3-C_7)$ cikloalkil;

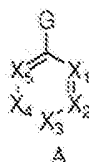
L jelentését egymástól függetlenül 1-6 T szubsztituált $-(C_1-C_5)$ alkil-csoportból választjuk;

T jelentése 1-6 szubsztituens, amelyet egymástól függetlenül az alábbiakból álló csoportból választunk: hidrogén, $-(C_1-C_6)$ alkil;

amely magában foglalja a 2.3 képletű vegyület reagáltatását



az A képletű aril vagy heteroaril boronsavval vagy észterrel réz katalizátor rendszer és egy bázis jelenlétében



ahol:

G jelentése boronsav, boronát (pl.: dimetil-boronát, pinakol-boronát és hasonló) és a keletkező reakció termék brómatomját ezután nukleofil helyettesítési reakciónak vetjük alá különböző csoportokkal a III.B képletű vegyület szolgáltatása érdekében.

A meghatalmazott:

Dr. Gal Melinda
 igazgató-helyettes
 Szent Szabolcs Utcai Iroda
 1052 Budapest, Andrássy út 113.
 Telefon: 461-1000 Fax: 461-1099
 Email: gal@szek.hu