



(51) International Patent Classification:

C07D 401/04 (2006.01) A01N 43/26 (2006.01)

C07D 403/04 (2006.01) A61P 33/00 (2006.01)

(21) International Application Number:

PCT/EP2020/059924

(22) International Filing Date:

07 April 2020 (07.04.2020)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

19168715.1 11 April 2019 (11.04.2019) EP

19196236.4 09 September 2019 (09.09.2019) EP

20152751.2 20 January 2020 (20.01.2020) EP

(71) Applicant: SYNGENTA CROP PROTECTION AG

[CH/CH]; Rosentalstrasse 67, 4058 Basel (CH).

(72) Inventors: SCHÄTZER, Jürgen, Harry; Syngenta Crop

Protection AG, Schaffhauserstrasse, 4332 Stein (CH).

EDMUNDS, Andrew; Syngenta Crop Protection AG,

Schaffhauserstrasse, 4332 Stein (CH). GAGNEPAIN,

Julien, Daniel, Henri; Syngenta Crop Protection AG,

Schaffhauserstrasse, 4332 Stein (CH). HALL, Roger,

Graham; Syngenta Crop Protection AG, Schaffhauser-

strasse, 4332 Stein (CH). JEANGUENAT, André; Syn-

genta Crop Protection AG, Schaffhauserstrasse, 4332 Stein

(CH). KOLLETH KRIEGER, Amandine; Syngenta Crop

Protection AG, Schaffhauserstrasse, 4332 Stein (CH). LE

CHAPELAIN, Camille; Syngenta Crop Protection AG,

Schaffhauserstrasse, 4332 Stein (CH). PALWE, Shrikant;

Syngenta Biosciences Pvt. Ltd., Santa Monica Plant, Cor-

lim Ilhas, Goa 403 110 (IN). PHADTE, Mangala; Syn-

genta Biosciences Private Limited, Santa Monica Works,

Corlim Ilhas, Goa 403 110 (IN). PITTERNA, Thomas;

Syngenta Crop Protection AG, Schaffhauserstrasse, 4332

Stein (CH). RENDLER, Sebastian; Syngenta Crop Pro-

tection AG, Schaffhauserstrasse, 4332 Stein (CH). SCAR-

BOROUGH, Christopher, Charles; Syngenta Crop Pro-

tection AG, Schaffhauserstrasse, 4332 Stein (CH).

(74) Agent: SYNGENTA IP; Rosentalstrasse 67, 4058 Basel (CH).

(81) Designated States (unless otherwise indicated, for every

kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP, KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, WS, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every

kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Declarations under Rule 4.17:

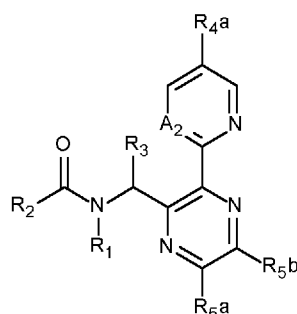
- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))
- of inventorship (Rule 4.17(iv))

Published:

- with international search report (Art. 21(3))

(54) Title: PESTICIDALLY ACTIVE DIAZINE-AMIDE COMPOUNDS

(57) Abstract: Compounds of formula (I) wherein the substituents are as defined in claim 1, and the agrochemically acceptable salts, stereoisomers, enantiomers, tautomers and N-oxides of those compounds, can be used as insecticides.



(I)

-1-

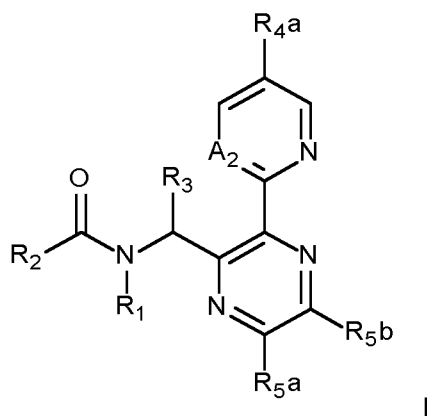
PESTICIDALLY ACTIVE DIAZINE-AMIDE COMPOUNDS

The present invention relates to pesticidally active, in particular insecticidally active diazine-amide compounds, to processes for their preparation, to compositions comprising those compounds, and to their use for controlling animal pests, including arthropods and in particular insects or representatives of the order *Acarina*.

WO2017192385 describes certain heteroaryl-1,2,4-triazole and heteroaryl-tetrazole compounds for use for controlling ectoparasites in animals (such as a mammal and a non-mammal animal).

There have now been found novel pesticidally active-diazine amide compounds.

The present invention accordingly relates, in a first aspect, to a compound of the formula I



wherein

R₁ is H, C₁-C₆alkyl, C₁-C₆cyanoalkyl, aminocarbonylC₁-C₆alkyl, hydroxycarbonylC₁-C₆alkyl, C₁-C₆nitroalkyl, trimethylsilaneC₁-C₆alkyl, C₁-C₆haloalkyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkynyl, C₂-C₆haloalkynyl, C₃-C₄cycloalkylC₁-C₂alkyl-, C₃-C₄cycloalkylC₁-C₂alkyl- wherein the C₃-C₄cycloalkyl group is substituted with 1 or 2 halogen atoms, oxetan-3-yl-CH₂-, benzyl or benzyl substituted with halogen or C₁-C₆haloalkyl;

R₂ is selected from phenyl, pyridine, pyrimidine, pyrazine, pyridazine, and phenyl, pyridine, pyrimidine, pyrazine and pyridazine, each of which is substituted with one to three substituents, provided the substituent(s) are not on either carbon adjacent to the carbon C=X is attached, and each substituent is independently selected from: C₁-C₃alkyl, C₁-C₃haloalkyl, C₁-C₃haloalkylthio, C₁-C₃alkoxy, C₁-C₃haloalkoxy, halogen, SF₅, CN, CONH₂, and C(S)NH₂; R₃ is C₁-C₃alkyl or C₁-C₃haloalkyl;

A₂ is CR_{4b} or N;

R_{4b} is hydrogen, or halogen;

R_{4a} is cyano, or C₁-C₃haloalkoxy;

R_{5a} and R_{5b} are, independently of each other, selected from hydrogen, halogen, CN, C₁-C₃alkyl, C₁-C₃haloalkyl, C₃-C₄cycloalkyl, C₁-C₃alkoxy, and C₁-C₃haloalkoxy; or agrochemically acceptable salts, stereoisomers, enantiomers, tautomers and N-oxides of the compounds of formula I.

-2-

Compounds of formula I which have at least one basic centre can form, for example, acid addition salts, for example with strong inorganic acids such as mineral acids, for example perchloric acid, sulfuric acid, nitric acid, nitrous acid, a phosphorus acid or a hydrohalic acid, with strong organic carboxylic acids, such as C₁-C₄alkanecarboxylic acids which are unsubstituted or substituted, for example by halogen, for example acetic acid, such as saturated or unsaturated dicarboxylic acids, for example oxalic acid, malonic acid, succinic acid, maleic acid, fumaric acid or phthalic acid, such as hydroxycarboxylic acids, for example ascorbic acid, lactic acid, malic acid, tartaric acid or citric acid, or such as benzoic acid, or with organic sulfonic acids, such as C₁-C₄alkane- or arylsulfonic acids which are unsubstituted or substituted, for example by halogen, for example methane- or p-toluenesulfonic acid. Compounds of formula I which have at least one acidic group can form, for example, salts with bases, for example mineral salts such as alkali metal or alkaline earth metal salts, for example sodium, potassium or magnesium salts, or salts with ammonia or an organic amine, such as morpholine, piperidine, pyrrolidine, a mono-, di- or tri-lower-alkylamine, for example ethyl-, diethyl-, triethyl- or dimethylpropylamine, or a mono-, di- or trihydroxy-lower-alkylamine, for example mono-, di- or triethanolamine.

In each case, the compounds of formula I according to the invention are in free form, in oxidized form as a N-oxide or in salt form, e.g. an agronomically usable salt form.

N-oxides are oxidized forms of tertiary amines or oxidized forms of nitrogen containing heteroaromatic compounds. They are described for instance in the book "*Heterocyclic N-oxides*" by A. Albini and S. Pietra, CRC Press, Boca Raton 1991.

The compounds of formula I according to the invention also include hydrates which may be formed during the salt formation.

The term "C₁-C_nalkyl" as used herein refers to a saturated straight-chain or branched hydrocarbon radical attached via any of the carbon atoms having 1 to n carbon atoms, for example, any one of the radicals methyl, ethyl, n-propyl, 1-methylbutyl, 2-methylbutyl, 3-methylbutyl, 2, 2-dimethylpropyl, 1-ethylpropyl, n-hexyl, n-pentyl, 1,1-dimethylpropyl, 1,2-dimethylpropyl, 1-methylpentyl, 2-methylpentyl, 3-methylpentyl, 4-methylpentyl, 1,1-dimethylbutyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,2-dimethylbutyl, 2,3-dimethylbutyl, 3,3-dimethylbutyl, 1-ethylbutyl, 2-ethylbutyl, 1,1,2-trimethylpropyl, 1,2,2-trimethylpropyl, 1-ethyl-1-methylpropyl, or 1-ethyl-2-methylpropyl.

The term "C₁-C_nhaloalkyl" as used herein refers to a straight-chain or branched saturated alkyl radical attached via any of the carbon atoms having 1 to n carbon atoms (as mentioned above), where some or all of the hydrogen atoms in these radicals may be replaced by fluorine, chlorine, bromine and/or iodine, i.e., for example, any one of chloromethyl, dichloromethyl, trichloromethyl, fluoromethyl, difluoromethyl, trifluoromethyl, chlorofluoromethyl, dichlorofluoromethyl, chlorodifluoromethyl, 2-

-3-

fluoroethyl, 2-chloroethyl, 2-bromoethyl, 2-iodoethyl, 2,2-difluoroethyl, 2,2,2-trifluoroethyl, 2-chloro-2-fluoroethyl, 2-chloro-2,2-difluoroethyl, 2,2-dichloro-2-fluoroethyl, 2,2,2-trichloroethyl, pentafluoroethyl, 2-fluoropropyl, 3-fluoropropyl, 2,2-difluoropropyl, 2,3-difluoropropyl, 2-chloropropyl, 3-chloropropyl, 2,3-dichloropropyl, 2-bromopropyl, 3-bromopropyl, 3,3,3-trifluoropropyl, 3,3,3-trichloropropyl, 2,2,3,3,3-pentafluoropropyl, heptafluoropropyl, 1-(fluoromethyl)-2-fluoroethyl, 1-(chloromethyl)-2-chloroethyl, 1-(bromomethyl)-2-bromoethyl, 4-fluorobutyl, 4-chlorobutyl, 4-bromobutyl or nonafluorobutyl. According to a term "C₁-C₂fluoroalkyl" would refer to a C₁-C₂alkyl radical which carries 1, 2, 3, 4, or 5 fluorine atoms, for example, any one of difluoromethyl, trifluoromethyl, 1-fluoroethyl, 2-fluoroethyl, 2,2-difluoroethyl, 2,2,2-trifluoroethyl, 1,1,2,2-tetrafluoroethyl or pentafluoroethyl.

The term "C₁-C_nalkoxy" as used herein refers to a straight-chain or branched saturated alkyl radical having 1 to n carbon atoms (as mentioned above) which is attached via an oxygen atom, i.e., for example, any one of the radicals methoxy, ethoxy, n-propoxy, 1-methylethoxy, n-butoxy, 1-methylpropoxy, 2-methylpropoxy or 1,1-dimethylethoxy. The term "haloC₁-C_nalkoxy" as used herein refers to a C₁-C_nalkoxy radical where one or more hydrogen atoms on the alkyl radical is replaced by the same or different halogen atom(s) - examples include trifluoromethoxy, 2-fluoroethoxy, 3-fluoropropoxy, 3,3,3-trifluoropropoxy, 4-chlorobutoxy.

The term "C₁-C_ncyanoalkyl" as used herein refers to a straight chain or branched saturated C₁-C_nalkyl radical having 1 to n carbon atoms (as mentioned above), where one of the hydrogen atoms in these radicals is replaced by a cyano group: for example, cyanomethyl, 2-cyanoethyl, 2-cyanopropyl, 3-cyanopropyl, 1-(cyanomethyl)-2-ethyl, 1-(methyl)-2-cyanoethyl, 4-cyanobutyl, and the like.

The term "C₃-C_ncycloalkyl" as used herein refers to 3-n membered cycloalkyl groups such as cyclopropane, cyclobutane, cyclopentane and cyclohexane.

The term "C₃-C_ncycloalkylC₁-C_nalkyl" as used herein refers to 3 to n membered cycloalkyl group with an alkyl radical, which alkyl radical is connected to the rest of the molecule. In the instance, the C₃-C_ncycloalkylC₁-C₂alkyl- group is substituted, the substituent(s) can be on the cycloalkyl group or alkyl radical.

The term "aminocarbonylC₁-C_nalkyl" as used herein refers to an alkyl radical where one of the hydrogen atoms in the radical is replaced by CONH₂ group.

The term "hydroxycarbonylC₁-C_nalkyl" as used herein refers to an alkyl radical where one of the hydrogen atoms in the radical is replaced by COOH group.

The term "C₁-C_nnitroalkyl" as used herein refers to an alkyl radical where one of the hydrogen atoms in the radical is replaced by NO₂ group.

The term "C₁-C_nalkylsulfanyl" or "C₁-C_nhaloalkylthio" as used herein refers to a C₁-C_nalkyl moiety linked through a sulfur atom. Similarly, the term "C₁-C_nhaloalkylsulfanyl" as used herein refers to a C₁-C_nhaloalkyl moiety linked through a sulfur atom.

5

The term "C₁-C_nalkylsulfinyl" as used herein refers to a C₁-C_nalkyl moiety linked through the sulfur atom of the S(=O) group. Similarly, the term "C₁-C_nhaloalkylsulfinyl" as used herein refers to a C₁-C_nhaloalkyl moiety linked through the sulfur atom of the S(=O) group.

- 10 The term "C₁-C_nalkylsulfonyl" as used herein refers to a C₁-C_nalkyl moiety linked through the sulfur atom of the S(=O)₂ group. Similarly, the term "C₁-C_nhaloalkylsulfonyl" as used herein refers to a C₁-C_nhaloalkyl moiety linked through the sulfur atom of the S(=O)₂ group

- 15 The term "trimethylsilaneC₁-C_nalkyl" as used herein refers to an alkyl radical where one of the hydrogen atoms in the radical is replaced by a -Si(CH₃)₃ group.

The term "C₂-C_nalkenyl" as used herein refers to a straight or branched alkenyl chain having from two to n carbon atoms and one or two double bonds, for example, ethenyl, prop-1-enyl, but-2-enyl.

- 20 The term "C₂-C_nhaloalkenyl" as used herein refers to a C₂-C_nalkenyl moiety substituted with one or more halogen atoms which may be the same or different.

The term "C₂-C_nalkynyl" as used herein refers to a straight or branched alkynyl chain having from two to n carbon atoms and one triple bond, for example, ethynyl, prop-2-ynyl, but-3-ynyl,

25

The term "C₂-C_nhaloalkynyl" as used herein refers to a C₂-C_nalkynyl moiety substituted with one or more halogen atoms which may be the same or different.

- 30 Halogen is generally fluorine, chlorine, bromine or iodine. This also applies, correspondingly, to halogen in combination with other meanings, such as haloalkyl

The pyridine, pyrimidine, pyrazine and pyridazine groups (unsubstituted or substituted) for R₂ and R₄ are each connected via a carbon atom on the respective ring to the rest of the compound.

- 35 As used herein, the term "controlling" refers to reducing the number of pests, eliminating pests and/or preventing further pest damage such that damage to a plant or to a plant derived product is reduced.

The staggered line as used herein, for example, in M-1 and L-1, represent the point of connection/attachment to the rest of the compound.

-5-

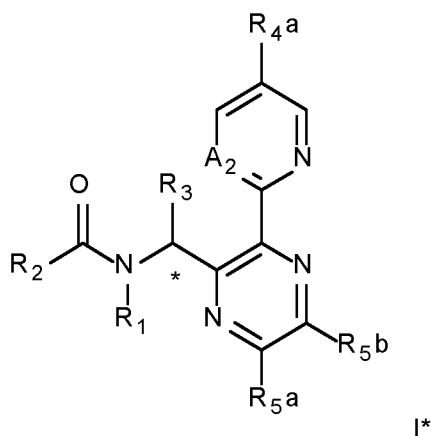
As used herein, the term "pest" refers to insects, acarines, nematodes and molluscs that are found in agriculture, horticulture, forestry, the storage of products of vegetable origin (such as fruit, grain and timber); and those pests associated with the damage of man-made structures. The term pest
 5 encompasses all stages in the life cycle of the pest.

As used herein, the term "effective amount" refers to the amount of the compound, or a salt thereof, which, upon single or multiple applications provides the desired effect.

10 An effective amount is readily determined by the skilled person in the art, by the use of known techniques and by observing results obtained under analogous circumstances. In determining the effective amount a number of factors are considered including, but not limited to: the type of plant or derived product to be applied; the pest to be controlled & its lifecycle; the particular compound applied; the type of application; and other relevant circumstances.

15

As one of ordinary skill in the art will appreciate, compounds of formula I contain a stereogenic centre which is indicated with an asterisk in the structure below:

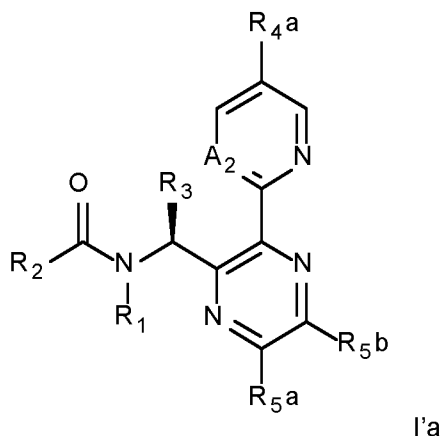


where R₁, R₂, R₃, R_{4a}, R_{5a}, R_{5b}, and A₂ are as defined in the first aspect.

20

The present invention contemplates both racemates and individual enantiomers. Compounds having preferred stereochemistry are set out below.

-6-



Particularly preferred compounds of the present invention are compounds of formula I'a:
 where R₁, R₂, R₃, R_{4a}, R_{5a}, R_{5b}, and A₂ are as defined in the first aspect, and stereoisomers,
 enantiomers, tautomers and N-oxides of the compounds of formula (I'a), and agrochemically
 5 acceptable salts thereof.

The term "optionally substituted" as used herein means that the group referenced is either
 unsubstituted or is substituted by a designated substituent, for example, "C₃-C₄cycloalkyl is optionally
 substituted with 1 or 2 halogen atoms" means C₃-C₄cycloalkyl, C₃-C₄cycloalkyl substituted with 1
 10 halogen atom and C₃-C₄cycloalkyl substituted with 2 halogen atoms.

Embodiments according to the invention are provided as set out below.

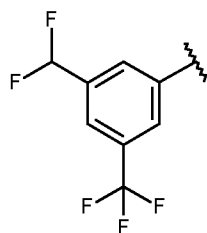
In an embodiment of each aspect of the invention, R₁ is

- 15 A. hydrogen, methyl, ethyl, n-propyl, isobutyl, cyclopropylmethyl or HCH≡CCH₂-; or
- B. hydrogen, methyl, or cyclopropylmethyl; or
- C. hydrogen; or
- D. methyl; or
- E. cyclopropylmethyl.

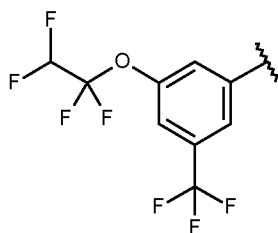
In an embodiment of each aspect of the invention, R₂ is

- 20 A. phenyl, pyridine, pyrimidine, pyrazine, pyridazine, or the phenyl, pyridine, pyrimidine, pyrazine
 and pyridazine, each of which is substituted with one to three substituents, provided the
 substituent(s) are not on either carbon adjacent to the carbon C=X is attached, and each
 25 substituent is independently selected from: C₁-C₃haloalkyl, C₁-C₃haloalkylthio, C₁-C₃alkoxy,
 C₁-C₃haloalkoxy, and halogen; or
- B. one of M-1 to M-13

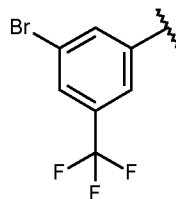
-7-



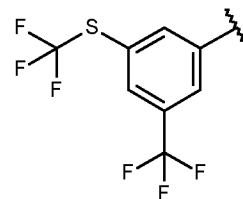
M-1



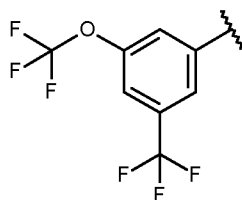
M-2



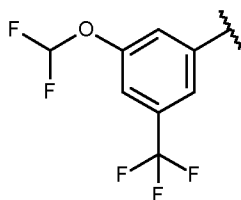
M-3



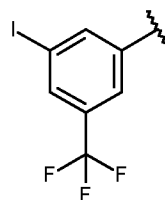
M-4



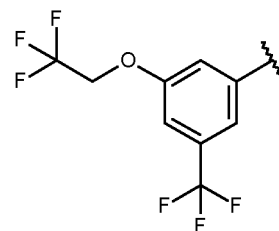
M-5



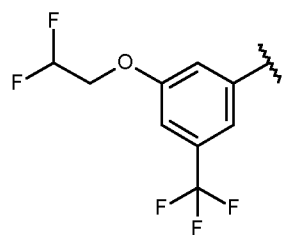
M-6



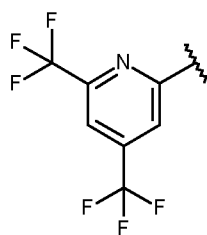
M-7



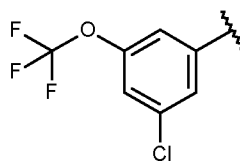
M-8



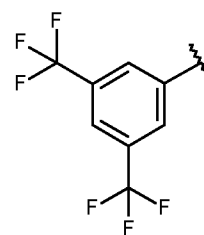
M-9



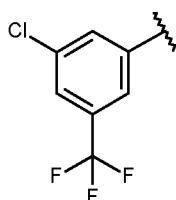
M-10



M-11



M-12



M-13

C. one of M-3, M-4, M-5, M-6, M-8, M-9, M-11 M-12 or M-13; or

D. one of M-3, M-4, M-5, M-6, M-8, M-9, M-12 or M-13; or

5 E. one of M-3, M-4, M-6, M-8; M-9, M-12 or M-13; or

F. one of M-3, M-6, M-8, M-9, M-12 or M-13; or

G. one of M-3, M-8, M-12 or M-13; or

H. one of M-6, M-9 or M-12; or

I. one of M-3, M-8 or M-12; or

10 J. one of M-3, M-12 or M-13 or

K. M-12 or M-13.

In an embodiment of each aspect of the invention, R₃ is

-8-

- A. C₁-C₃alkyl or C₁-C₃haloalkyl; or
- B. methyl.

In an embodiment of each aspect of the invention, A₂ is

- 5 A. N; or
- B. C-R_{4b}, where R_{4b} is hydrogen or halogen (such as Cl, F, Br and I); preferably R_{4b} is hydrogen.

In an embodiment of each aspect of the invention, R_{4a} is

- A. cyano, or C₁-C₃fluoroalkoxy; or
- 10 B. cyano, trifluoromethoxy, difluoromethoxy, 2,2,2-trifluoroethoxy, or 2,2-difluoroethoxy.

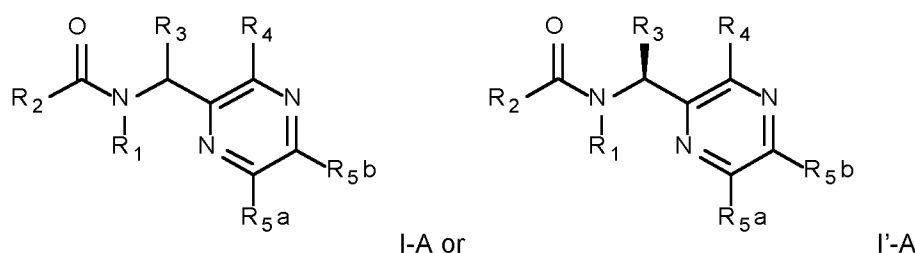
In an embodiment of each aspect of the invention, R_{5a} and R_{5b}, independent of each other, are

- A. hydrogen, halogen, C₁-C₃alkyl, or C₁-C₃alkoxy; or
- B. selected from hydrogen, bromo, chloro, methyl, and methoxy; or
- 15 C. hydrogen.

The present invention, accordingly, makes available a compound of formula I having the substituents R₁, R_{2a}, R_{2b}, R₃, R_{4a}, R_{5a}, R_{5b}, A₁ and A₂ as defined above in all combinations / each permutation.

Accordingly, made available, for example, is a compound of formula I with R₁ being embodiment B (i.e. hydrogen, methyl, cyclopropylmethyl); R₂ being an embodiment C (i.e. one of M-3, M-4, M-5, M-6, M-8, M-9, M-11 M-12 or M-13); R₃ being embodiment B (i.e. methyl); A₂ being embodiment B (i.e. C-R_{4b}, where R_{4b} is hydrogen or halogen (such as Cl, F, Br and I); preferably R_{4b} is hydrogen); R_{4a} being embodiment A (i.e. cyano, or C₁-C₃fluoroalkoxy); and R_{5a} being embodiment A (i.e. selected from hydrogen, halogen, C₁-C₃alkyl, and C₁-C₃alkoxy); and R_{5b} being embodiment C (i.e. hydrogen).

In an embodiment, the compound of formula I can be represented as

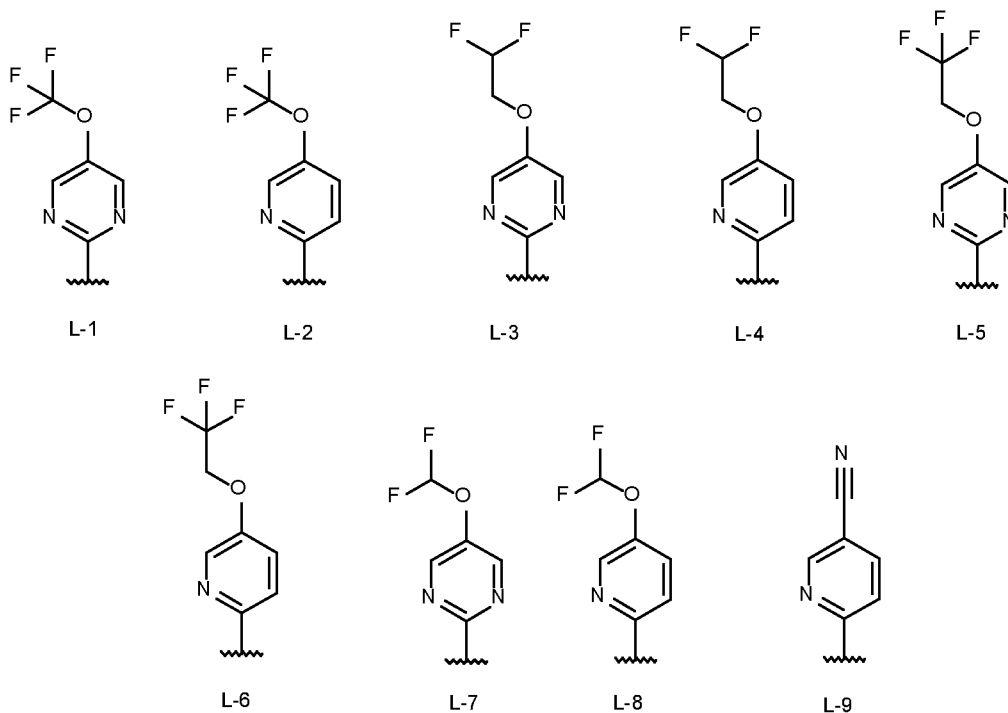


wherein R₁, R₂, R_{5a}, and R_{5b} are as defined in the first aspect, and R₄ is the cyclic group containing A₂ and the substituent R_{4a} as defined in the first aspect.

In an embodiment of each aspect of the invention, the R₄ (the cyclic group containing A₂ and the substituent R_{4a}) is

- A. selected from L-1 to L-9

-9-



B. selected from L-1, L-2, L-3, L-4, L-5, L-6, L-7, L-8, and L-9; or

C. selected from L-1, L-3, L-5, L-6, L-7, L-8 and L-9; or

D. selected from L-1, L-2, L-7, L-8 and L-9; or

5 E. selected from L-3, L-4, L-7, L-8 and L-9; or

F. selected from L-1, L-3, L-5, L-7, L-8 and L-9; or

G. selected from L-1, L-5, L-7, L-8 and L-9; or

H. L-7, L-8 or L-9; or

I. L-3, L-7 or L-9; or

10 J. L-8 or L-9; or

K. L-5 or L-9; or

L. L-7 or L-9; or

M. L-1 or L-9; or

N. L-9.

15

L-1, L-5, L-7, L-8 and L-9

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, ethyl, n-propyl, isobutyl, cyclopropylmethyl or HCH≡CCH₂-; as R₂ one of M-1 to M-13; as R₃

20 methyl; as R₄ one of L-1 to L-9; and as R_{5a} and R_{5b}, independently selected from hydrogen, halogen, C₁-C₃alkyl, and C₁-C₃alkoxy.

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-1 to M-13; as R₃ methyl; as R₄ one of L-1 to L-9; and as

25 R_{5a} and R_{5b}, independently selected from hydrogen, halogen, C₁-C₃alkyl, and C₁-C₃alkoxy.

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen; as R₂ one of M-1 to M-13; as R₃ methyl; as R₄ one of L-1 to L-9; and as R_{5a} and R_{5b}, independently selected from hydrogen, halogen, C₁-C₃alkyl, and C₁-C₃alkoxy.

5

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-4, M-5, M-6, M-8, M-9, M-11 M-12 or M-13; as R₃ methyl; as R₄ one of L-1 to L-9; and as R_{5a} and R_{5b}, independently selected from hydrogen, halogen, C₁-C₃alkyl, and C₁-C₃alkoxy.

10

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-4, M-5, M-6, M-8, M-9, M-11 M-12 or M-13; as R₃ methyl; as R₄ one of L-1, L-2, L-3, L-4, L-5, L-6, L-7, L-8, or L-9; and as R_{5a} and R_{5b}, independently selected from hydrogen, halogen, C₁-C₃alkyl, and C₁-C₃alkoxy.

15

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-4, M-5, M-6, M-8, M-9, M-11, M-12 or M-13; as R₃ methyl; as R₄ one of L-1, L-2, L-3, L-4, L-5, L-6, L-7, L-8, or L-9; and as R_{5a} and R_{5b}, each hydrogen.

20

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-6, M-8, M-9, M-12 or M-13; as R₃ methyl; as R₄ one of L-1 to L-9; and as R_{5a} and R_{5b}, each hydrogen.

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-6, M-8, M-9, M-12 or M-13; as R₃ methyl; as R₄ one of L-1, L-2, L-3, L-4, L-5, L-6, L-7, L-8, or L-9; and as R_{5a} and R_{5b}, each hydrogen.

25

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-4, M-6, M-8, M-9, or M-12; as R₃ methyl; as R₄ one of L-1, L-2, L-7, L-8 and L-9; and as R_{5a} and R_{5b}, each hydrogen.

30

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-4, M-6, M-8, M-9, or M-12; as R₃ methyl; as R₄ one of L-3, L-4, L-7, L-8 and L-9; and as R_{5a} and R_{5b}, each hydrogen.

35

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-8, M-12, or M-13; as R₃ methyl; as R₄ one of L1 to L9; and as R_{5a} and R_{5b}, each hydrogen.

-11-

In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-8, M-12, or M-13; as R₃ methyl; as R₄ one of L-1, L-2, L-7, L-8 and L-9; and as R_{5a} and R_{5b}, each hydrogen.

- 5 In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-8, M-12, or M-13; as R₃ methyl; as R₄ one of L-3, L-4, L-7, L-8 and L-9; and as R_{5a} and R_{5b}, each hydrogen.

- 10 In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-8, M-12, or M-13; as R₃ methyl; as R₄ one of L7, L8 or L9; and as R_{5a} and R_{5b}, each hydrogen.

- 15 In an embodiment of each aspect of the invention, the compound of formula I has as R₁ hydrogen, methyl, or cyclopropylmethyl; as R₂ one of M-3, M-8, or M-12; as R₃ methyl; as R₄ L9; and as R_{5a} and R_{5b}, each hydrogen.

- 20 In a second aspect, the present invention makes available a composition comprising a compound of formula I as defined in the first aspect, one or more auxiliaries and diluent, and optionally one more other active ingredient.

- In a third aspect, the present invention makes available a method of combating and controlling insects, acarines, nematodes or molluscs which comprises applying to a pest, to a locus of a pest, or to a plant susceptible to attack by a pest an insecticidally, acaricidally, nematocidally or molluscicidally effective amount of a compound as defined in the first aspect or a composition as defined in the second aspect.

- 25 In a fourth aspect, the present invention makes available a method for the protection of plant propagation material from the attack by insects, acarines, nematodes or molluscs, which comprises treating the propagation material or the site, where the propagation material is planted, with an effective amount of a compound of formula I as defined in the first aspect or a composition as defined in the second aspect.

- 30 In a fifth aspect, the present invention makes available a plant propagation material, such as a seed, comprising, or treated with or adhered thereto, a compound of formula I as defined in the first aspect or a composition as defined in the second aspect.

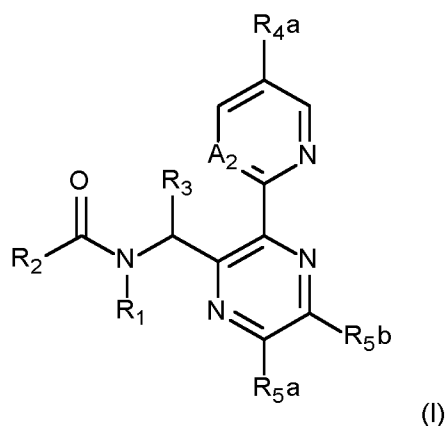
- 35 The present invention in a further aspect provides a method of controlling parasites in or on an animal in need thereof comprising administering an effective amount of a compound of the first aspect. The present invention further provides a method of controlling ectoparasites on an animal in need thereof comprising administering an effective amount of a compound of formula I as defined on the first

-12-

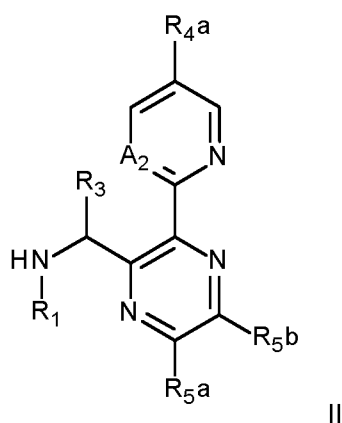
aspect. The present invention further provides a method for preventing and/or treating diseases transmitted by ectoparasites comprising administering an effective amount of a compound of formula I as defined in the first aspect, to an animal in need thereof.

- 5 Compounds of formula I can be prepared by those skilled in the art following known methods. More specifically compounds of formulae I, and I'a, and intermediates therefor can be prepared as described below in the schemes and examples. Certain stereogenic centers have been left unspecified for the clarity and are not intended to limit the teaching of the schemes in any way.
- 10 The process according to the invention for preparing compounds of formula I is carried out by methods known to those skilled in the art.

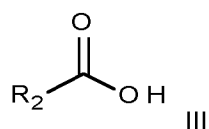
Compounds of formula I



- 15 can be prepared by reaction of an amine of formula II



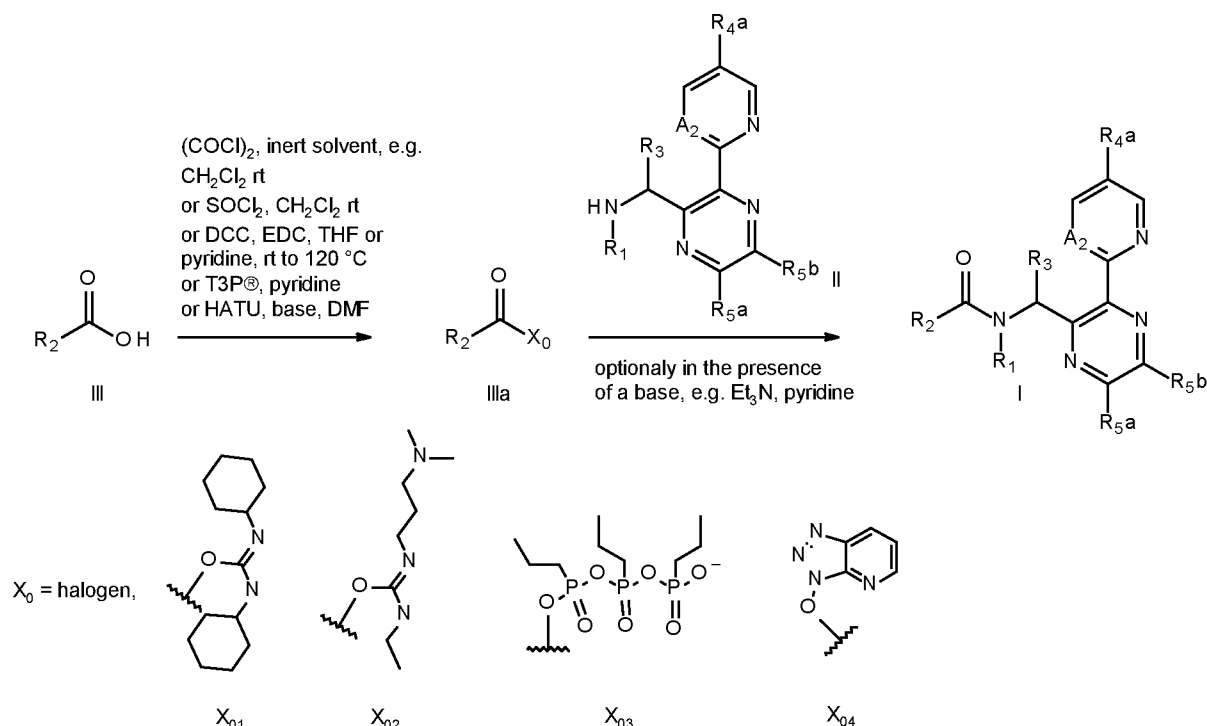
wherein R₁, R₃, A₂, R_{4a}, R_{5a}, and R_{5b} are as defined in formula I, with a carboxylic acid derivative of formula III



- 20 wherein R₂ is as defined in formula I. The chemistry is described in more detail in Scheme 1.

-13-

Scheme 1:

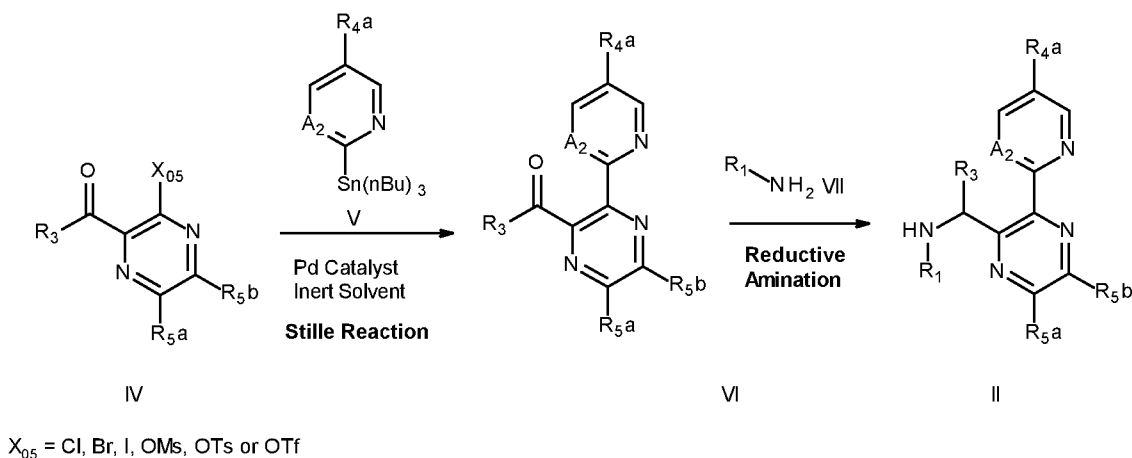


In Scheme 1 compounds of formula III, wherein R₂ is as defined in formula I, are activated to compounds of formula IIIa by methods known to those skilled in the art and described for example in Tetrahedron, 61 (46) , 10827-10852, 2005. For example, compounds where X₀ is halogen are formed by treatment of compounds of formula III with for example, oxalyl chloride or thionyl chloride in the presence of catalytic quantities of DMF in inert solvents such as methylene dichloride or THF at temperatures between 20 °C to 100 °C., preferably 25 °C. Treatment of IIIa with compounds of formula II, wherein R₁, R₃, A₂, R_{4a}, R_{5a}, and R_{5b}, are as defined in formula I, optionally in the presence of a base, e.g. triethylamine or pyridine leads to compounds of formula I. Alternatively, compounds of formula I can be prepared by treatment of compounds of formula III with dicyclohexyl carbodiimide (DCC) or 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDC) to give the activated species IIIa, wherein X₀ is X₀₁ or X₀₂, in an inert solvent, e.g. pyridine, or THF optionally in the presence of a base, e.g. triethylamine, at temperatures between 50-180 °C. In addition, an acid of the formula III can also be activated by reaction with a coupling reagent such as propanephosphonic acid anhydride (T3P®) or O-(7-Aza-1-benzotriazolyl)-N,N,N',N'-tetramethyluronium-hexafluorophosphat (HATU) to provide compounds of formula IIIa wherein X₀ is X₀₃ and X₀₄ as described for example in Synthesis 2013, 45, 1569 and Journal Prakt. Chemie 1998, 340, 581. Subsequent reaction with an amine of the formula II provides compounds of formula I.

Intermediates of formula II, wherein R₁, R₃, A₂, R_{4a}, R_{5a} and R_{5b} are as defined in formula I, can be prepared according to Scheme 2:

-14-

Scheme 2:



In Scheme 2, compounds of formula II, wherein R_1 , R_3 , A_2 , R_{4a} , R_{5a} and R_{5b} are as defined in formula I, can be prepared by treatment of compounds of formula VI, wherein R_3 , A_2 , R_{4a} , R_{5a} and R_{5b} are as defined in formula I, with compounds of formula VII (wherein R_1 is as defined in formula I), e.g. in the presence of $\text{NaBH}(\text{OAc})_3$ or NaBH_3CN , preferably with NaBH_3CN as reductive reagent, in a suitable solvent, preferably in acetic acid at room temperature analog to WO2002/088073, page 35.

Alternatively, another reagent system for the reductive amination uses a combination of $\text{Ti}(\text{i-OPr})_4$ and NaBH_4 in the presence of an amine of formula VII can also provide compounds of formula II (see Synthesis 2003 (14), 2206).

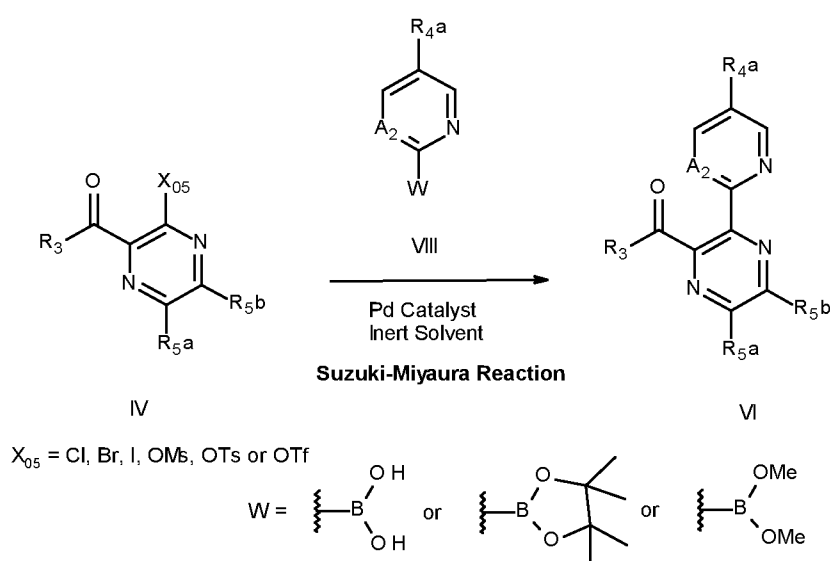
Compounds of formula VI, wherein R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I, can be prepared by a Stille reaction between compounds of formula IV, wherein X_{05} is a leaving group, such as chlorine, bromine, iodine, arylsulfonate, alkylsulfonate or trifluoromethanesulfonate and R_3 , R_{5a} and R_{5b} are as defined in formula I, and tin compounds of formula V (wherein A_2 and R_{4a} are as defined in formula I), in the presence of a palladium catalyst, for example tetrakis(triphenylphosphine)palladium(0), or (1,1'-bis(diphenylphosphino)-ferrocene)dichloropalladium-dichloromethane (1:1 complex), in an inert solvent, such as DMF, acetonitrile, or dioxane, optionally in the presence of an additive, such as potassium, cesium fluoride, or lithium chloride, and optionally in the presence of a further catalyst, for example copper(I)iodide. Such Stille coupling reactions are well known to those skilled in the art, and have been described in for example J. Org. Chem., 2005, 70, 8601, J. Org. Chem., 2009, 74, 5599, Angew. Chem. Int. Ed., 2004, 43, 1132, Heterocycles 2010, 80, 1215 and J. Am. Chem. Soc. 2004, 126, 16433.

Alternatively, compounds of formula VI (wherein R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I) can also be prepared by a Suzuki reaction (Scheme 3), which involves for example, reacting compounds of formula IV (wherein R_3 , R_{5a} , and R_{5b} are as defined in formula I and X_{05} is a leaving group like, for example, chlorine, bromine, iodine, arylsulfonate, alkylsulfonate or trifluoromethanesulfonate) with compounds of formula VIII, wherein W can be a boron-derived functional group, as for example $\text{B}(\text{OH})_2$ or a pinacol boronic ester. The reaction can be catalyzed by a palladium based catalyst, for example

-15-

tetrakis(triphenylphosphine)-palladium or (1,1'-bis(diphenylphosphino)-ferrocene)dichloropalladium-dichloromethane (1:1 complex), in presence of a base, like sodium carbonate or cesium fluoride, in a solvent or a solvent mixture, like, for example a mixture of 1,2-dimethoxyethane and water, dioxane and water, or DMF and water preferably under inert atmosphere. The reaction temperature can preferentially range from room temperature to the boiling point of the reaction mixture. Such Suzuki reactions are well known to those skilled in the art and have been reported, for example in J. Organomet. Chem. 576, 1999, 147–168, Science of Synthesis 2010, 45b, 547, Eur. J. Org. Chem. 2012, (31), 6248 and Synthesis 2017, 49, 4372.

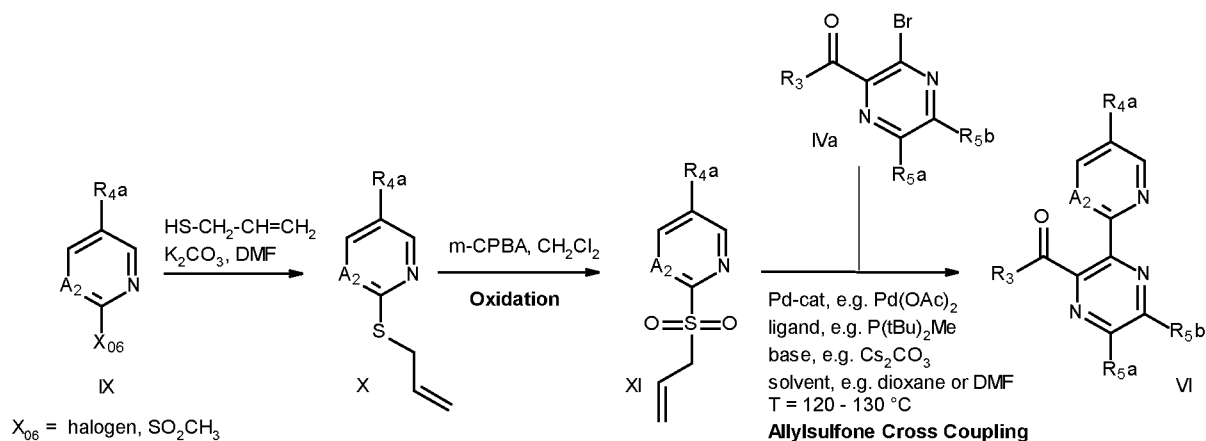
10 Scheme 3:



Compounds of formula IV are generally commercially available.

Alternatively compounds of formula VI, wherein R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I, is outlined in Scheme 4.

Scheme 4:



-16-

For example, compounds of formula VI, wherein R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I, may be prepared by allyl sulfone coupling reaction of compounds of formula IVa (wherein R_3 , R_{5a} , and R_{5b} are as defined in formula I), with compounds of formula XI, wherein A_2 and R_{4a} are defined in formula I, in suitable solvents, preferable dioxane or DMF, in the presence of a Pd-catalyst, preferable palladium acetate, a ligand, e.g. ditert-butyl(methyl)phosphane, and a base, e.g. Cs_2CO_3 usually upon heating at temperatures between 120 to 130 °C. Such processes have been described, for example, in J. Am. Chem. Soc. 2018, 140, 15916.

The required intermediates of formula XI can be obtained from compounds of formula IX (wherein A_2 and R_{4a} are defined as for formula I and X_{06} is halogen or methyl sulfone) through nucleophilic substitution with prop-2-ene-1-thiol and subsequent oxidation with mCPBA. Such transformation are well known and reported, for instance, in J. Am. Chem. Soc. 2018, 140, 15916.

Alternatively compounds of formula VI, wherein R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I, is outlined in Scheme 5.

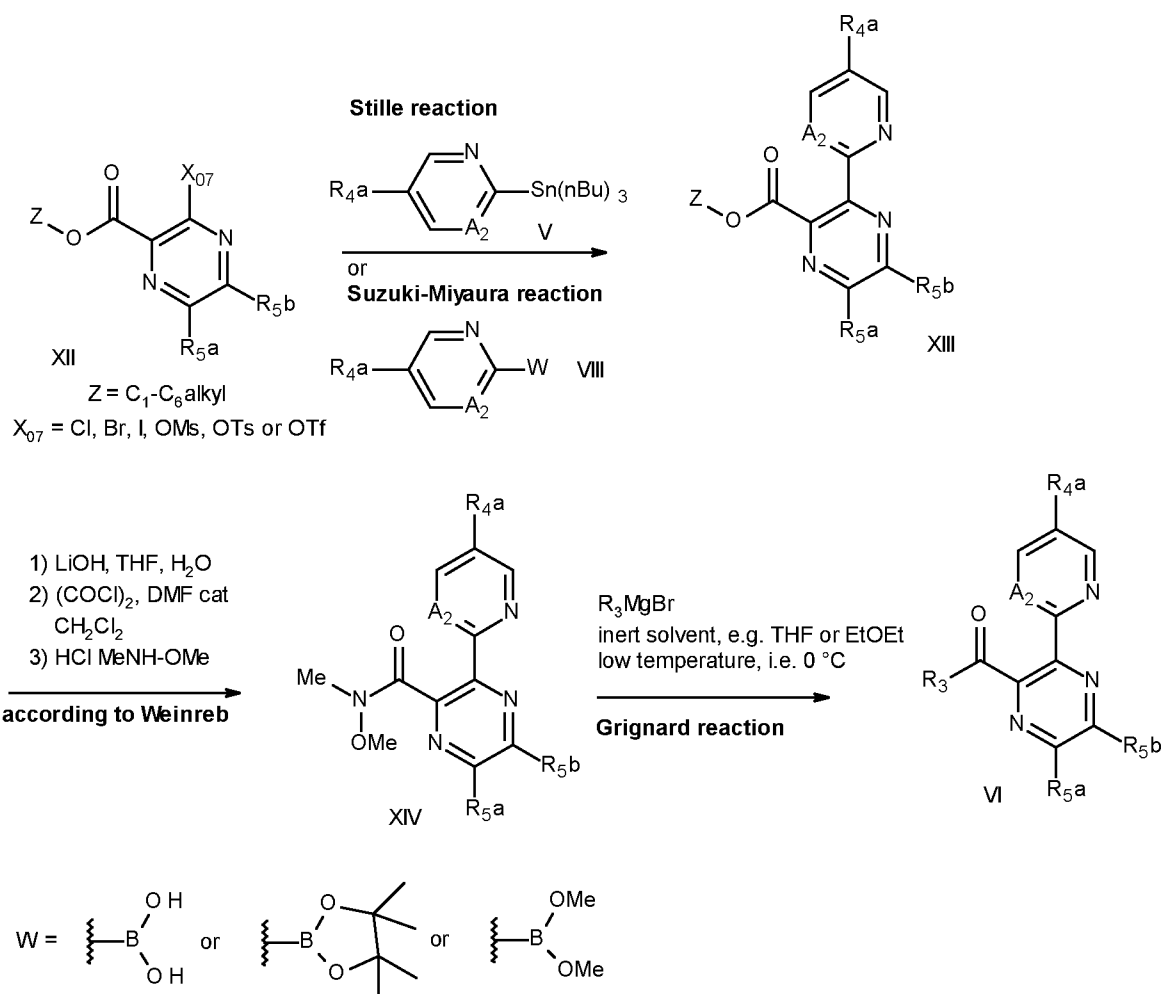
Compounds of formula VI (wherein R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I) may be prepared by treatment of compounds of formula XIV with a Grignard reagent R_3MgBr , e.g. $MeMgBr$ at lower temperatures, preferable at 0 to 25 °C in a suitable solvent such as THF or diethyl ether.

Weinreb amides of formula XIV (wherein A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I) can be prepared in three steps from compounds of formula XIII, wherein A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I and Z is C_1 - C_6 alkyl. Compounds of formula XIII are converted to carboxylic acids by methods known in the art (see e.g. WO2011/143365, page 138) and activation (see Scheme 1) of the subsequent carboxylic acids followed by treatment with N-methoxy-N-methylamine (according to Weinreb et al. Tet. Lett. 1981, 39, 3815) lead to compound of formula XIV.

Compounds of formula XIII, wherein A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I and Z is C_1 - C_6 alkyl, can be prepared by reaction of compounds of formula XII (wherein R_{5a} , and R_{5b} are as defined in formula I, Z is C_1 - C_6 alkyl, and X_{07} is a leaving group like, for example, chlorine, bromine, iodine) with compounds of formula V (Stille reaction) or compounds of formula VIII (Suzuki-Miyaura reaction) in the presence of a palladium catalyst as described in detail in Schemes 2 and 3.

Scheme 5:

-17-



In an alternative process (Scheme 6), compounds of formula II, wherein R₁, R₃, A₂, R_{4a}, R_{5a}, and R_{5b} are defined as in formula I, may be prepared by nucleophilic substitution reaction of compound of formula XVI, wherein R₃, A₂, R_{4a}, R_{5a}, and R_{5b} are as defined in formula I and X₀₈ is Cl, Br, OMs, OTs or OTf, with amines of formula VII, in suitable solvents that may include, for example, acetonitrile or DMF, in the presence of a suitable base, such as sodium, potassium or caesium carbonate, usually upon heating at temperatures between room temperature and 200°C, preferably between 40°C to the boiling point of the reaction mixture, optionally under microwave heating conditions.

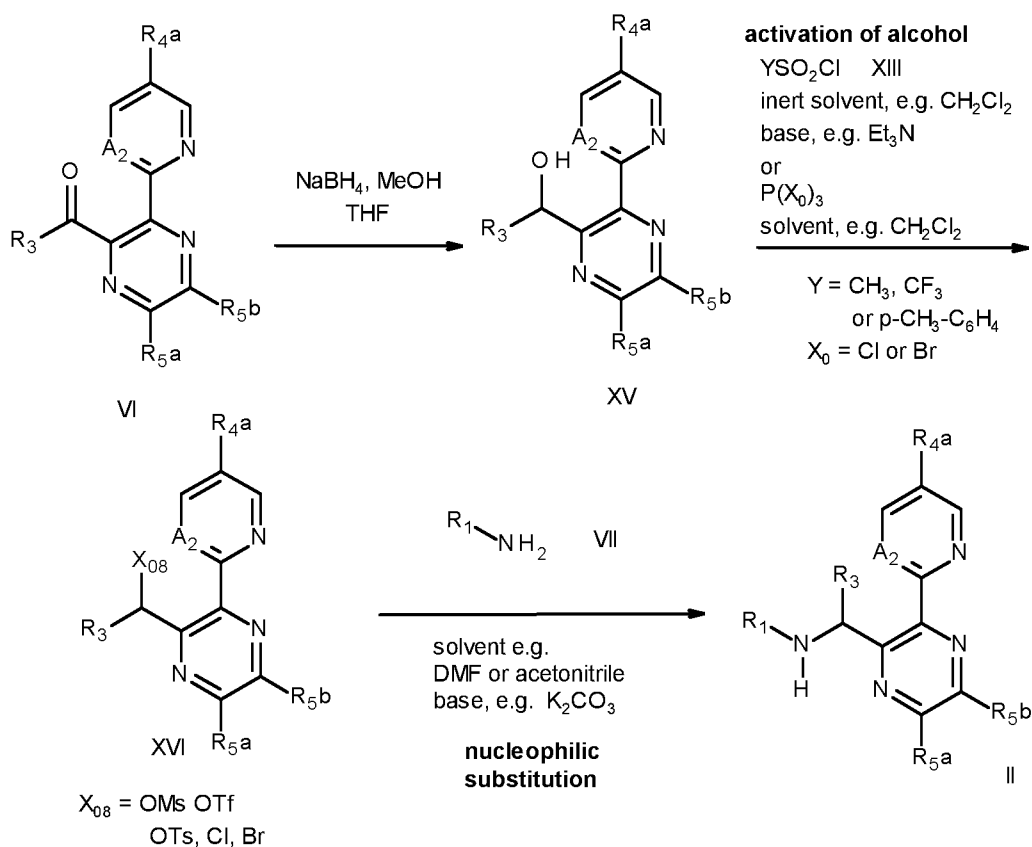
- 10 Compounds of formula XVI, wherein R₃, A₂, R_{4a}, R_{5a}, R_{5b} are defined as in formula I and X₀₈ is Cl, Br, OMs, OTs or OTf, may be prepared by activation of the alcohols of formula XV, wherein R₃, A₂, R_{4a}, R_{5a} and R_{5b} are defined as in formula I, with compounds of formula XIII, wherein Y is CH₃, CF₃ or p-CH₃-C₆H₄, in an inert solvent, preferably in methylene dichloride and in the presence of a base, e.g. triethyl amine. Alcohols of formula XV may be also be activated to alkyl halides XVI (wherein X₀₈ is Cl or Br) by
- 15 treatment with phosphorous compounds, e.g. P(X₀)₃, wherein X₀ is chlorine or bromine by methods known to those skilled in the art. Such general functional group transformations are described for example in *Organische Chemie*. 4. Auflage, Wiley-VCH Verlag, Weinheim 2005, p. 393 ff and *Chem Commun.* 2014, 50, 5756.

-18-

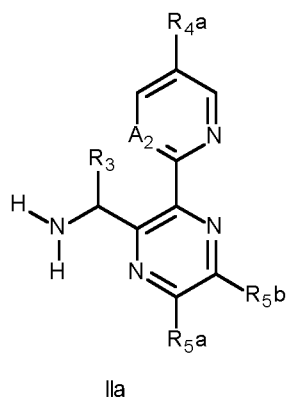
Compounds of formula XV, wherein R_3 , A_2 , R_{4a} , R_{5a} and R_{5b} are defined as in formula I, may be prepared by reduction of ketones VI (wherein R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I) for example with NaBH_4 in the usual manner (see e.g. WO2012/082997, page 141), preferably in MeOH as solvent.

5

Scheme 6:



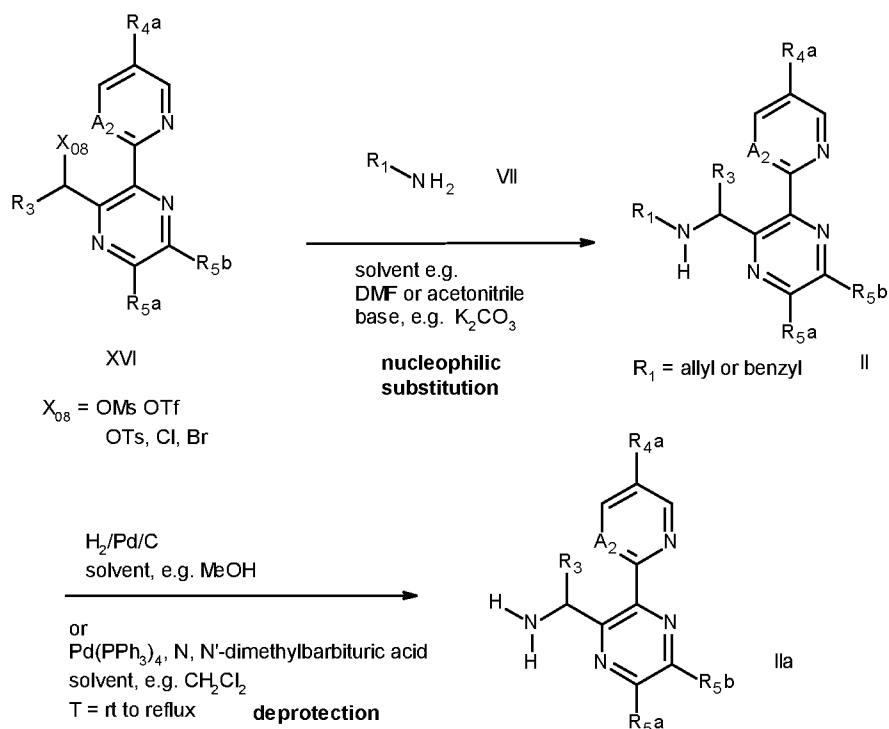
Yet another methodology to prepare compounds of general formula of IIa is outlined in Scheme 7.



10

Scheme 7:

-19-



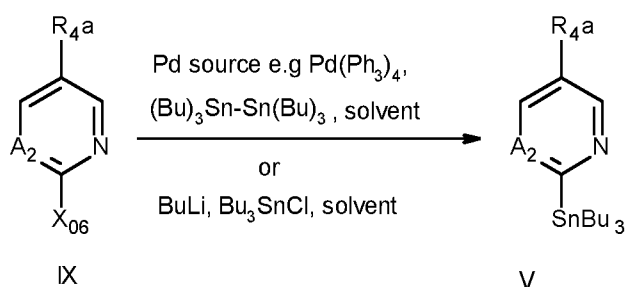
Thus, nucleophilic substitution reaction of compound of formula XVI with amines of formula VII furnishes compounds of formula II (wherein R_1 , R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are defined as in formula I) as already described in detail in Scheme 6. Compounds of formula II suited with a protecting group, e.g. R_1 is benzyl, can be hydrogenated with hydrogen in the presence of palladium (on charcoal) in a solvent, e.g. MeOH or EtOH, to give compounds of formula IIa wherein R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} is defined as in formula I (see e.g. Synlett, 2010, (18), page 2708). Compounds of formula II wherein R_1 is allyl, and R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are as defined in formula I can also be converted to compounds of formula IIa by reaction with N,N'-dimethylbarbituric acid in the presence of a Pd-catalyst, preferable tetrakis(triphenylphosphine)palladium(0), in a suitable solvent, for example CH_2Cl_2 to provide compounds of formula IIa according to J. Org. Chem. 1993, 58, 6109.

Formation of compounds of formula V, wherein A_2 and R_{4a} are as defined in formula I, is outlined in Scheme 8.

Compounds of formula V, wherein A_2 and R_{4a} are as defined in formula I, may be prepared by treatment of compounds of formula IX with a palladium source such as for example $\text{Pd}(\text{Ph}_3)_4$ and bis(tributyltin) in a suitable solvent such as DMF usually upon heating at temperatures between 100 to 130 °C. Such processes have been described, for example, in Molecular Pharmacology, 90(3), 177-187; 2016. Alternatively compounds of formula V can also be prepared by treatment of compounds of formula IX with n-Butyllithium and tributyltin chloride in a suitable solvent such as THF, usually at lower temperature such as -78°C to 0°C. Such processes have been described, for example, in US20180273562.

Scheme 8:

-20-

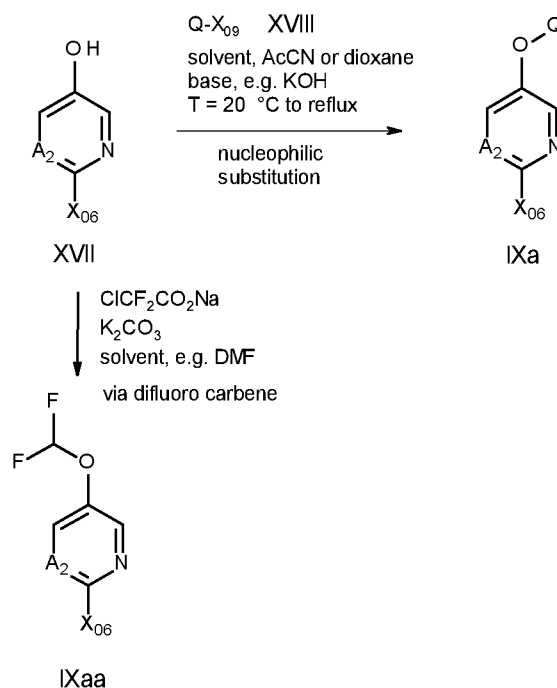


$X_{06} = \text{Cl, Br}$

Compounds of formula IXa and IXaa, wherein A_2 is as defined in formula I, X_{06} is a leaving group such as Cl, Br and Q is C_1 - C_3 haloalkyl, are either commercially available or can be prepared according to well-known methods as shown in Scheme 9.

5

Scheme 9:



Thus compounds of formula IXa, wherein A_2 is as defined in formula I, X_{06} is a leaving group such as Cl or Br, and Q is C_1 - C_3 haloalkyl, can be obtained by alkylation of compounds XVII with a compound of formula XVIII, wherein Q is C_1 - C_3 haloalkyl and X_{09} is a leaving group, such as Cl, Br, F, I, OSO_2CF_3 , or OSO_2CH_3 , in the presence of a suitable base, for example cesium or potassium carbonate, in a solvent such as acetonitrile or DMF at temperatures between 20-80 °C. Such reactions are well known to those skilled in the art and have been reported for example in see e.g. Med. Chem. Letts., 2017, 8(5), p543-548 and Bio. Med. Chem. Letts., 2017, 27(11), 2420-2423.

15

Compounds of formula IXaa, wherein A_2 is as defined in formula I, X_{06} is a leaving group such as Cl or Br, can be prepared from compounds of formula XVII by treatment with a difluorocarbene source e.g. $\text{ClCF}_2\text{CO}_2\text{Na}$ or $\text{CF}_2\text{SO}_2\text{OCHF}_2$ in the presence of a base such as KOH, potassium carbonate and the

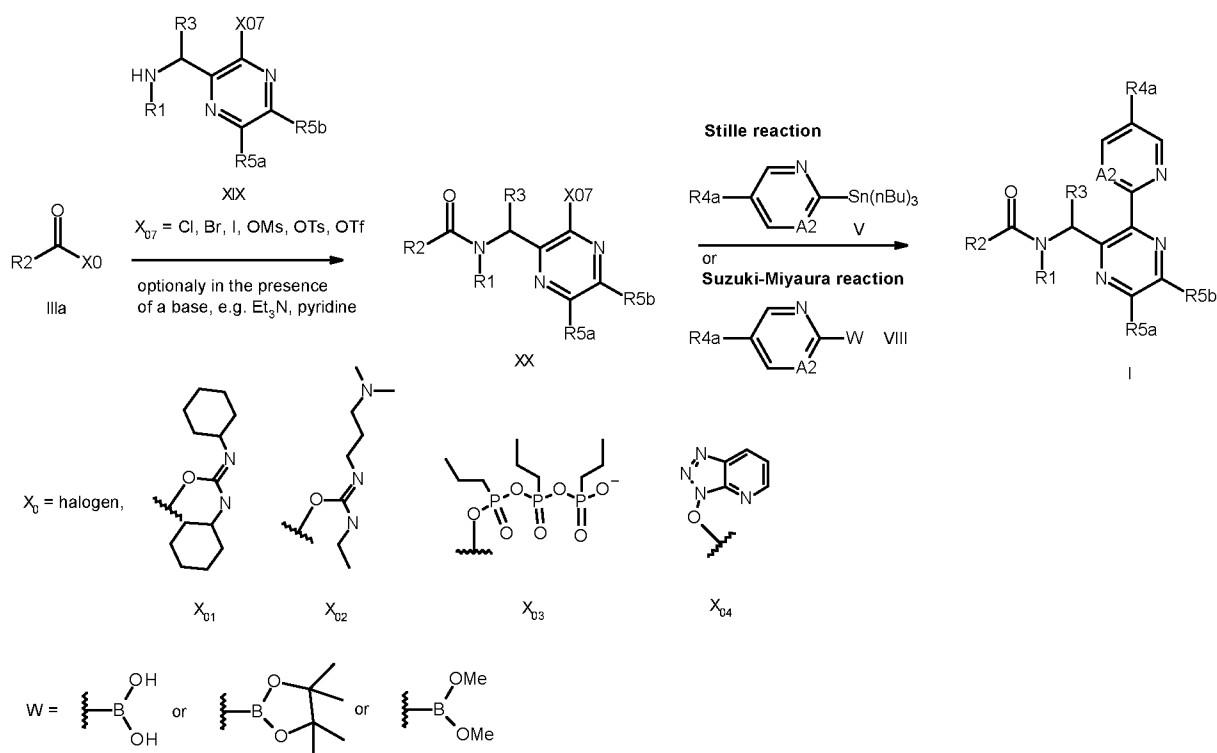
-21-

like, in an inert solvent at temperatures between 20-80 °C. Such procedures have been described for example in J. Fluor. Chem. 2017, 203, 155, and US2013/0225552, page 128, and Org. Process Res. Dev., 2011, 15, 721.

5 Compounds of formula XVII are commercially available.

Alternatively, compounds of formula I may be prepared from compounds of formula XX, wherein R₁, R₂, R₃, R_{5a}, and R_{5b} are defined in formula I and X₀₇ is defined in Scheme 5, following the synthesis outlined in Scheme 10.

10 Scheme 10:

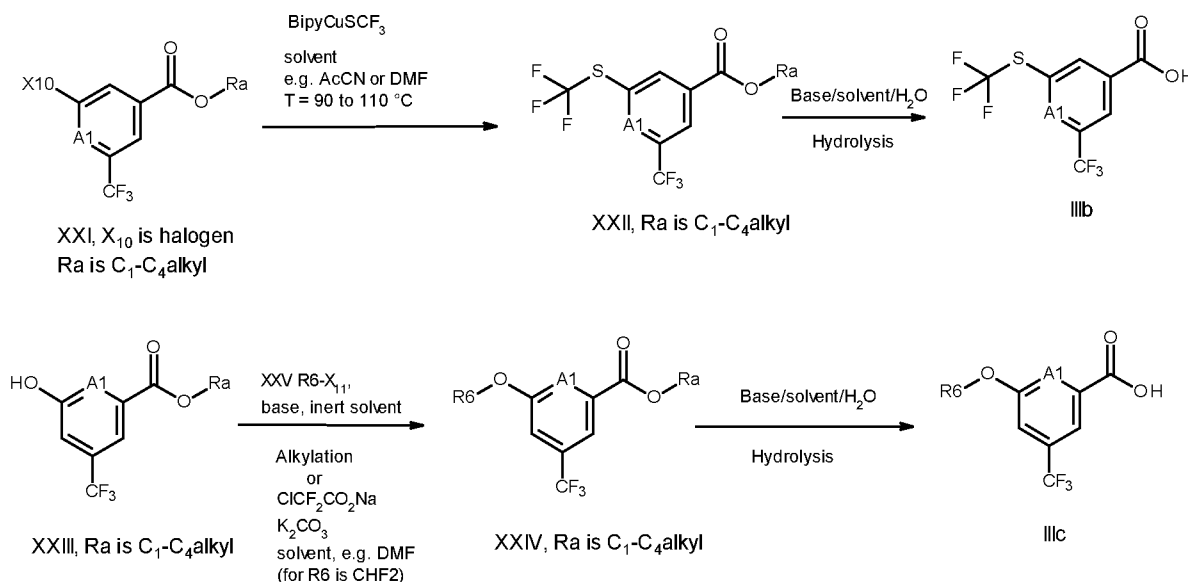


Compounds of formula I can be prepared by reaction of compounds of to compounds of formula XX with compounds of formula V (Stille reaction) or compounds of formula VIII (Suzuki Miyaura reaction) in the presence of a palladium catalyst as described in detail in Schemes 2 and 3. Compounds of
 15 formula XX can be synthesized by reaction of an activated compound of formula IIIa with an amine of formula XIX, wherein R₁, R₃, R_{5a}, and R_{5b} are defined in formula I and X₀₇ is defined in Scheme 5, following the conditions detailed in Scheme 1.

Compounds of formula III are known, for example 3-(difluoromethyl)-5-(trifluoromethyl)benzoic acid
 20 CAS: [2248290-21-1], 3-bromo-5-(trifluoromethyl)benzoic acid CAS: [328-67-6], 3-iodo-5-(trifluoromethyl)benzoic acid CAS: [28186-62-1] are commercially available, or can be prepared by those skilled in the art. For example see also WO 2013/171712, WO 2012/117000, WO2017/192385. Some

compounds of formula IIIb and IIIc are novel and have been especially prepared to enable preparation of compounds of formula I. Synthesis of such compounds is shown in scheme 11:

Scheme 11



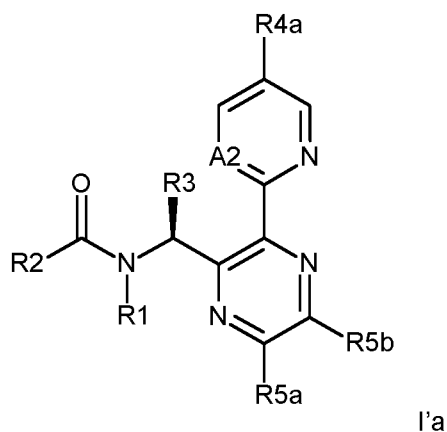
As shown in scheme 11, compounds of formula XXI, wherein X_{10} is halogen, preferably bromine or iodine, A_1 is nitrogen or methane, and Ra is C_1 - C_4 alkyl, can be treated with a bipyridine copper reagent (bpy)CuSCF₃, wherein bpy is bipyridyl, in an inert solvent such as acetonitrile or DMF, at temperatures between room temperature and 120°C, optionally under microwave heating, to give compounds of formula XXII, wherein A_1 and Ra are as defined for formula XXI. Such chemistry is known and has been described in the literature, for example, in Angew. Chem. Int. Ed. 2013, 52, 1548-1552. A preferred reagent for this transformation is (bpy)CuSCF₃ (CAS [1413732-47-4]).

Compounds of formula XXII are converted to compounds of IIIb by ester hydrolysis, for example reaction with a base e.g. Lithium, potassium, or sodium hydroxide, in water optionally in the presence of a water miscible solvent such as THF, acetone, dioxane and the like. Such reactions are well known to those skilled in the art.

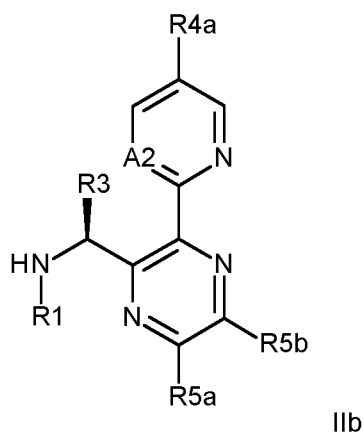
Compounds of formula XXIV, wherein R_6 is C_1 - C_3 alkyl, or C_1 - C_3 haloalkyl, and Ra is C_1 - C_4 alkyl, can be prepared by reaction of a compound of formula XXIII with a compound of formula XXV, wherein X_{11} is Cl, Br, F, I, OSO₂CF₃, or OSO₂CH₃, in the presence of a base, such as sodium hydride, K₂CO₃, or Cs₂CO₃, in an inert solvent such as THF, DMF, or acetonitrile, to give compounds of formula XXIV. Compounds of formula XXIV, wherein R_6 is CHF₂, are prepared by treatment with in situ generated difluorocarbene analogous to the procedures described for preparation of IXaa in scheme 9. Hydrolysis of compounds of formula XXIV give compounds of formula IIIc as is described vide supra.

Compounds of formula I'a

-23-



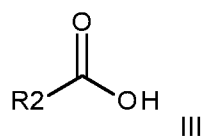
can be prepared by reaction of an amine of formula IIb



wherein R₁, R₃, A₂, R_{4a}, R_{5a}, and R_{5b} are as described in formula I, with a carboxylic acid derivative of

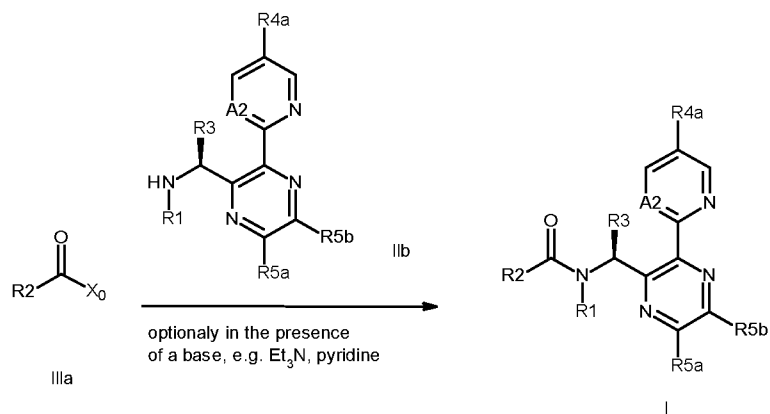
5

formula III wherein R₂ is described as above under formula I.



The chemistry is described in in more detail in Scheme 12.

Scheme 12:



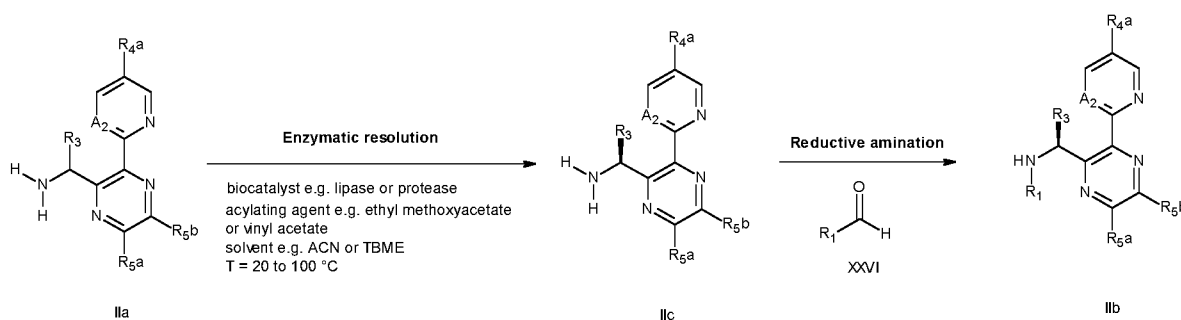
Compounds of formula IIIa, wherein R_2 is described in formula I and XO is described in Scheme 1, can be treated with compounds of formula IIb, wherein R_1 , R_3 , A_2 , R_{4a} , R_{5a} , and R_{5b} are described in formula I, under the conditions described in detail in Scheme 1. The formation of compounds of formula IIIa from compounds of formula III is described in Scheme 1.

5

The formation of compounds of formula IIb is outlined in Scheme 13. Compounds of formula IIb can be prepared by treatment of compounds of formula IIc, wherein wherein A_2 , R_3 , R_{4a} , R_{5a} , and R_{5b} are described in formula I, with compounds of formula XXVI (wherein R_1 is as defined in formula I), e.g. in the presence of $NaBH(OAc)_3$ or $NaBH_3CN$, in a suitable solvent, preferably in acetic acid at room temperature analog to WO2002/088073, page 35. Alternatively, another reagent system for the reductive amination uses a combination of $Ti(i-OiPr)_4$ and $NaBH_4$ (see Synthesis 2003 (14), 2206). Compounds of formula XXVI are generally commercially available.

Amines of formula IIc may be obtained by biocatalyzed deracemization of amines of formula IIa. This may be done for instance using a lipase, e.g. *Candida Antarctica* lipase B or *Pseudomonas fluorescens* lipase, eventually in immobilized form (e.g. Novozym® 435) in presence of an acyl donor, e.g. ethyl methoxyacetate or vinyl acetate, in a suitable solvent such as acetonitrile or methyl tert-butyl ether at temperatures between 20 °C to 100 °C. Such processes are described for instance in *J. Org. Chem.* **2007**, 72, 6918-6923 or *Adv. Synth. Catal.* **2007**, 349, 1481-1488. The expected stereochemical outcome of such enzymatic deracemization is known of those skilled in the art and is documented in the literature, for instance in *J. Org. Chem.* **1991**, 56, 2656-2665 or *J. Am. Chem. Soc.* **2015**, 137, 3996-4009.

Scheme 13:

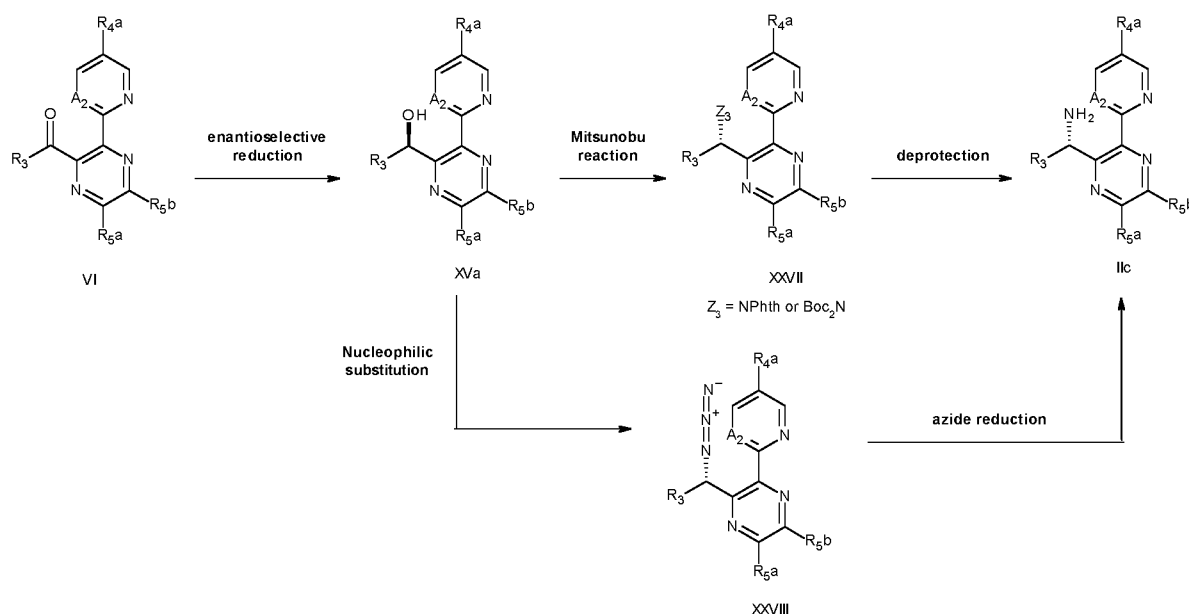


25

In an alternative process, compounds of formula IIc can be obtained from XVa, herein A_2 , R_3 , R_{4a} , R_{5a} , and R_{5b} are as described in formula I, following the synthesis described in Scheme 14.

Scheme 14:

-25-



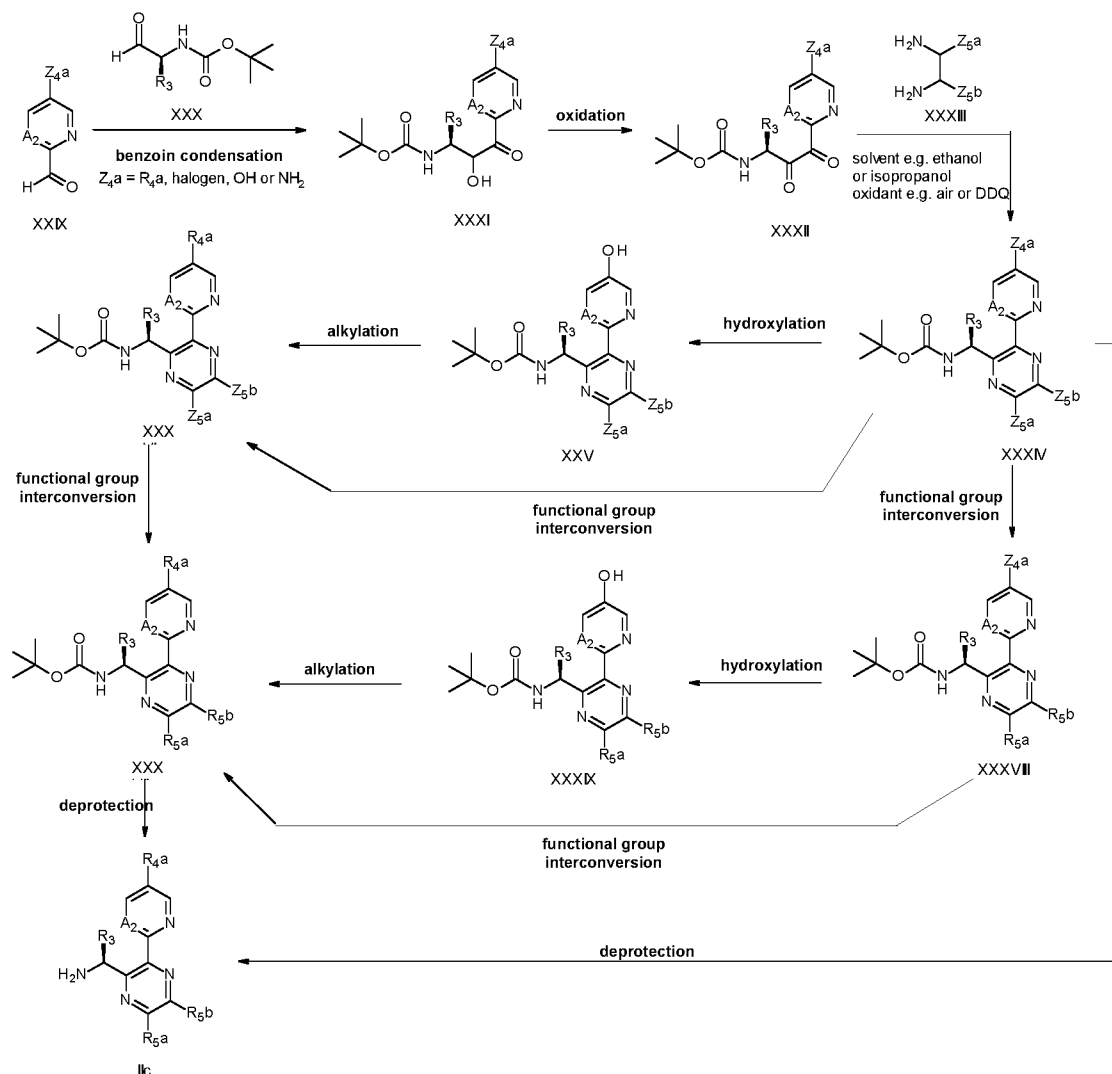
Amines of formula IIc may be obtained from intermediates of formula XXVII, wherein A₂, R₃, R_{4a}, R_{5a}, and R_{5b} are described in formula I and Z₃ is NPhth or NBoc₂. Such intermediates can be obtained from alcohols of formula XVa by a Mitsunobu reaction, which involves treating alcohols of formula XVa by diisopropyl azodicarboxylate in the presence of a phosphine such as triphenylphosphine or tributylphosphine and of an amine such as phthalimide or bis(*tert*-butoxycarbonyl)amine. Mitsunobu reactions are known by those skilled in the art to proceed with inversion of the stereocenter, as described for instance in *Chem. Rev.* **2009**, 109, 2551-2651. Amines of formula XXVII can then be transformed into amines of formula IIc by treatment with hydrazine if Z₃ = NPhth or with TFA if Z₃ = NBoc₂.

Alternatively, amines of formula IIc may be obtained by reduction of azides of formula XXVIII, wherein A₂, R₃, R_{4a}, R_{5a}, and R_{5b} are described in formula I, by treatment with triphenylphosphine and water (Staudinger reaction) or by hydrogenation for example using a palladium catalyst in the presence of hydrogen. Azides of formula XXVIII may be obtained by treatment of alcohols of formula XVa with an azidation reagent such as diphenyl phosphoryl azide in a solvent such as toluene or THF in presence of a base such as DBU. Such processes are known by those skilled in the art to proceed with inversion of the stereocenter and are described in the literature for instance in *Adv. Synth. Catal.* **2018**, 360, 2157-2165.

Alcohols of formula XVa may be obtained by enantioselective reduction of ketones of formula VI, wherein A₂, R₃, R_{4a}, R_{5a}, and R_{5b} are as described in formula I. Such reductions can be done using a catalyst, for instance a ruthenium or a rhodium catalyst with a chiral ligand such as RuCl[(*R,R*)-TsDPEN](mesitylene) or RuBF₄[(*R,R*)-TsDPEN](*p*-cymene) in the presence of a hydrogen donor system such as for example HCOOH/Et₃N or HCO₂NH₄. Such processes are described in the literature for instance in *J. Org. Chem.* **2017**, 82, 5607.

Alternatively, compounds of formula IIc may also be prepared as outlined in Scheme 15.

Scheme 15:



Amines of formula IIc can be prepared by deprotection of amines of formula XXXVII, wherein A_2 , R_3 , R_{4a} , R_{5a} and R_{5b} are described in formula I, for instance using an acid such as trifluoroacetic acid or hydrochloric acid. Compounds of formula XXXVII may be synthesized from compounds of formula XXXVI, wherein A_2 , R_3 and R_{4a} are described in formula I and Z_{5a} and Z_{5b} are, independently of each other, selected from R_{5a} , R_{5b} , halogen, NH_2 or OH. Such functional group interconversions are known to those skilled in the art and examples of such transformations have been described in the literature, for instance in *Eur. J. Org. Chem.* **2005**, 19, 4141-4153 or in *J. Org. Chem.* **2008**, 73, 7481-7485.

Compounds of formula XXXVI can be obtained from compounds of formula XXXV by alkylation, for instance using a base and an electrophile, e.g. chlorodifluoroacetic acid. Compounds of formula XXXV may be prepared by hydroxylation of compounds of formula XXXIV, wherein A_2 and R_3 are described in formula I, Z_{4a} is selected from R_{4a} , halogen or NH_2 , and Z_{5a} and Z_{5b} are, independently of each other, selected from R_{5a} , R_{5b} , halogen, NH_2 or OH. Conversion of XXXIV to XXXV can be done following methods referenced in the literature, for instance in *Org. Lett.* **2016**, 18, 2244-2247 or *Tetrahedron* **2009**, 65, 757-764. Amines of formula XXXIV can be obtained by condensation of diamines of formula XXXIII, wherein Z_{5a} and Z_{5b} are, independently of each other, selected from R_{5a} , R_{5b} , halogen, NH_2 or OH, on

diketones of formula XXXII, wherein A_2 and R_3 are described in formula I and Z_{4a} is selected from R_{4a} , halogen or NH_2 . This condensation can take place in the presence of a suitable solvent such as ethanol or isopropanol in presence of an oxidant such as air or DDQ. Diketones of formula XXXII may be formed by oxidation of hydroxyketones of formula XXXI, A_2 and R_3 are described in formula I and Z_{4a} is selected from R_{4a} , halogen or NH_2 . This oxidation can involve for instance SO_3 -pyridine in presence of DMSO and a base, for instance triethylamine or also sodium hypochlorite in presence of a catalyst such as TEMPO/ Bu_4NHSO_4 . Examples of such oxidations can be found in the literature, for instance in *Synlett*, **2014**, 25, 596 or *J. Am. Chem. Soc.* **1990**, 112, 5290-5313. Hydroxyketones of formula XXXI may be synthesized by cross-benzoin condensation between aldehydes of formula XXIX, wherein A_2 is described in formula I and Z_{4a} is selected from R_{4a} , halogen or NH_2 , and aldehydes of formula XXX, wherein R_3 is as described in formula I. Aldehydes of formula XXX are commercially available in chiral form, like for instance Boc-L-alaninal (CAS 79069-50-4) or tert-butyl N-[(1S)-1-(cyclopropylmethyl)-2-oxo-ethyl]carbamate (CAS 881902-36-9). Cross-benzoin condensations are done in the usual way by employing an organocatalyst such as a triazolium salt or a thiazolium salt in the presence of a base such as potassium tert-butoxide or isopropyl-diethylamine in a suitable solvent such as dichloromethane or tetrahydrofuran at a temperature between -20 °C and the boiling point of the solvent. Examples of catalysts for such transformations have been described in the literature for instance in *J. Am. Chem. Soc.* **2014**, 136, 7539-7542 or in *Org. Lett.* **2016**, 18, 4518-4521.

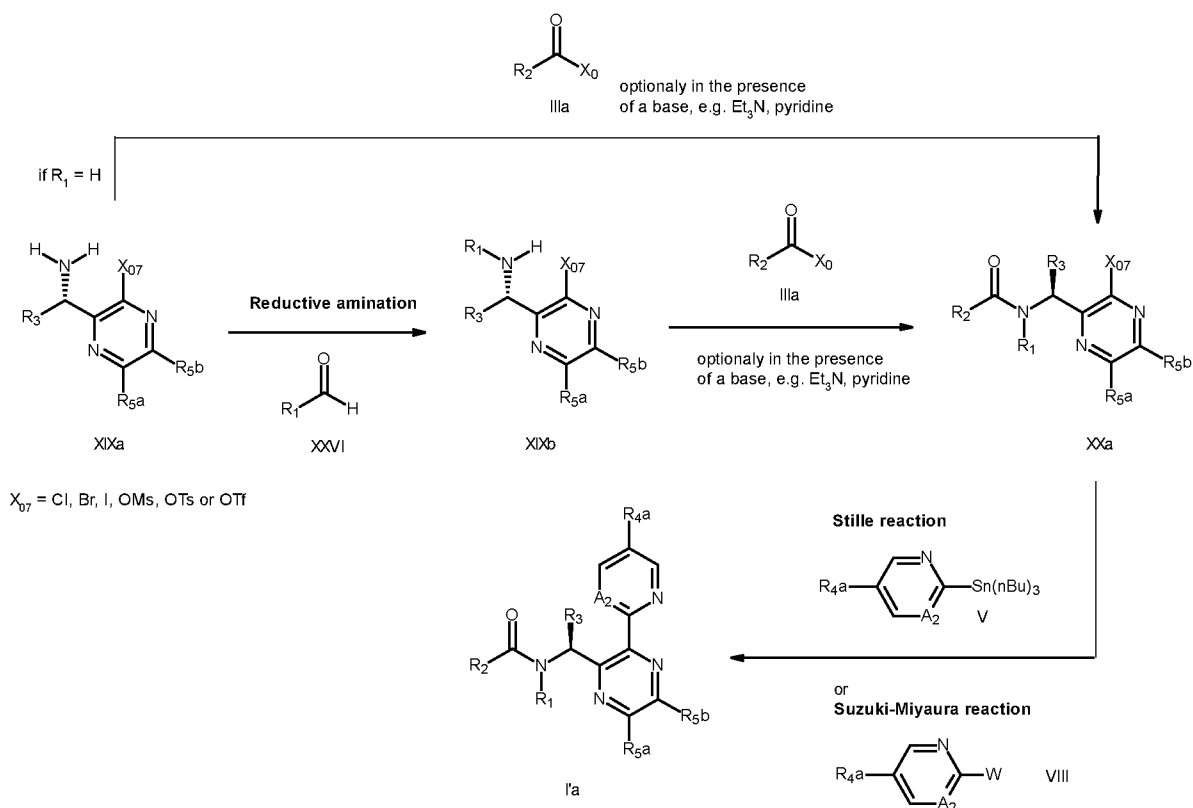
Compounds of formula XXXVI may also be obtained directly from compounds of formula XXXIV, for instance using transition metal catalysis or diazonium chemistry. Such functional group interconversions are known to those skilled in the art and examples can be found in the literature, for instance in *J. Am. Chem. Soc.* **2019**, 141, 19257-19262, *Angew. Chem. Int. Ed.* **2015**, 54, 5736-5739 or *Heterocycles*, 2004, 63, 2735-2746.

Compounds of formula XXXVII may be obtained from compounds of formula XXXIX, wherein A_2 , R_3 , R_{5a} and R_{5b} are described in formula I, by alkylation for instance using a base and an electrophile, e.g. chlorodifluoroacetic acid. Compounds of formula XXXIX may be synthesized from compounds of formula XXXVIII (wherein A_2 , R_3 , R_{5a} and R_{5b} are described in formula I and Z_{4a} is selected from R_{4a} , halogen or NH_2) by hydroxylation reaction, following methods referenced in the literature, for instance in *Org. Lett.* **2016**, 18, 2244-2247 or *Tetrahedron* **2009**, 65, 757-764. Alternatively, compounds of formula XXXVII may be obtained from compounds of formula XXXVIII. Such functional group interconversions are known to those skilled in the art and examples can be found in the literature, for instance in *J. Am. Chem. Soc.* **2019**, 141, 19257-19262, *Angew. Chem. Int. Ed.* **2015**, 54, 5736-5739 or *Heterocycles*, 2004, 63, 2735-2746.

As shown in Scheme 16, compounds of formula I'a can be alternatively prepared by reaction of compounds of formula XXa (wherein R_1 , R_3 , R_{5a} , and R_{5b} are defined in formula I, and X_{07} is a leaving group like, for example, chlorine, bromine, iodine) with compounds of formula V (Stille reaction) or

compounds of formula VIII (Suzuki-Miyaura reaction) in the presence of a palladium catalyst as described in detail in Schemes 2 and 3.

Scheme 16:



5

Compounds of formula I'a can be prepared by reaction of compounds of formula XXa with compounds of formula V (Stille reaction) or compounds of formula VIII (Suzuki-Miyaura reaction) in the presence of a palladium catalyst as described in detail in Schemes 2 and 3.

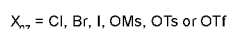
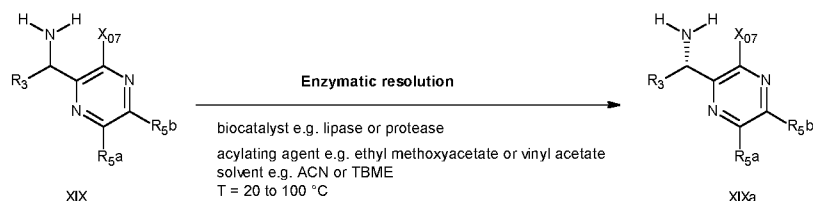
- 10 Compounds of formula XXa can be prepared by coupling of amines of formula XIXb (wherein R_1 , R_3 , R_{5a} , and R_{5b} are defined in formula I, and X_{07} is a leaving group like, for example, chlorine, bromine, iodine) and compounds of formula IIIa, wherein R_2 is described in formula I and X_0 is described in Scheme 1, under the conditions described in detail in Scheme 1. Under the same conditions, if $R_1 = H$, compounds of formula XXa may be obtained directly from compounds of formula XIXa (wherein R_3 , R_{5a} , and R_{5b} are defined in formula I, and X_{07} is a leaving group like, for example, chlorine, bromine, iodine).
- 15

Compounds of formula XIXb can be prepared by treatment of compounds of formula XIXa, with compounds of formula XXVI (wherein R_1 is defined in formula I), e.g. in the presence of $NaBH(OAc)_3$ or $NaBH_3CN$, in a suitable solvent, preferably in acetic acid at room temperature analog to

20 WO2002/088073, page 35. Alternatively, another reagent system for the reductive amination uses a combination of $Ti(OiPr)_4$ and $NaBH_4$ (see *Synthesis* 2003 (14), 2206).

Amines of formula XIXa can be prepared by deracemization procedure method, which involves for example, a selective acylation of one enantiomer. Such an example is described more in details in Scheme 17.

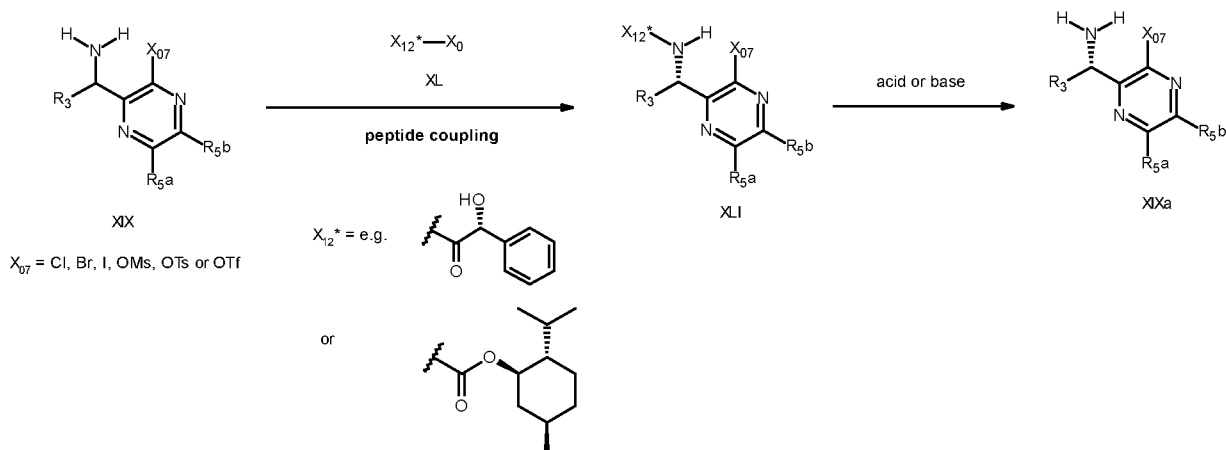
5 Scheme 17:



Amines of formula XIXa may be obtained by biocatalyzed deracemization of amines of formula XIX, wherein R_3 , R_{5a} , and R_{5b} are described in formula I and X_{07} is a leaving group such as bromine, chlorine, iodine, mesylate, tosylate or triflate. This may be done for instance using a lipase, e.g. *Candida* *Antarctica* lipase B or *Pseudomonas fluorescens* lipase, eventually in immobilized form (e.g. Novozym® 435) in presence of an acyl donor, e.g. ethyl methoxyacetate or vinyl acetate, in a suitable solvent such as acetonitrile or methyl tert-butyl ether at temperatures between 20 °C to 100 °C. Such processes are described for instance in *J. Org. Chem.* **2007**, 72, 6918-6923 or *Adv. Synth. Catal.* **2007**, 349, 1481-1488. The expected stereochemical outcome of such enzymatic deracemization are known of those skilled in the art and are documented in the literature, for instance in *J. Org. Chem.* **1991**, 56, 2656-2665 or *J. Am. Chem. Soc.* **2015**, 137, 3996-4009.

Alternatively, resolution of amines of formula XIXa may be achieved using a chiral auxiliary, as described in Scheme 18.

Scheme 18



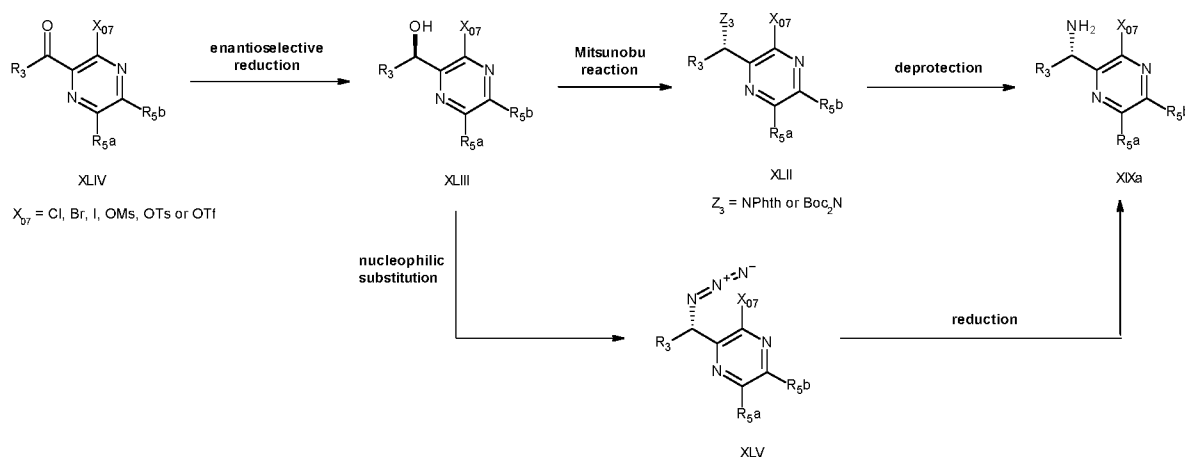
Amines of formula XIXa can be prepared for intermediates of formula XLI, wherein R_3 , R_{5a} , and R_{5b} are described in formula I, X_{07} is a leaving group such as bromine, chlorine or iodine and X_{12}^* is a chiral

-30-

auxiliary, by treatment with acids such as HCl or bases such as NaOH. Amines of formula XLI can be formed by coupling of a chiral compound of formula XL, wherein X_0 is described in Scheme 1 and X_{12}^* is a chiral moiety of known chirality, with amines of formula XIX following the conditions detailed in Scheme 1. Chiral auxiliaries of formula XL are for instance derived from mandelic acid or (1R)-menthylchloroformate. Examples of such deracemization processes are reported in the literature, for instance in *J. Org. Chem.* **2007**, *72*, 485-493.

Alternatively, amines of formula XLVIIIId (wherein R_3 , R_{5a} , and R_{5b} are defined in formula I, and X_{07} is a leaving group like, for example, chlorine, bromine, iodine), can be formed as described in Scheme 19.

Scheme 19:



Amines of formula XIXa may be obtained from intermediates of formula XLII, wherein R_3 , R_{5a} , and R_{5b} are as described in formula I, X_{07} is a leaving group as described in Scheme 5 and Z_3 is NPhth or NBoc₂.

Such intermediates can be obtained from alcohols of formula XLIII, wherein R_3 , R_{5a} , and R_{5b} are described in formula I and X_{07} is a leaving group as described in Scheme 5, by a Mitsunobu reaction, which involves treating alcohols of formula XLIII by diisopropyl azodicarboxylate in the presence of a phosphine such as triphenylphosphine or tributylphosphine and of an amine such as phthalimide or bis(*tert*-butoxycarbonyl)amine. Mitsunobu reactions are known by those skilled in the art to proceed with inversion of the stereocenter, as described for instance in *Chem. Rev.* **2009**, *109*, 2551-2651. Amines of formula XLII can then be transformed into amines of formula XIXa by treatment with hydrazine if $Z_3 = \text{NPhth}$ or with TFA if $Z_3 = \text{NBoc}_2$.

Alternatively, amines of formula XIXa may be obtained by reduction of azides of formula XLV, wherein R_3 , R_{5a} , and R_{5b} are as described in formula I and X_{07} is a leaving group as described in Scheme 5, by treatment with triphenylphosphine and water (Staudinger reaction) or by hydrogenation for example using a palladium catalyst in the presence of hydrogen. Azides of formula XLV may be obtained by treatment of alcohols of formula XLIII with an azidation reagent such as diphenyl phosphoryl azide in a solvent such as toluene or THF in presence of a base such as DBU. Such processes are known by

those skilled in the art to proceed with inversion of the stereocenter and are described in the literature for instance in *Adv. Synth. Catal.* **2018**, 360, 2157–2165.

Alcohols of formula XLIII may be obtained by enantioselective reduction of ketones of formula XLIV, wherein R_3 , R_{5a} , and R_{5b} are as described in formula I and X_{07} is a leaving group as described in Scheme 5. Such reductions can be done using catalysts, for instance a ruthenium or a rhodium catalyst with a chiral ligand such as $RuCl[(R,R)\text{-TsDPEN}](\text{mesitylene})$ or $RuBF_4[(R,R)\text{-TsDPEN}](p\text{-cymene})$ in the presence of a hydrogen donor system such as for example $HCOOH/Et_3N$ or HCO_2NH_4 . Such processes are described in the literature for instance in *J. Org. Chem.* **2017**, 82, 5607.

Depending on the procedure or the reaction conditions, the reactants can be reacted in the presence of a base. Examples of suitable bases are alkali metal or alkaline earth metal hydroxides, alkali metal or alkaline earth metal hydrides, alkali metal or alkaline earth metal amides, alkali metal or alkaline earth metal alkoxides, alkali metal or alkaline earth metal acetates, alkali metal or alkaline earth metal carbonates, alkali metal or alkaline earth metal dialkylamides or alkali metal or alkaline earth metal alkylsilylamides, alkylamines, alkylenediamines, free or N-alkylated saturated or unsaturated cycloalkylamines, basic heterocycles, ammonium hydroxides and carbocyclic amines. Examples which may be mentioned are sodium hydroxide, sodium hydride, sodium amide, sodium methoxide, sodium acetate, sodium carbonate, potassium tert-butoxide, potassium hydroxide, potassium carbonate, potassium hydride, lithium diisopropylamide, potassium bis(trimethylsilyl)amide, calcium hydride, triethylamine, diisopropylethylamine, triethylenediamine, cyclohexylamine, N-cyclohexyl-N,N-dimethylamine, N,N-diethylaniline, pyridine, 4-(N,N-dimethylamino)pyridine, quinuclidine, N-methylmorpholine, benzyltrimethylammonium hydroxide and 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU).

The reactants can be reacted with each other as such, i.e. without adding a solvent or diluent. In most cases, however, it is advantageous to add an inert solvent or diluent or a mixture of these. If the reaction is carried out in the presence of a base, bases which are employed in excess, such as triethylamine, pyridine, N-methylmorpholine or N,N-diethylaniline, may also act as solvents or diluents.

The reactions are advantageously carried out in a temperature range from approximately -80°C to approximately $+140^\circ\text{C}$, preferably from approximately -30°C to approximately $+100^\circ\text{C}$, in many cases in the range between ambient temperature and approximately $+80^\circ\text{C}$.

Depending on the choice of the reaction conditions and starting materials which are suitable in each case, it is possible, for example, in one reaction step only to replace one substituent by another substituent according to the invention, or a plurality of substituents can be replaced by other substituents according to the invention in the same reaction step.

Salts of compounds of formula I can be prepared in a manner known *per se*. Thus, for example, acid addition salts of compounds of formula I are obtained by treatment with a suitable acid or a suitable ion exchanger reagent and salts with bases are obtained by treatment with a suitable base or with a suitable ion exchanger reagent.

5

Salts of compounds of formula I can be converted in the customary manner into the free compounds I, acid addition salts, for example, by treatment with a suitable basic compound or with a suitable ion exchanger reagent and salts with bases, for example, by treatment with a suitable acid or with a suitable ion exchanger reagent.

10

Salts of compounds of formula I can be converted in a manner known *per se* into other salts of compounds of formula I, acid addition salts, for example, into other acid addition salts, for example by treatment of a salt of inorganic acid such as hydrochloride with a suitable metal salt such as a sodium, barium or silver salt, of an acid, for example with silver acetate, in a suitable solvent in which an inorganic salt which forms, for example silver chloride, is insoluble and thus precipitates from the reaction mixture.

15

Depending on the procedure or the reaction conditions, the compounds of formula I, which have salt-forming properties can be obtained in free form or in the form of salts.

20

The compounds of formula I and, where appropriate, the tautomers thereof, in each case in free form or in salt form, can be present in the form of one of the isomers which are possible or as a mixture of these, for example in the form of pure isomers, such as antipodes and/or diastereomers, or as isomer mixtures, such as enantiomer mixtures, for example racemates, diastereomer mixtures or racemate mixtures, depending on the number, absolute and relative configuration of asymmetric carbon atoms which occur in the molecule and/or depending on the configuration of non-aromatic double bonds which occur in the molecule; the invention relates to the pure isomers and also to all isomer mixtures which are possible and is to be understood in each case in this sense hereinabove and hereinbelow, even when stereochemical details are not mentioned specifically in each case.

25

30

Diastereomer mixtures or racemate mixtures of compounds of formula I, in free form or in salt form, which can be obtained depending on which starting materials and procedures have been chosen can be separated in a known manner into the pure diastereomers or racemates on the basis of the physicochemical differences of the components, for example by fractional crystallization, distillation and/or chromatography.

35

Enantiomer mixtures, such as racemates, which can be obtained in a similar manner can be resolved into the optical antipodes by known methods, for example by recrystallization from an optically active solvent, by chromatography on chiral adsorbents, for example high-performance liquid chromatography (HPLC) on acetyl cellulose, with the aid of suitable microorganisms, by cleavage with specific,

-33-

immobilized enzymes, via the formation of inclusion compounds, for example using chiral crown ethers, where only one enantiomer is complexed, or by conversion into diastereomeric salts, for example by reacting a basic end-product racemate with an optically active acid, such as a carboxylic acid, for example camphor, tartaric or malic acid, or sulfonic acid, for example camphorsulfonic acid, and
5 separating the diastereomer mixture which can be obtained in this manner, for example by fractional crystallization based on their differing solubilities, to give the diastereomers, from which the desired enantiomer can be set free by the action of suitable agents, for example basic agents.

Pure diastereomers or enantiomers can be obtained according to the invention not only by separating
10 suitable isomer mixtures, but also by generally known methods of diastereoselective or enantioselective synthesis, for example by carrying out the process according to the invention with starting materials of a suitable stereochemistry.

N-oxides can be prepared by reacting a compound of the formula I with a suitable oxidizing agent, for
15 example the H_2O_2 /urea adduct in the presence of an acid anhydride, e.g. trifluoroacetic anhydride. Such oxidations are known from the literature, for example from *J. Med. Chem.*, 32 (12), 2561-73, 1989 or WO 2000/15615.

It is advantageous to isolate or synthesize in each case the biologically more effective isomer, for
20 example enantiomer or diastereomer, or isomer mixture, for example enantiomer mixture or diastereomer mixture, if the individual components have a different biological activity.

The compounds of formula I and, where appropriate, the tautomers thereof, in each case in free form or in salt form, can, if appropriate, also be obtained in the form of hydrates and/or include other solvents,
25 for example those which may have been used for the crystallization of compounds which are present in solid form.

The compounds of formula I according to the following Tables A-1 to A-27 can be prepared according to the methods described above. The examples which follow are intended to illustrate the invention
30 and show preferred compounds of formula I, in the form of a compound of formula J-1.

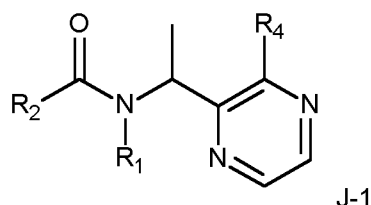
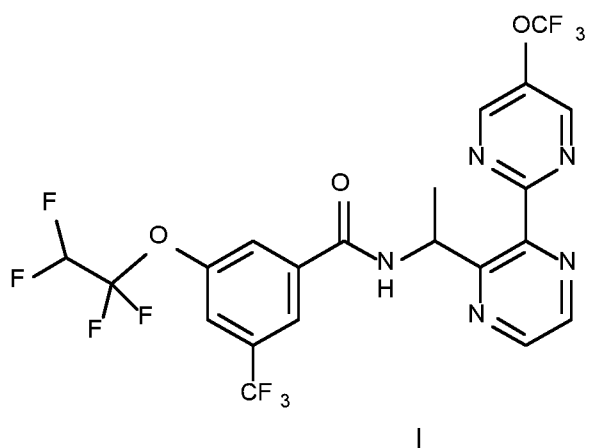


Table A-1 provides 13 compounds A-1.001 to A-1.013 of formula J-1 wherein R_1 is H, R_4 is [5-(trifluoromethoxy)pyrimidin-2-yl] and R_2 is as defined in table Z. For example, A-1.002 is

-34-

Table Z: Substituent definitions of R₂:

Index	R ₂	Index	R ₂
1		7	
2		8	
3		9	

Index	R ₂	Index	R ₂
4		10	
5		11	
6		12	
13			

Table A-2 provides 13 compounds A-2.001 to A-2.013 of formula J-1 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-3 provides 13 compounds A-3.001 to A-3.013 of formula J-1 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table A-4 provides 13 compounds A-4.001 to A-4.013 of formula J-1 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-5 provides 13 compounds A-5.001 to A-5.013 of formula J-1 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table A-6 provides 13 compounds A-6.001 to A-6.013 of formula J-1 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-7 provides 13 compounds A-7.001 to A-7.013 of formula J-1 wherein R₁ is H, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

- 5 Table A-8 provides 13 compounds A-8.001 to A-8.013 of formula J-1 wherein R₁ is H, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-9 provides 13 compounds A-9.001 to A-9.013 of formula J-1 wherein R₁ is H, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

- 10 Table A-10 provides 13 compounds A-10.001 to A-10.013 of formula J-1 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table A-11 provides 13 compounds A-11.001 to A-11.013 of formula J-1 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-12 provides 13 compounds A-12.001 to A-12.013 of formula J-1 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

- 15 Table A-13 provides 13 compounds A-13.001 to A-13.013 of formula J-1 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-14 provides 13 compounds A-14.001 to A-14.013 of formula J-1 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

- 20 Table A-15 provides 13 compounds A-15.001 to A-15.013 of formula J-1 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-16 provides 13 compounds A-16.001 to A-16.013 of formula J-1 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table A-17 provides 13 compounds A-17.001 to A-17.013 of formula J-1 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

- 25 Table A-18 provides 13 compounds A-18.001 to A-18.013 of formula J-1 wherein R₁ is CH₃, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

Table A-19 provides 13 compounds A-19.001 to A-19.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

- 30 Table A-20 provides 13 compounds A-20.001 to A-20.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-21 provides 13 compounds A-21.001 to A-21.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table A-22 provides 13 compounds A-22.001 to A-22.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

- 35 Table A-23 provides 13 compounds A-23.001 to A-23.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table A-24 provides 13 compounds A-24.001 to A-24.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table A-25 provides 13 compounds A-25.001 to A-25.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table A-26 provides 13 compounds A-26.001 to A-26.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

- 5 Table A-27 provides 13 compounds A-27.001 to A-27.013 of formula J-1 wherein R₁ is CH₂Cyp, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

The compounds of formula I according to the following Tables B-1 to B-27 can be prepared according to the methods described above. The examples which follow are intended to illustrate the invention and show preferred compounds of formula I, in the form of a compound of formula J-2.

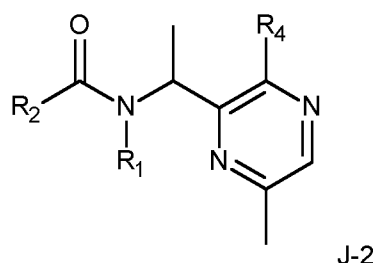


Table B-1 provides 13 compounds B-1.001 to B-1.013 of formula J-2 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-2 provides 13 compounds B-2.001 to B-2.013 of formula J-2 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-3 provides 13 compounds B-3.001 to B-3.013 of formula J-2 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-4 provides 13 compounds B-4.001 to B-4.013 of formula J-2 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-5 provides 13 compounds B-5.001 to B-5.013 of formula J-2 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-6 provides 13 compounds B-6.001 to B-6.013 of formula J-2 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-7 provides 13 compounds B-7.001 to B-7.013 of formula J-2 wherein R₁ is H, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-8 provides 13 compounds B-8.001 to B-8.013 of formula J-2 wherein R₁ is H, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-9 provides 13 compounds B-9.001 to B-9.013 of formula J-2 wherein R₁ is H, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

Table B-10 provides 13 compounds B-10.001 to B-10.013 of formula J-2 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-11 provides 13 compounds B-11.001 to B-11.013 of formula J-2 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-12 provides 13 compounds B-12.001 to B-12.013 of formula J-2 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-13 provides 13 compounds B-13.001 to B-13.013 of formula J-2 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

5 Table B-14 provides 13 compounds B-14.001 to B-14.013 of formula J-2 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-15 provides 13 compounds B-15.001 to B-15.013 of formula J-2 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

10 Table B-16 provides 13 compounds B-16.001 to B-16.013 of formula J-2 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-17 provides 13 compounds B-17.001 to B-17.013 of formula J-2 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-18 provides 13 compounds B-18.001 to B-18.013 of formula J-2 wherein R₁ is CH₃, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

15 Table B-19 provides 13 compounds B-19.001 to B-19.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-20 provides 13 compounds B-20.001 to B-20.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

20 Table B-21 provides 13 compounds B-21.001 to B-21.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table B-22 provides 13 compounds B-22.001 to B-22.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-23 provides 13 compounds B-23.001 to B-23.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

25 Table B-24 provides 13 compounds B-24.001 to B-24.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-25 provides 13 compounds B-25.001 to B-25.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

30 Table B-26 provides 13 compounds B-26.001 to B-26.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table B-27 provides 13 compounds B-27.001 to B-27.013 of formula J-2 wherein R₁ is CH₂Cyp, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

35 The compounds of formula I according to the following Tables C-1 to C-27 can be prepared according to the methods described above. The examples which follow are intended to illustrate the invention and show preferred compounds of formula I, in the form of a compound of formula J-3.

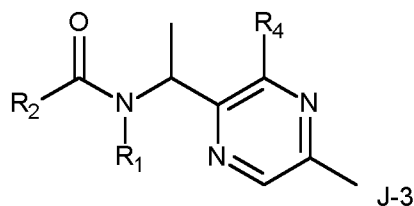


Table C-1 provides 13 compounds C-1.001 to C-1.013 of formula J-3 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-2 provides 13 compounds C-2.001 to C-2.013 of formula J-3 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-3 provides 13 compounds C-3.001 to C-3.013 of formula J-3 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-4 provides 13 compounds C-4.001 to C-4.013 of formula J-3 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-5 provides 13 compounds C-5.001 to C-5.013 of formula J-3 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-6 provides 13 compounds C-6.001 to C-6.013 of formula J-3 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-7 provides 13 compounds C-7.001 to C-7.013 of formula J-3 wherein R₁ is H, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-8 provides 13 compounds C-8.001 to C-8.013 of formula J-3 wherein R₁ is H, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-9 provides 13 compounds C-9.001 to C-9.013 of formula J-3 wherein R₁ is H, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

Table C-10 provides 13 compounds C-10.001 to C-10.013 of formula J-3 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-11 provides 13 compounds C-11.001 to C-11.013 of formula J-3 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-12 provides 13 compounds C-12.001 to C-12.013 of formula J-3 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-13 provides 13 compounds C-13.001 to C-13.013 of formula J-3 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-14 provides 13 compounds C-14.001 to C-14.013 of formula J-3 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-15 provides 13 compounds C-15.001 to C-15.013 of formula J-3 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-16 provides 13 compounds C-16.001 to C-16.013 of formula J-3 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-17 provides 13 compounds C-17.001 to C-17.013 of formula J-3 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-18 provides 13 compounds C-18.001 to C-18.013 of formula J-3 wherein R₁ is CH₃, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

5 Table C-19 provides 13 compounds C-19.001 to C-19.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-20 provides 13 compounds C-20.001 to C-20.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

10 Table C-21 provides 13 compounds C-21.001 to C-21.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table C-22 provides 13 compounds C-22.001 to C-22.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-23 provides 13 compounds C-23.001 to C-23.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

15 Table C-24 provides 13 compounds C-24.001 to C-24.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-25 provides 13 compounds C-25.001 to C-25.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

20 Table C-26 provides 13 compounds C-26.001 to C-26.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table C-27 provides 13 compounds C-27.001 to C-27.013 of formula J-3 wherein R₁ is CH₂Cyp, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

25 The compounds of formula I according to the following Tables D-1 to D-27 can be prepared according to the methods described above. The examples which follow are intended to illustrate the invention and show preferred compounds of formula I, in the form of a compound of formula J-4.

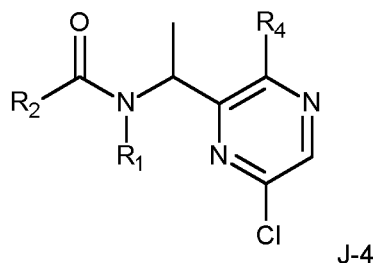


Table D-1 provides 13 compounds D-1.001 to D-1.013 of formula J-4 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

30 Table D-2 provides 13 compounds D-2.001 to D-2.013 of formula J-4 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table D-3 provides 13 compounds D-3.001 to D-3.013 of formula J-4 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table D-4 provides 13 compounds D-4.001 to D-4.013 of formula J-4 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table D-5 provides 13 compounds D-5.001 to D-5.013 of formula J-4 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

5 Table D-6 provides 13 compounds D-6.001 to D-6.013 of formula J-4 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table D-7 provides 13 compounds D-7.001 to D-7.013 of formula J-4 wherein R₁ is H, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

10 Table D-8 provides 13 compounds D-8.001 to D-8.013 of formula J-4 wherein R₁ is H, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table D-9 provides 13 compounds D-9.001 to D-9.013 of formula J-4 wherein R₁ is H, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

Table D-10 provides 13 compounds D-10.001 to D-10.013 of formula J-4 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

15 Table D-11 provides 13 compounds D-11.001 to D-11.013 of formula J-4 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table D-12 provides 13 compounds D-12.001 to D-12.013 of formula J-4 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

20 Table D-13 provides 13 compounds D-13.001 to D-13.013 of formula J-4 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table D-14 provides 13 compounds D-14.001 to D-14.013 of formula J-4 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table D-15 provides 13 compounds D-15.001 to D-15.013 of formula J-4 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

25 Table D-16 provides 13 compounds D-16.001 to D-16.013 of formula J-4 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table D-17 provides 13 compounds D-17.001 to D-17.013 of formula J-4 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

30 Table D-18 provides 13 compounds D-18.001 to D-18.013 of formula J-4 wherein R₁ is CH₃, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

Table D-19 provides 13 compounds D-19.001 to D-19.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table D-20 provides 13 compounds D-20.001 to D-20.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

35 Table D-21 provides 13 compounds D-21.001 to D-21.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table D-22 provides 13 compounds D-22.001 to D-22.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

-42-

Table D-23 provides 13 compounds D-23.001 to D-23.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

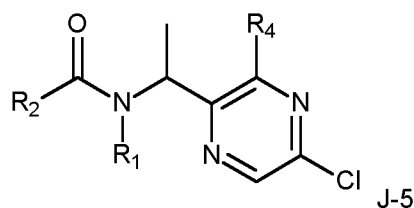
Table D-24 provides 13 compounds D-24.001 to D-24.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

5 Table D-25 provides 13 compounds D-25.001 to D-25.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table D-26 provides 13 compounds D-26.001 to D-26.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

10 Table D-27 provides 13 compounds D-27.001 to D-27.013 of formula J-4 wherein R₁ is CH₂Cyp, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

The compounds of formula I according to the following Tables E-1 to E-27 can be prepared according to the methods described above. The examples which follow are intended to illustrate the invention and show preferred compounds of formula I, in the form of a compound of formula J-5.



15

Table E-1 provides 13 compounds E-1.001 to E-1.013 of formula J-5 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table E-2 provides 13 compounds E-2.001 to E-2.013 of formula J-5 wherein R₁ is H, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

20 Table E-3 provides 13 compounds E-3.001 to E-3.013 of formula J-5 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table E-4 provides 13 compounds E-4.001 to E-4.013 of formula J-5 wherein R₁ is H, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

25 Table E-5 provides 13 compounds E-5.001 to E-5.013 of formula J-5 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table E-6 provides 13 compounds E-6.001 to E-6.013 of formula J-5 wherein R₁ is H, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table E-7 provides 13 compounds E-7.001 to E-7.013 of formula J-5 wherein R₁ is H, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

30 Table E-8 provides 13 compounds E-8.001 to E-8.013 of formula J-5 wherein R₁ is H, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table E-9 provides 13 compounds E-9.001 to E-9.013 of formula J-5 wherein R₁ is H, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

Table E-10 provides 13 compounds E-10.001 to E-10.013 of formula J-5 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table E-11 provides 13 compounds E-11.001 to E-11.013 of formula J-5 wherein R₁ is CH₃, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

5 Table E-12 provides 13 compounds E-12.001 to E-12.013 of formula J-5 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table E-13 provides 13 compounds E-13.001 to E-13.013 of formula J-5 wherein R₁ is CH₃, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

10 Table E-14 provides 13 compounds E-14.001 to E-14.013 of formula J-5 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table E-15 provides 13 compounds E-15.001 to E-15.013 of formula J-5 wherein R₁ is CH₃, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table E-16 provides 13 compounds E-16.001 to E-16.013 of formula J-5 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

15 Table E-17 provides 13 compounds E-17.001 to E-17.013 of formula J-5 wherein R₁ is CH₃, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table E-18 provides 13 compounds E-18.001 to E-18.013 of formula J-5 wherein R₁ is CH₃, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

20 Table E-19 provides 13 compounds E-19.001 to E-19.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table E-20 provides 13 compounds E-20.001 to E-20.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is [5-(trifluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table E-21 provides 13 compounds E-21.001 to E-21.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

25 Table E-22 provides 13 compounds E-22.001 to E-22.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2-difluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

Table E-23 provides 13 compounds E-23.001 to E-23.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

30 Table E-24 provides 13 compounds E-24.001 to E-24.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is [5-(2,2,2-trifluoroethoxy)-2-pyridyl] and R₂ is as defined in table Z.

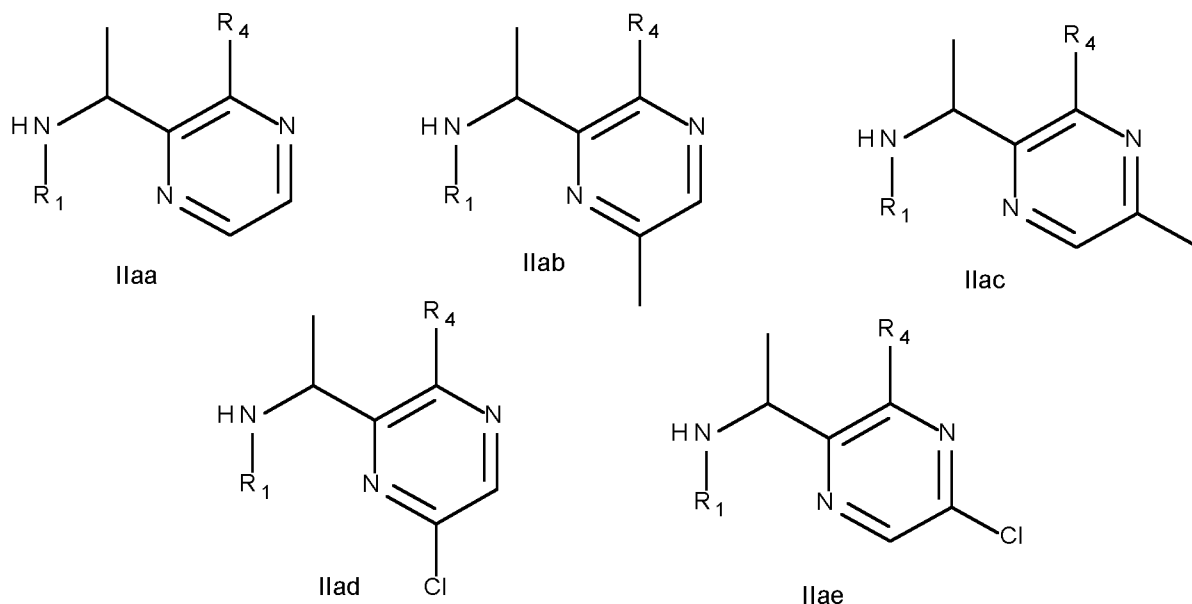
Table E-25 provides 13 compounds E-25.001 to E-25.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)pyrimidin-2-yl] and R₂ is as defined in table Z.

Table E-26 provides 13 compounds E-26.001 to E-26.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is [5-(difluoromethoxy)-2-pyridyl] and R₂ is as defined in table Z.

35 Table E-27 provides 13 compounds E-27.001 to E-27.013 of formula J-5 wherein R₁ is CH₂Cyp, R₄ is (5-cyano-2-pyridyl) and R₂ is as defined in table Z.

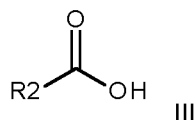
-44-

Also made available are certain intermediate compounds of the amine of formulae IIa to IIe, some of which are novel.



Specific examples of compounds of formula IIa to IIe are where R₁ and R₄ are as defined for a compound in Tables A-1 to A-27.

Further made available are certain intermediate compounds of the acid of formulae III, some of which are novel.



Specific examples of compounds of formula III are where R₂ is as defined in table Z.

Also made available are compounds of formulae VI, XV, XVa, XLII, XLV, XXXI, XXXII, XXXIV, XXV, XXX, XXXVIII, XXXIX, XIXa, XIXb, XXa, XLII, XLIII, and XLV, wherein, as applicable, the substituents R₁, R₂, R₃, R_{5a}, R_{5b} and R₄ (corresponding to the ring having R_{4a} and A₂), are as defined in any one of the compounds in Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27. An especially preferred enantiomer of the compounds of formulae VI, XV, XVa, XLII, XLV, XXXI, XXXII, XXXIV, XXV, XXX, XXXVIII, XXXIX, XIXa, XIXb, XXa, XLII, XLIII, and XLV, , as applicable, is the enantiomer having the same spatial arrangement at the stereogenic centre as depicted in formula I'a or I'-A.

The present invention also makes available

- a compound of formula II, wherein A₂, R₁, R₃, R_{4a}, R_{5a}, and R_{5b} are as defined for formula I; accordingly preferred embodiments of A₂, R₁, R₃, R_{4a}, R_{5a}, and R_{5b} for a compound of formula

-45-

I are likewise preferred embodiments of A₂, R₁, R₃, R_{4a}, R_{5a}, and R_{5b} for a compound of formula II;

- a compound of formulae IIaa, IIab, IIac, IIad and IIae, wherein R₁ is as defined formula I. and R₄ is the cyclic group containing A₂ and the substituent R_{4a} in formula I, wherein A₂ and R_{4a} are as defined as defined formula I; accordingly preferred embodiments of R₁, A₂ and R_{4a} for a compound of formula I are likewise preferred embodiments of R₁, A₂ and R_{4a} for a any one of a compound of formulae IIaa, IIab, IIac, IIad and IIae;
- a compound of formula III, wherein R₂ is as defined for formula I; accordingly preferred embodiments of R₂ for a compound of formula I are likewise preferred embodiments of R₂ for a compound of formula III;
- a compound of formula VI, wherein A₂, R₃, R_{4a}, R_{5a}, and R_{5b} are as defined for formula I; accordingly preferred embodiments of A₂, R₃, R_{4a}, R_{5a}, and R_{5b} for a compound of formula I are likewise preferred embodiments of A₂, R₃, R_{4a}, R_{5a}, and R_{5b} for a compound of a compound of formula XV, wherein A₂, R₃, R_{4a}, R_{5a}, and R_{5b} are as defined for formula I; accordingly preferred embodiments of A₂, R₃, R_{4a}, R_{5a}, and R_{5b} for a compound of formula I are likewise preferred embodiments of A₂, R₃, R_{4a}, R_{5a}, and R_{5b} for a compound of formula XV;
- a compound of formula XVa, wherein A₂, R₃, R_{4a}, R_{5a}, and R_{5b} are as defined for formula I; accordingly preferred embodiments of A₂, R₃, R_{4a}, R_{5a}, and R_{5b} for a compound of formula I are likewise preferred embodiments of A₂, R₃, R_{4a}, R_{5a}, and R_{5b} for a compound of formula XVa;
- a compound of formula XLII, wherein R₃, R_{5a}, and R_{5b} are as defined for formula I, X₀₇ is a leaving group like, for example, chlorine, bromine, iodine, and Z₃ is NPhth or NBoc₂; accordingly preferred embodiments of R₃, R_{5a}, and R_{5b} for a compound of formula I are likewise preferred embodiments of R₃, R_{5a}, and R_{5b} for a compound of formula XLII;
- a compound of formula XLV, wherein R₃, R_{5a}, and R_{5b} are as defined for formula I and X₀₇ is a leaving group like, for example, chlorine, bromine, iodine; accordingly preferred embodiments of R₃, R_{5a}, and R_{5b} for a compound of formula I are likewise preferred embodiments of R₃, R_{5a}, and R_{5b} for a compound of formula XLV;
- a compound of formula XXXI, wherein A₂ and R₃ are as defined for formula I and Z_{4a} is R_{4a}, halogen or NH₂; accordingly preferred embodiments of A₂, R₃, and R_{4a} for a compound of formula I are likewise preferred embodiments of A₂, R₃ and R_{4a} for a compound of formula XXXI;
- a compound of formula XXXII, wherein A₂ and R₃ are as defined for formula I and Z_{4a} is R_{4a}, halogen or NH₂; accordingly preferred embodiments of A₂, R₃, and R_{4a} for a compound of formula I are likewise preferred embodiments of A₂, R₃ and R_{4a} for a compound of formula XXXII;
- a compound of formula XXXIV, wherein A₂ and R₃ are as defined for formula I, Z_{4a} is R_{4a}, halogen or NH₂, and Z_{5a} and Z_{5b} are independent of each other selected from R_{5a}, R_{5b}, halogen, NH₂ and OH; accordingly preferred embodiments of A₂, R₃, R_{4a}, R_{5a} and R_{5b} for a

compound of formula I are likewise preferred embodiments of A₂, R₃, R_{4a}, R_{5a} and R_{5b} for a compound of formula XXXIV;

- a compound of formula XXV, wherein A₂ and R₃ are as defined for formula I, and Z_{5a} and Z_{5b} are independent of each other selected from R_{5a}, R_{5b}, halogen, NH₂ and OH; accordingly preferred embodiments of A₂, R₃, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments A₂, R₃, R_{5a} and R_{5b} for a compound of formula XXV;

- a compound of formula XXX, wherein A₂, R₃ and R_{4a} are as defined for formula I, and Z_{5a} and Z_{5b} are independent of each other selected from R_{5a}, R_{5b}, halogen, NH₂ and OH; accordingly preferred embodiments of A₂, R₃, R_{4a}, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments A₂, R₃, R_{4a}, R_{5a} and R_{5b} for a compound of formula XXX;

- a compound of formula XXXVIII, wherein A₂, R₃, R_{5a} and R_{5b} are as defined for formula I, and Z_{4a} is R_{4a}, halogen or NH₂; accordingly preferred embodiments of A₂, R₃, R_{4a}, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments A₂, R₃, R_{4a}, R_{5a} and R_{5b} for a compound of formula XXXVIII;

- a compound of formula XXXIX, wherein A₂, R₃, R_{5a} and R_{5b} are as defined for formula I, and Z_{4a} is R_{4a}, halogen or NH₂; accordingly preferred embodiments of A₂, R₃, R_{4a}, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments A₂, R₃, R_{4a}, R_{5a} and R_{5b} for a compound of formula XXXIX;

- a compound of formula XIXa, wherein R₃, R_{5a} and R_{5b} are as defined for formula I, and X₀₇ is a leaving group, for example, chlorine, bromine, iodine; accordingly preferred embodiments of R₃, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments R₃, R_{5a} and R_{5b} for a compound of formula XIXa;

- a compound of formula XIXb, wherein R₁, R₃, R_{5a} and R_{5b} are as defined for formula I, and X₀₇ is a leaving group, for example, chlorine, bromine, iodine; accordingly preferred embodiments of R₁, R₃, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments R₁, R₃, R_{5a} and R_{5b} for a compound of formula XIXb;

- a compound of formula XXa, wherein A₁, R_{2a}, R_{2b}, R₁, R₃, R_{5a} and R_{5b} are as defined for formula I, and X₀₇ is a leaving group, for example, chlorine, bromine, iodine; accordingly preferred embodiments of A₁, R_{2a}, R_{2b}, R₁, R₃, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments A₁, R_{2a}, R_{2b}, R₁, R₃, R_{5a} and R_{5b} for a compound of formula XXa;

- a compound of formula XLII, wherein R₃, R_{5a} and R_{5b} are as defined for formula I, Z₃ is NPhth or NBoc₂ and X₀₇ is a leaving group, for example, chlorine, bromine, or iodine; accordingly preferred embodiments of R₃, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments R₃, R_{5a} and R_{5b} for a compound of formula XLII;

- a compound of formula XLIII, wherein R₃, R_{5a} and R_{5b} are as defined for formula I and X₀₇ is a leaving group, for example, chlorine, bromine, or iodine; accordingly preferred embodiments of R₃, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments R₃, R_{5a} and R_{5b} for a compound of formula XLIII; and

- a compound of formula XLV, wherein R₃, R_{5a} and R_{5b} are as defined for formula I and X₀₇ is a leaving group, for example, chlorine, bromine, or iodine; accordingly preferred embodiments of R₃, R_{5a} and R_{5b} for a compound of formula I are likewise preferred embodiments R₃, R_{5a} and R_{5b} for a compound of formula XLV.

5

The compounds of formula I according to the invention are preventively and/or curatively valuable active ingredients in the field of pest control, even at low rates of application, which have a very favorable biocidal spectrum and are well tolerated by warm-blooded species, fish and plants. The active ingredients according to the invention act against all or individual developmental stages of normally sensitive, but also resistant, animal pests, such as insects or representatives of the order Acarina. The insecticidal or acaricidal activity of the active ingredients according to the invention can manifest itself directly, i. e. in destruction of the pests, which takes place either immediately or only after some time has elapsed, for example during ecdysis, or indirectly, for example in a reduced oviposition and/or hatching rate.

15

Examples of the above mentioned animal pests are:

from the order *Acarina*, for example,

Acalitus spp, Aculus spp, Acaricalus spp, Aceria spp, Acarus siro, Amblyomma spp., Argas spp., Boophilus spp., Brevipalpus spp., Bryobia spp, Calipitimerus spp., Chorioptes spp., Dermanyssus gallinae, Dermatophagoides spp, Eotetranychus spp, Eriophyes spp., Hemitarsonemus spp, Hyalomma spp., Ixodes spp., Olygonychus spp, Ornithodoros spp., Polyphagotarsonus latus, Panonychus spp., Phyllocoptruta oleivora, Phytionemus spp, Polyphagotarsonemus spp, Psoroptes spp., Rhipicephalus spp., Rhizoglyphus spp., Sarcoptes spp., Steneotarsonemus spp, Tarsonemus spp. and Tetranychus spp.;

25

from the order *Anoplura*, for example,

Haematopinus spp., Linognathus spp., Pediculus spp., Pemphigus spp. and Phylloxera spp.;

from the order *Coleoptera*, for example,

Agriotes spp., Amphimallon majale, Anomala orientalis, Anthonomus spp., Aphodius spp, Astylus atromaculatus, Ataenius spp, Atomaria linearis, Chaetocnema tibialis, Cerotoma spp, Conoderus spp, Cosmopolites spp., Cotinis nitida, Curculio spp., Cyclocephala spp, Dermestes spp., Diabrotica spp., Diloboderus abderus, Epilachna spp., Eremnus spp., Heteronychus arator, Hypothenemus hampei, Lagria vilosa, Leptinotarsa decemlineata, Lissorhoptrus spp., Liogenys spp, Maecolaspis spp, Maladera castanea, Megascelis spp, Meligethes aeneus, Melolontha spp., Myochrous armatus, Orycaephilus spp., Otiorhynchus spp., Phyllophaga spp, Phlyctinus spp., Popillia spp., Psylliodes spp., Rhyssomatus aubtilis, Rhizopertha spp., Scarabeidae, Sitophilus spp., Sitotroga spp., Somaticus spp, Sphenophorus spp, Sternechus subsignatus, Tenebrio spp., Tribolium spp. and Trogoderma spp.;

35

from the order *Diptera*, for example,

Aedes spp., Anopheles spp, Antherigona soccata, Bactrocea oleae, Bibio hortulanus, Bradysia spp, Calliphora erythrocephala, Ceratitis spp., Chrysomyia spp., Culex spp., Cuterebra spp., Dacus spp.,

-48-

Delia spp, Drosophila melanogaster, Fannia spp., Gastrophilus spp., Geomyza tripunctata, Glossina spp., Hypoderma spp., Hyppobosca spp., Liriomyza spp., Lucilia spp., Melanagromyza spp., Musca spp., Oestrus spp., Orseolia spp., Oscinella frit, Pegomyia hyoscyami, Phorbia spp., Rhagoletis spp, Rivelia quadrifasciata, Scatella spp, Sciara spp., Stomoxys spp., Tabanus spp., Tannia spp. and

5 Tipula spp.;

from the order *Hemiptera*, for example,

Acanthocoris scabrator, Acrosternum spp, Adelphocoris lineolatus, Aleurodes spp., Amblypelta nitida, Bathycoelia thalassina, Blissus spp, Cimex spp., Clavigralla tomentosicollis, Creontiades spp, Distantiella theobroma, Dichelops furcatus, Dysdercus spp., Edessa spp, Euchistus spp., Eurydema pulchrum, Eurygaster spp., Halyomorpha halys, Horcias nobilellus, Leptocoris spp., Lygus spp, Margarodes spp, Murgantia histrionic, Neomegalotomus spp, Nesidiocoris tenuis, Nezara spp., Nysius simulans, Oebalus insularis, Piesma spp., Piezodorus spp, Rhodnius spp., Sahlbergella singularis, Scaptocoris castanea, Scotinophara spp. , Thyanta spp , Triatoma spp., Vatiga illudens;

10 Acyrthosium pisum, Adalges spp, Agalliana ensigera, Agonoscena targionii, Aleurodicus spp, Aleurocanthus spp, Aleurolobus barodensis, Aleurothrixus floccosus, Aleyrodes brassicae, Amarasca biguttula, Amritodus atkinsoni, Aonidiella spp., Aphididae, Aphis spp., Aspidiotus spp., Aulacorthum solani, Bactericera cockerelli, Bemisia spp, Brachycaudus spp, Brevicoryne brassicae, Cacopsylla spp, Cavariella aegopodii Scop., Ceroplaster spp., Chrysomphalus aonidium, Chrysomphalus dictyospermi, Cicadella spp, Cofana spectra, Cryptomyzus spp, Cicadulina spp, Coccus hesperidum,

15 Dalbulus maidis, Dialeurodes spp, Diaphorina citri, Diuraphis noxia, Dysaphis spp, Empoasca spp., Eriosoma larigerum, Erythroneura spp., Gascardia spp., Glycaspis brimblecombei, Hyadaphis pseudobrassicae, Hyalopterus spp, Hyperomyzus pallidus, Idioscopus clypealis, Jacobiasca lybica, Laodelphax spp., Lecanium corni, Lepidosaphes spp., Lopaphis erysimi, Lyogenys maidis, Macrosiphum spp., Mahanarva spp, Metcalfa pruinosa, Metopolophium dirhodum, Myndus crudus,

20 Myzus spp., Neotoxoptera sp, Nephotettix spp., Nilaparvata spp., Nippolachnus piri Mats, Odonaspis ruthae, Oregma lanigera Zehnter, Parabemisia myricae, Paratrioza cockerelli, Parlatoria spp., Pemphigus spp., Peregrinus maidis, Perkinsiella spp, Phorodon humuli, Phylloxera spp, Planococcus spp., Pseudaulacaspis spp., Pseudococcus spp., Pseudatomoscelis seriatus, Psylla spp., Pulvinaria aethiopica, Quadraspidiotus spp., Quesada gigas, Recilia dorsalis, Rhopalosiphum spp., Saissetia spp., Scaphoideus spp., Schizaphis spp., Sitobion spp., Sogatella furcifera, Spissistilus festinus,

30 Tarophagus Proserpina, Toxoptera spp, Trialeurodes spp, Tridiscus sporoboli, Trionymus spp, Trioza erytrae , Unaspis citri, Zygina flammigera, Zyginidia scutellaris, ;

from the order *Hymenoptera*, for example,

Acromyrmex, Arge spp, Atta spp., Cephus spp., Diprion spp., Diprionidae, Gilpinia polytoma, Hoplocampa spp., Lasius spp., Monomorium pharaonis, Neodiprion spp., Pogonomymex spp, Slenopsis invicta, Solenopsis spp. and Vespa spp.;

from the order *Isoptera*, for example,

Coptotermes spp, Cornitermes cumulans, Incisitermes spp, Macrotermes spp, Mastotermes spp, Microtermes spp, Reticulitermes spp.; Solenopsis geminate

from the order *Lepidoptera*, for example,

Acleris spp., Adoxophyes spp., Aegeria spp., Agrotis spp., Alabama argillaceae, Amylois spp.,
 Anticarsia gemmatilis, Archips spp., Argyroresthia spp, Argyrotaenia spp., Autographa spp., Bucculatrix
 thurberiella, Busseola fusca, Cadra cautella, Carposina nipponensis, Chilo spp., Choristoneura spp.,
 5 Chrysoteuchia topiaria, Clysia ambiguella, Cnaphalocrocis spp., Cnephasia spp., Cochylis spp.,
 Coleophora spp., Colias lesbia, Cosmophila flava, Crambus spp, Crocidolomia binotalis, Cryptophlebia
 leucotreta, Cydalima perspectalis, Cydia spp., Diaphania perspectalis, Diatraea spp., Diparopsis
 castanea, Earias spp., Elasmopalpus lignosellus, Eldana saccharina, Ephestia spp., Epinotia spp.,
 Estigmene acrea, Etiella zinckenella, Eucosma spp., Eupoecilia ambiguella, Euproctis spp., Euxoa
 10 spp., Feltia jaculiferia, Grapholita spp., Hedya nubiferana, Heliothis spp., Hellula undalis,
 Herpetogramma spp, Hyphantria cunea, Keiferia lycopersicella, Lasmopalpus lignosellus, Leucoptera
 scitella, Lithocolletis spp., Lobesia botrana, Loxostege bifidalis, Lymantria spp., Lyonetia spp.,
 Malacosoma spp., Mamestra brassicae, Manduca sexta, Mythimna spp, Noctua spp, Operophtera
 spp., Orniodes indica, Ostrinia nubilalis, Pammene spp., Pandemis spp., Panolis flammea, Papaipema
 15 nebris, Pectinophora gossypiella, Perileucoptera coffeella, Pseudaletia unipuncta, Phthorimaea
 operculella, Pieris rapae, Pieris spp., Plutella xylostella, Prays spp., Pseudoplusia spp, Rachiplusia nu,
 Richia albicosta, Scirpophaga spp., Sesamia spp., Sparganothis spp., Spodoptera spp., Sylepta
 derogate, Synanthedon spp., Thaumetopoea spp., Tortrix spp., Trichoplusia ni, Tuta absoluta, and
 Yponomeuta spp.;

20 from the order *Mallophaga*, for example,

Damalinea spp. and Trichodectes spp.;

from the order *Orthoptera*, for example,

Blatta spp., Blattella spp., Gryllotalpa spp., Leucophaea maderae, Locusta spp., Neocurtilla
 hexadactyla, Periplaneta spp. , Scapteriscus spp, and Schistocerca spp.;

25 from the order *Psocoptera*, for example,

Liposcelis spp.;

from the order *Siphonaptera*, for example,

Ceratophyllus spp., Ctenocephalides spp. and Xenopsylla cheopis;

from the order *Thysanoptera*, for example,

30 Calliothrips phaseoli, Frankliniella spp., Heliothrips spp, Hercinothrips spp., Parthenothrips spp,
 Scirtothrips aurantii, Sericothrips variabilis, Taeniothrips spp., Thrips spp;

from the order *Thysanura*, for example, Lepisma saccharina.

In a further aspect, the invention may also relate to a method of controlling damage to plant and parts

35 thereof by plant parasitic nematodes (Endoparasitic-, Semiendoparasitic- and Ectoparasitic
 nematodes), especially plant parasitic nematodes such as root knot nematodes, Meloidogyne hapla,
 Meloidogyne incognita, Meloidogyne javanica, Meloidogyne arenaria and other Meloidogyne species;
 cyst-forming nematodes, Globodera rostochiensis and other Globodera species; Heterodera avenae,
 Heterodera glycines, Heterodera schachtii, Heterodera trifolii, and other Heterodera species; Seed gall

- nematodes, *Anguina* species; Stem and foliar nematodes, *Aphelenchoides* species; Sting nematodes, *Belonolaimus longicaudatus* and other *Belonolaimus* species; Pine nematodes, *Bursaphelenchus xylophilus* and other *Bursaphelenchus* species; Ring nematodes, *Criconema* species, *Criconemella* species, *Criconemoides* species, *Mesocriconema* species; Stem and bulb nematodes, *Ditylenchus destructor*, *Ditylenchus dipsaci* and other *Ditylenchus* species; Awl nematodes, *Dolichodorus* species; Spiral nematodes, *Heliocotylenchus multicinctus* and other *Helicotylenchus* species; Sheath and sheathoid nematodes, *Hemicyclophora* species and *Hemicriconemoides* species; *Hirshmanniella* species; Lance nematodes, *Hoploaimus* species; false rootknot nematodes, *Nacobbus* species; Needle nematodes, *Longidorus elongatus* and other *Longidorus* species; Pin nematodes, *Pratylenchus* species; Lesion nematodes, *Pratylenchus neglectus*, *Pratylenchus penetrans*, *Pratylenchus curvatus*, *Pratylenchus goodeyi* and other *Pratylenchus* species; Burrowing nematodes, *Radopholus similis* and other *Radopholus* species; Reniform nematodes, *Rotylenchus robustus*, *Rotylenchus reniformis* and other *Rotylenchus* species; *Scutellonema* species; Stubby root nematodes, *Trichodorus primitivus* and other *Trichodorus* species, *Paratrichodorus* species; Stunt nematodes, *Tylenchorhynchus claytoni*, *Tylenchorhynchus dubius* and other *Tylenchorhynchus* species; Citrus nematodes, *Tylenchulus* species; Dagger nematodes, *Xiphinema* species; and other plant parasitic nematode species, such as *Subanguina* spp., *Hypsoperine* spp., *Macroposthonia* spp., *Melinius* spp., *Punctodera* spp., and *Quinisulcius* spp..
- The compounds of the invention may also have activity against the molluscs. Examples of which include, for example, *Ampullariidae*; *Arion* (*A. ater*, *A. circumscriptus*, *A. hortensis*, *A. rufus*); *Bradybaenidae* (*Bradybaena fruticum*); *Cepaea* (*C. hortensis*, *C. Nemoralis*); *ochlodina*; *Deroceras* (*D. agrestis*, *D. empiricorum*, *D. laeve*, *D. reticulatum*); *Discus* (*D. rotundatus*); *Euomphalia*; *Galba* (*G. trunculata*); *Helicelia* (*H. itala*, *H. obvia*); *Helicidae* *Helicigona arbustorum*); *Helicodiscus*; *Helix* (*H. aperta*); *Limax* (*L. cinereoniger*, *L. flavus*, *L. marginatus*, *L. maximus*, *L. tenellus*); *Lymnaea*; *Milax* (*M. gagates*, *M. marginatus*, *M. sowerbyi*); *Opeas*; *Pomacea* (*P. canaticulata*); *Vallonia* and *Zanitoides*.

The active ingredients according to the invention can be used for controlling, i. e. containing or destroying, pests of the abovementioned type which occur in particular on plants, especially on useful plants and ornamentals in agriculture, in horticulture and in forests, or on organs, such as fruits, flowers, foliage, stalks, tubers or roots, of such plants, and in some cases even plant organs which are formed at a later point in time remain protected against these pests.

Suitable target crops are, in particular, cereals, such as wheat, barley, rye, oats, rice, maize or sorghum; beet, such as sugar or fodder beet; fruit, for example pomaceous fruit, stone fruit or soft fruit, such as apples, pears, plums, peaches, almonds, cherries or berries, for example strawberries, raspberries or blackberries; leguminous crops, such as beans, lentils, peas or soya; oil crops, such as oilseed rape, mustard, poppies, olives, sunflowers, coconut, castor, cocoa or ground nuts; cucurbits, such as pumpkins, cucumbers or melons; fibre plants, such as cotton, flax, hemp or jute; citrus fruit,

such as oranges, lemons, grapefruit or tangerines; vegetables, such as spinach, lettuce, asparagus, cabbages, carrots, onions, tomatoes, potatoes or bell peppers; Lauraceae, such as avocado, Cinnamomum or camphor; and also tobacco, nuts, coffee, eggplants, sugarcane, tea, pepper, grapevines, hops, the plantain family and latex plants.

5

The compositions and/or methods of the present invention may be also used on any ornamental and/or vegetable crops, including flowers, shrubs, broad-leaved trees and evergreens.

For example the invention may be used on any of the following ornamental species: *Ageratum* spp.,

- 10 *Alonsoa* spp., *Anemone* spp., *Anisodonteia capensis*, *Anthemis* spp., *Antirrhinum* spp., *Aster* spp., *Begonia* spp. (e.g. *B. elatior*, *B. semperflorens*, *B. tuberosa*), *Bougainvillea* spp., *Brachycome* spp., *Brassica* spp. (ornamental), *Calceolaria* spp., *Capsicum annuum*, *Catharanthus roseus*, *Canna* spp., *Centaurea* spp., *Chrysanthemum* spp., *Cineraria* spp. (*C. maritime*), *Coreopsis* spp., *Crassula coccinea*, *Cuphea ignea*, *Dahlia* spp., *Delphinium* spp., *Dicentra spectabilis*, *Dorotheanthus* spp.,
- 15 *Eustoma grandiflorum*, *Forsythia* spp., *Fuchsia* spp., *Geranium gnaphalium*, *Gerbera* spp., *Gomphrena globosa*, *Heliotropium* spp., *Helianthus* spp., *Hibiscus* spp., *Hortensia* spp., *Hydrangea* spp., *Hypoestes phyllostachya*, *Impatiens* spp. (*I. Walleriana*), *Iresines* spp., *Kalanchoe* spp., *Lantana camara*, *Lavatera trimestris*, *Leonotis leonurus*, *Lilium* spp., *Mesembryanthemum* spp., *Mimulus* spp., *Monarda* spp., *Nemesia* spp., *Tagetes* spp., *Dianthus* spp. (carnation), *Canna* spp., *Oxalis* spp., *Bellis*
- 20 spp., *Pelargonium* spp. (*P. peltatum*, *P. Zonale*), *Viola* spp. (pansy), *Petunia* spp., *Phlox* spp., *Plectranthus* spp., *Poinsettia* spp., *Parthenocissus* spp. (*P. quinquefolia*, *P. tricuspidata*), *Primula* spp., *Ranunculus* spp., *Rhododendron* spp., *Rosa* spp. (rose), *Rudbeckia* spp., *Saintpaulia* spp., *Salvia* spp., *Scaevola aemola*, *Schizanthus wisetonensis*, *Sedum* spp., *Solanum* spp., *Surfinia* spp., *Tagetes* spp., *Nicotinia* spp., *Verbena* spp., *Zinnia* spp. and other bedding plants.

25

For example the invention may be used on any of the following vegetable species: *Allium* spp. (*A. sativum*, *A. cepa*, *A. oschaninii*, *A. Porrum*, *A. ascalonicum*, *A. fistulosum*), *Anthriscus cerefolium*, *Apium graveolus*, *Asparagus officinalis*, *Beta vulgaris*, *Brassica* spp. (*B. Oleracea*, *B. Pekinensis*, *B. rapa*), *Capsicum annuum*, *Cicer arietinum*, *Cichorium endivia*, *Cichorium* spp. (*C. intybus*, *C. endivia*),

30 *Citrullus lanatus*, *Cucumis* spp. (*C. sativus*, *C. melo*), *Cucurbita* spp. (*C. pepo*, *C. maxima*), *Cyanara* spp. (*C. scolymus*, *C. cardunculus*), *Daucus carota*, *Foeniculum vulgare*, *Hypericum* spp., *Lactuca sativa*, *Lycopersicon* spp. (*L. esculentum*, *L. lycopersicum*), *Mentha* spp., *Ocimum basilicum*, *Petroselinum crispum*, *Phaseolus* spp. (*P. vulgaris*, *P. coccineus*), *Pisum sativum*, *Raphanus sativus*, *Rheum raphaniticum*, *Rosemarinus* spp., *Salvia* spp., *Scorzonera hispanica*, *Solanum melongena*,

35 *Spinacea oleracea*, *Valerianella* spp. (*V. locusta*, *V. eriocarpa*) and *Vicia faba*.

Preferred ornamental species include African violet, *Begonia*, *Dahlia*, *Gerbera*, *Hydrangea*, *Verbena*, *Rosa*, *Kalanchoe*, *Poinsettia*, *Aster*, *Centaurea*, *Coreopsis*, *Delphinium*, *Monarda*, *Phlox*, *Rudbeckia*,

-52-

Sedum, Petunia, Viola, Impatiens, Geranium, Chrysanthemum, Ranunculus, Fuchsia, Salvia, Hortensia, rosemary, sage, St. Johnswort, mint, sweet pepper, tomato and cucumber.

The active ingredients according to the invention are especially suitable for controlling *Aphis craccivora*, *Diabrotica balteata*, *Heliothis virescens*, *Myzus persicae*, *Plutella xylostella* and *Spodoptera littoralis* in cotton, vegetable, maize, rice and soya crops. The active ingredients according to the invention are further especially suitable for controlling *Mamestra* (preferably in vegetables), *Cydia pomonella* (preferably in apples), *Empoasca* (preferably in vegetables, vineyards), *Leptinotarsa* (preferably in potatoes) and *Chilo suppressalis* (preferably in rice).

The compounds of formula I are particularly suitable for control of

- a pest of the order Hemiptera, for example, one or more of the species *Bemisia tabaci*, *Aphis craccivora*, *Myzus persicae*, *Rhopalosiphum padi*, *Nilaparvata lugens*, and *Euschistus heros* (preferably in vegetables, soybeans, and sugarcane);
- a pest of the order Lepidoptera, for example, one or more of the species *Spodoptera littoralis*, *Spodoptera frugiperda*, *Plutella xylostella*, *Cnaphalocrocis medinalis*, *Cydia pomonella*, *Chrysodeixis includens*, *Chilo suppressalis*, *Elasmopalpus lignosellus*, *Pseudoplusia includens*, and *Tuta absoluta* (preferably in vegetables and corn);
- a pest of the order Thysanoptera, such as the family Thripidae, for example, one or more of *Thrips tabaci* and *Frankliniella occidentalis* (preferably in vegetables); and
- soil pests (such as of the order Coleoptera), for example, the species *Diabrotica balteata*, *Agriotes* spp. and *Leptinotarsa decemlineata* (preferably in vegetables and corn).

The term "crops" is to be understood as including also crop plants which have been so transformed by the use of recombinant DNA techniques that they are capable of synthesising one or more selectively acting toxins, such as are known, for example, from toxin-producing bacteria, especially those of the genus *Bacillus*.

Toxins that can be expressed by such transgenic plants include, for example, insecticidal proteins, for example insecticidal proteins from *Bacillus cereus* or *Bacillus popilliae*; or insecticidal proteins from *Bacillus thuringiensis*, such as δ -endotoxins, e.g. Cry1Ab, Cry1Ac, Cry1F, Cry1Fa2, Cry2Ab, Cry3A, Cry3Bb1 or Cry9C, or vegetative insecticidal proteins (Vip), e.g. Vip1, Vip2, Vip3 or Vip3A; or insecticidal proteins of bacteria colonising nematodes, for example *Photorhabdus* spp. or *Xenorhabdus* spp., such as *Photorhabdus luminescens*, *Xenorhabdus nematophilus*; toxins produced by animals, such as scorpion toxins, arachnid toxins, wasp toxins and other insect-specific neurotoxins; toxins produced by fungi, such as *Streptomyces* toxins, plant lectins, such as pea lectins, barley lectins or snowdrop lectins; agglutinins; proteinase inhibitors, such as trypsin inhibitors, serine protease inhibitors, patatin, cystatin, papain inhibitors; ribosome-inactivating proteins (RIP), such as ricin, maize-RIP, abrin, luffin, saporin or bryodin; steroid metabolism enzymes, such as

3-hydroxysteroidoxidase, ecdysteroid-UDP-glycosyl-transferase, cholesterol oxidases, ecdysone inhibitors, HMG-COA-reductase, ion channel blockers, such as blockers of sodium or calcium channels, juvenile hormone esterase, diuretic hormone receptors, stilbene synthase, bibenzyl synthase, chitinases and glucanases.

5

In the context of the present invention there are to be understood by δ -endotoxins, for example Cry1Ab, Cry1Ac, Cry1F, Cry1Fa2, Cry2Ab, Cry3A, Cry3Bb1 or Cry9C, or vegetative insecticidal proteins (Vip), for example Vip1, Vip2, Vip3 or Vip3A, expressly also hybrid toxins, truncated toxins and modified toxins. Hybrid toxins are produced recombinantly by a new combination of different domains of those proteins (see, for example, WO 02/15701). Truncated toxins, for example a truncated Cry1Ab, are known. In the case of modified toxins, one or more amino acids of the naturally occurring toxin are replaced. In such amino acid replacements, preferably non-naturally present protease recognition sequences are inserted into the toxin, such as, for example, in the case of Cry3A055, a cathepsin-G-recognition sequence is inserted into a Cry3A toxin (see WO 03/018810).

15

Examples of such toxins or transgenic plants capable of synthesising such toxins are disclosed, for example, in EP-A-0 374 753, WO 93/07278, WO 95/34656, EP-A-0 427 529, EP-A-451 878 and WO 03/052073.

20

The processes for the preparation of such transgenic plants are generally known to the person skilled in the art and are described, for example, in the publications mentioned above. CryI-type deoxyribonucleic acids and their preparation are known, for example, from WO 95/34656, EP-A-0 367 474, EP-A-0 401 979 and WO 90/13651.

25

The toxin contained in the transgenic plants imparts to the plants tolerance to harmful insects. Such insects can occur in any taxonomic group of insects, but are especially commonly found in the beetles (Coleoptera), two-winged insects (Diptera) and moths (Lepidoptera).

30

Transgenic plants containing one or more genes that code for an insecticidal resistance and express one or more toxins are known and some of them are commercially available. Examples of such plants are: YieldGard® (maize variety that expresses a Cry1Ab toxin); YieldGard Rootworm® (maize variety that expresses a Cry3Bb1 toxin); YieldGard Plus® (maize variety that expresses a Cry1Ab and a Cry3Bb1 toxin); Starlink® (maize variety that expresses a Cry9C toxin); Herculex I® (maize variety that expresses a Cry1Fa2 toxin and the enzyme phosphinothricine N-acetyltransferase (PAT) to achieve tolerance to the herbicide glufosinate ammonium); NuCOTN 33B® (cotton variety that expresses a Cry1Ac toxin); Bollgard I® (cotton variety that expresses a Cry1Ac toxin); Bollgard II® (cotton variety that expresses a Cry1Ac and a Cry2Ab toxin); VipCot® (cotton variety that expresses a Vip3A and a Cry1Ab toxin); NewLeaf® (potato variety that expresses a Cry3A toxin); NatureGard®,

35

Agrisure® GT Advantage (GA21 glyphosate-tolerant trait), Agrisure® CB Advantage (Bt11 corn borer (CB) trait) and Protecta®.

Further examples of such transgenic crops are:

- 5 1. **Bt11 Maize** from Syngenta Seeds SAS, Chemin de l'Hobit 27, F-31 790 St. Sauveur, France, registration number C/FR/96/05/10. Genetically modified *Zea mays* which has been rendered resistant to attack by the European corn borer (*Ostrinia nubilalis* and *Sesamia nonagrioides*) by transgenic expression of a truncated Cry1Ab toxin. Bt11 maize also transgenically expresses the enzyme PAT to achieve tolerance to the herbicide glufosinate ammonium.
- 10 2. **Bt176 Maize** from Syngenta Seeds SAS, Chemin de l'Hobit 27, F-31 790 St. Sauveur, France, registration number C/FR/96/05/10. Genetically modified *Zea mays* which has been rendered resistant to attack by the European corn borer (*Ostrinia nubilalis* and *Sesamia nonagrioides*) by transgenic expression of a Cry1Ab toxin. Bt176 maize also transgenically expresses the enzyme PAT to achieve tolerance to the herbicide glufosinate ammonium.
- 15 3. **MIR604 Maize** from Syngenta Seeds SAS, Chemin de l'Hobit 27, F-31 790 St. Sauveur, France, registration number C/FR/96/05/10. Maize which has been rendered insect-resistant by transgenic expression of a modified Cry3A toxin. This toxin is Cry3A055 modified by insertion of a cathepsin-G-protease recognition sequence. The preparation of such transgenic maize plants is described in WO 03/018810.
- 20 4. **MON 863 Maize** from Monsanto Europe S.A. 270-272 Avenue de Tervuren, B-1150 Brussels, Belgium, registration number C/DE/02/9. MON 863 expresses a Cry3Bb1 toxin and has resistance to certain Coleoptera insects.
5. **IPC 531 Cotton** from Monsanto Europe S.A. 270-272 Avenue de Tervuren, B-1150 Brussels, Belgium, registration number C/ES/96/02.
- 25 6. **1507 Maize** from Pioneer Overseas Corporation, Avenue Tedesco, 7 B-1160 Brussels, Belgium, registration number C/NL/00/10. Genetically modified maize for the expression of the protein Cry1F for achieving resistance to certain Lepidoptera insects and of the PAT protein for achieving tolerance to the herbicide glufosinate ammonium.
7. **NK603 × MON 810 Maize** from Monsanto Europe S.A. 270-272 Avenue de Tervuren, B-1150 Brussels, Belgium, registration number C/GB/02/M3/03. Consists of conventionally bred hybrid maize varieties by crossing the genetically modified varieties NK603 and MON 810. NK603 × MON 810 Maize transgenically expresses the protein CP4 EPSPS, obtained from *Agrobacterium sp.* strain CP4, which imparts tolerance to the herbicide Roundup® (contains glyphosate), and also a Cry1Ab toxin obtained from *Bacillus thuringiensis subsp. kurstaki* which brings about tolerance to certain
- 35 Lepidoptera, include the European corn borer.

Transgenic crops of insect-resistant plants are also described in BATS (Zentrum für Biosicherheit und Nachhaltigkeit, Zentrum BATS, Clarastrasse 13, 4058 Basel, Switzerland) Report 2003, (<http://bats.ch>).

-55-

The term "crops" is to be understood as including also crop plants which have been so transformed by the use of recombinant DNA techniques that they are capable of synthesising antipathogenic substances having a selective action, such as, for example, the so-called "pathogenesis-related proteins" (PRPs, see e.g. EP-A-0 392 225). Examples of such antipathogenic substances and transgenic plants capable of synthesising such antipathogenic substances are known, for example, from EP-A-0 392 225, WO 95/33818 and EP-A-0 353 191. The methods of producing such transgenic plants are generally known to the person skilled in the art and are described, for example, in the publications mentioned above.

Crops may also be modified for enhanced resistance to fungal (for example Fusarium, Anthracnose, or Phytophthora), bacterial (for example Pseudomonas) or viral (for example potato leafroll virus, tomato spotted wilt virus, cucumber mosaic virus) pathogens.

Crops also include those that have enhanced resistance to nematodes, such as the soybean cyst nematode.

Crops that are tolerance to abiotic stress include those that have enhanced tolerance to drought, high salt, high temperature, chill, frost, or light radiation, for example through expression of NF-YB or other proteins known in the art.

Antipathogenic substances which can be expressed by such transgenic plants include, for example, ion channel blockers, such as blockers for sodium and calcium channels, for example the viral KP1, KP4 or KP6 toxins; stilbene synthases; bibenzyl synthases; chitinases; glucanases; the so-called "pathogenesis-related proteins" (PRPs; see e.g. EP-A-0 392 225); antipathogenic substances produced by microorganisms, for example peptide antibiotics or heterocyclic antibiotics (see e.g. WO 95/33818) or protein or polypeptide factors involved in plant pathogen defence (so-called "plant disease resistance genes", as described in WO 03/000906).

Further areas of use of the compositions according to the invention are the protection of stored goods and store rooms and the protection of raw materials, such as wood, textiles, floor coverings or buildings, and also in the hygiene sector, especially the protection of humans, domestic animals and productive livestock against pests of the mentioned type.

The present invention provides a compound of the first aspect for use in therapy. The present invention provides a compound of the first aspect, for use in controlling parasites in or on an animal. The present invention further provides a compound of the first aspect, for use in controlling ectoparasites on an animal. The present invention further provides a compound of the first aspect, for use in preventing and/or treating diseases transmitted by ectoparasites.

-56-

The present invention provides the use of a compound of the first aspect, for the manufacture of a medicament for controlling parasites in or on an animal. The present invention further provides the use of a compound of the first aspect, for the manufacture of a medicament for controlling ectoparasites on an animal. The present invention further provides the use of a compound of the first aspect, for the manufacture of a medicament for preventing and/or treating diseases transmitted by ectoparasites.

The present invention provides the use of a compound of the first aspect, in controlling parasites in or on an animal. The present invention further provides the use of a compound of the first aspect, in controlling ectoparasites on an animal.

The term "controlling" when used in context of parasites in or on an animal refers to reducing the number of pests or parasites, eliminating pests or parasites and/or preventing further pest or parasite infestation.

The term "treating" when used in context of parasites in or on an animal refers to restraining, slowing, stopping or reversing the progression or severity of an existing symptom or disease. The term "preventing" when used in context of parasites in or on an animal refers to the avoidance of a symptom or disease developing in the animal.

The term "animal" when used in context of parasites in or on an animal may refer to a mammal and a non-mammal, such as a bird or fish. In the case of a mammal, it may be a human or non-human mammal. Non-human mammals include, but are not limited to, livestock animals and companion animals. Livestock animals include, but are not limited to, cattle, camellids, pigs, sheep, goats and horses. Companion animals include, but are not limited to, dogs, cats and rabbits.

A "parasite" is a pest which lives in or on the host animal and benefits by deriving nutrients at the host animal's expense. An "endoparasite" is a parasite which lives in the host animal. An "ectoparasite" is a parasite which lives on the host animal. Ectoparasites include, but are not limited to, acari, insects and crustaceans (e.g. sea lice). The Acari (or Acarina) sub-class comprises ticks and mites. Ticks include, but are not limited to, members of the following genera: *Rhipicaphalus*, for example, *Rhipicaphalus (Boophilus) microplus* and *Rhipicephalus sanguineus*; *Amblyomrna*; *Dermacentor*; *Haemaphysalis*; *Hyalomma*; *Ixodes*; *Rhipicentor*; *Margaropus*; *Argas*; *Otobius*; and *Ornithodoros*. Mites include, but are not limited to, members of the following genera: *Chorioptes*, for example *Chorioptes bovis*; *Psoroptes*, for example *Psoroptes ovis*; *Cheyletiella*; *Dermanyssus*; for example *Dermanyssus gallinae*; *Ornithonyssus*; *Demodex*, for example *Demodex canis*; *Sarcoptes*, for example *Sarcoptes scabiei*; and *Psorergates*. Insects include, but are not limited to, members of the orders: Siphonaptera, Diptera, Phthiraptera, Lepidoptera, Coleoptera and Homoptera. Members of the Siphonaptera order include, but are not limited to, *Ctenocephalides felis* and *Ctenocephalides canis*. Members of the Diptera order include, but are not limited to, *Musca spp.*; bot fly, for example *Gasterophilus intestinalis* and *Oestrus*

-57-

ovis; biting flies; horse flies, for example *Haematopota* spp. and *Tabanus* spp.; *haematobia*, for example *haematobia irritans*; *Stomoxys*; *Lucilia*; midges; and mosquitoes. Members of the Phthiraptera class include, but are not limited to, blood sucking lice and chewing lice, for example *Bovicola Ovis* and *Bovicola Bovis*.

5

The term "effective amount" when used in context of parasites in or on an animal refers to the amount or dose of the compound of the invention, or a salt thereof, which, upon single or multiple dose administration to the animal, provides the desired effect in or on the animal. The effective amount can be readily determined by the attending diagnostician, as one skilled in the art, by the use of known techniques and by observing results obtained under analogous circumstances. In determining the effective amount a number of factors are considered by the attending diagnostician, including, but not limited to: the species of mammal; its size, age, and general health; the parasite to be controlled and the degree of infestation; the specific disease or disorder involved; the degree of or involvement or the severity of the disease or disorder; the response of the individual; the particular compound administered; the mode of administration; the bioavailability characteristics of the preparation administered; the dose regimen selected; the use of concomitant medication; and other relevant circumstances.

The compounds of the invention may be administered to the animal by any route which has the desired effect including, but not limited to topically, orally, parenterally and subcutaneously. Topical administration is preferred. Formulations suitable for topical administration include, for example, solutions, emulsions and suspensions and may take the form of a pour-on, spot-on, spray-on, spray race or dip. In the alternative, the compounds of the invention may be administered by means of an ear tag or collar.

25

Salt forms of the compounds of the invention include both pharmaceutically acceptable salts and veterinary acceptable salts, which can be different to agrochemically acceptable salts. Pharmaceutically and veterinary acceptable salts and common methodology for preparing them are well known in the art. See, for example, Gould, P.L., "Salt selection for basic drugs", International Journal of Pharmaceutics, 33: 201 -217 (1986); Bastin, R.J., *et al.* "Salt Selection and Optimization Procedures for Pharmaceutical New Chemical Entities", Organic Process Research and Development, 4: 427-435 (2000); and Berge, S.M., *et al.*, "Pharmaceutical Salts", Journal of Pharmaceutical Sciences, 66: 1-19, (1977). One skilled in the art of synthesis will appreciate that the compounds of the invention are readily converted to and may be isolated as a salt, such as a hydrochloride salt, using techniques and conditions well known to one of ordinary skill in the art. In addition, one skilled in the art of synthesis will appreciate that the compounds of the invention are readily converted to and may be isolated as the corresponding free base from the corresponding salt.

35

The present invention also provides a method for controlling pests (such as mosquitoes and other disease vectors; see also http://www.who.int/malaria/vector_control/irs/en/). In one embodiment, the method for controlling pests comprises applying the compositions of the invention to the target pests, to their locus or to a surface or substrate by brushing, rolling, spraying, spreading or dipping. By way of example, an IRS (indoor residual spraying) application of a surface such as a wall, ceiling or floor surface is contemplated by the method of the invention. In another embodiment, it is contemplated to apply such compositions to a substrate such as non-woven or a fabric material in the form of (or which can be used in the manufacture of) netting, clothing, bedding, curtains and tents.

In one embodiment, the method for controlling such pests comprises applying a pesticidally effective amount of the compositions of the invention to the target pests, to their locus, or to a surface or substrate so as to provide effective residual pesticidal activity on the surface or substrate. Such application may be made by brushing, rolling, spraying, spreading or dipping the pesticidal composition of the invention. By way of example, an IRS application of a surface such as a wall, ceiling or floor surface is contemplated by the method of the invention so as to provide effective residual pesticidal activity on the surface. In another embodiment, it is contemplated to apply such compositions for residual control of pests on a substrate such as a fabric material in the form of (or which can be used in the manufacture of) netting, clothing, bedding, curtains and tents.

Substrates including non-woven, fabrics or netting to be treated may be made of natural fibres such as cotton, raffia, jute, flax, sisal, hessian, or wool, or synthetic fibres such as polyamide, polyester, polypropylene, polyacrylonitrile or the like. The polyesters are particularly suitable. The methods of textile treatment are known, e.g. WO 2008/151984, WO 2003/034823, US 5631072, WO 2005/64072, WO2006/128870, EP 1724392, WO 2005113886 or WO 2007/090739.

Further areas of use of the compositions according to the invention are the field of tree injection/trunk treatment for all ornamental trees as well all sort of fruit and nut trees.

In the field of tree injection/trunk treatment, the compounds according to the present invention are especially suitable against wood-boring insects from the order *Lepidoptera* as mentioned above and from the order *Coleoptera*, especially against woodborers listed in the following tables A and B:

Table A. Examples of exotic woodborers of economic importance.

Family	Species	Host or Crop Infested
Buprestidae	<i>Agrilus planipennis</i>	Ash
Cerambycidae	<i>Anoplua glabripennis</i>	Hardwoods
Scolytidae	<i>Xylosandrus crassiusculus</i>	Hardwoods
	<i>X. mutilatus</i>	Hardwoods
	<i>Tomicus piniperda</i>	Conifers

Table B. Examples of native woodborers of economic importance.

Family	Species	Host or Crop Infested
Buprestidae	<i>Agrilus anxius</i>	Birch
	<i>Agrilus politus</i>	Willow, Maple
	<i>Agrilus sayi</i>	Bayberry, Sweetfern
	<i>Agrilus vittaticollis</i>	Apple, Pear, Cranberry, Serviceberry, Hawthorn
	<i>Chrysobothris femorata</i>	Apple, Apricot, Beech, Boxelder, Cherry, Chestnut, Currant, Elm, Hawthorn, Hackberry, Hickory, Horsechestnut, Linden, Maple, Mountain-ash, Oak, Pecan, Pear, Peach, Persimmon, Plum, Poplar, Quince, Redbud, Serviceberry, Sycamore, Walnut, Willow
	<i>Texania campestris</i>	Basswood, Beech, Maple, Oak, Sycamore, Willow, Yellow-poplar
Cerambycidae	<i>Goes pulverulentus</i>	Beech, Elm, Nuttall, Willow, Black oak, Cherrybark oak, Water oak, Sycamore
	<i>Goes tigrinus</i>	Oak
	<i>Neoclytus acuminatus</i>	Ash, Hickory, Oak, Walnut, Birch, Beech, Maple, Eastern hophornbeam, Dogwood, Persimmon, Redbud, Holly, Hackberry, Black locust, Honeylocust, Yellow-poplar, Chestnut, Osage-orange, Sassafras, Lilac, Mountain-mahogany, Pear, Cherry, Plum, Peach, Apple, Elm, Basswood, Sweetgum
	<i>Neoptychodes trilineatus</i>	Fig, Alder, Mulberry, Willow, Nettle hackberry
	<i>Oberea ocellata</i>	Sumac, Apple, Peach, Plum, Pear, Currant, Blackberry
	<i>Oberea tripunctata</i>	Dogwood, Viburnum, Elm, Sourwood, Blueberry,

Family	Species	Host or Crop Infested
		Rhododendron, Azalea, Laurel, Poplar, Willow, Mulberry
	<i>Oncideres cingulata</i>	Hickory, Pecan, Persimmon, Elm, Sourwood, Basswood, Honeylocust, Dogwood, Eucalyptus, Oak, Hackberry, Maple, Fruit trees
	<i>Saperda calcarata</i>	Poplar
	<i>Strophiona nitens</i>	Chestnut, Oak, Hickory, Walnut, Beech, Maple
Scolytidae	<i>Corthylus columbianus</i>	Maple, Oak, Yellow-poplar, Beech, Boxelder, Sycamore, Birch, Basswood, Chestnut, Elm
	<i>Dendroctonus frontalis</i>	Pine
	<i>Dryocoetes betulae</i>	Birch, Sweetgum, Wild cherry, Beech, Pear
	<i>Monarthrum fasciatum</i>	Oak, Maple, Birch, Chestnut, Sweetgum, Blackgum, Poplar, Hickory, Mimosa, Apple, Peach, Pine
	<i>Phloeotribus liminaris</i>	Peach, Cherry, Plum, Black cherry, Elm, Mulberry, Mountain-ash
	<i>Pseudopityophthorus pruinosis</i>	Oak, American beech, Black cherry, Chickasaw plum, Chestnut, Maple, Hickory, Hornbeam, Hophornbeam
Sesiidae	<i>Paranthrene simulans</i>	Oak, American chestnut
	<i>Sannina uroceriformis</i>	Persimmon
	<i>Synanthedon exitiosa</i>	Peach, Plum, Nectarine, Cherry, Apricot, Almond, Black cherry
	<i>Synanthedon pictipes</i>	Peach, Plum, Cherry, Beach, Black Cherry
	<i>Synanthedon rubrofascia</i>	Tupelo
	<i>Synanthedon scitula</i>	Dogwood, Pecan, Hickory, Oak, Chestnut, Beech, Birch, Black cherry, Elm, Mountain-ash, Viburnum, Willow, Apple, Loquat, Ninebark, Bayberry
	<i>Vitacea polistiformis</i>	Grape

The present invention may be also used to control any insect pests that may be present in turfgrass, including for example beetles, caterpillars, fire ants, ground pearls, millipedes, sow bugs, mites, mole crickets, scales, mealybugs, ticks, spittlebugs, southern chinch bugs and white grubs. The present invention may be used to control insect pests at various stages of their life cycle, including eggs, larvae, nymphs and adults.

In particular, the present invention may be used to control insect pests that feed on the roots of turfgrass including white grubs (such as *Cyclocephala* spp. (e.g. masked chafer, *C. lurida*), *Rhizotrogus* spp. (e.g. European chafer, *R. majalis*), *Cotinus* spp. (e.g. Green June beetle, *C. nitida*), *Popillia* spp. (e.g. Japanese beetle, *P. japonica*), *Phyllophaga* spp. (e.g. May/June beetle), *Ataenius* spp. (e.g. Black turfgrass ataenius, *A. spretulus*), *Maladera* spp. (e.g. Asiatic garden beetle, *M. castanea*) and *Tomarus* spp.), ground pearls (*Margarodes* spp.), mole crickets (tawny, southern, and short-winged; *Scapteriscus* spp., *Gryllotalpa africana*) and leatherjackets (European crane fly, *Tipula* spp.).

The present invention may also be used to control insect pests of turfgrass that are thatch dwelling, including armyworms (such as fall armyworm *Spodoptera frugiperda*, and common armyworm *Pseudaletia unipuncta*), cutworms, billbugs (*Sphenophorus* spp., such as *S. venatus verstitus* and *S. parvulus*), and sod webworms (such as *Crambus* spp. and the tropical sod webworm, *Herpetogramma phaeopteralis*).

The present invention may also be used to control insect pests of turfgrass that live above the ground and feed on the turfgrass leaves, including chinch bugs (such as southern chinch bugs, *Blissus insularis*), Bermudagrass mite (*Eriophyes cynodoniensis*), rhodesgrass mealybug (*Antonina graminis*), two-lined spittlebug (*Prospapia bicincta*), leafhoppers, cutworms (*Noctuidae* family), and greenbugs.

The present invention may also be used to control other pests of turfgrass such as red imported fire ants (*Solenopsis invicta*) that create ant mounds in turf.

In the hygiene sector, the compositions according to the invention are active against ectoparasites such as hard ticks, soft ticks, mange mites, harvest mites, flies (biting and licking), parasitic fly larvae, lice, hair lice, bird lice and fleas.

Examples of such parasites are:

Of the order Anoplurida: *Haematopinus* spp., *Linognathus* spp., *Pediculus* spp. and *Phtirus* spp., *Solenopotes* spp..

Of the order Mallophagida: *Trimenopon* spp., *Menopon* spp., *Trinoton* spp., *Bovicola* spp., *Werneckiella* spp., *Lepikentron* spp., *Damalina* spp., *Trichodectes* spp. and *Felicola* spp..

Of the order Diptera and the suborders Nematocera and Brachycera, for example *Aedes* spp., *Anopheles* spp., *Culex* spp., *Simulium* spp., *Eusimulium* spp., *Phlebotomus* spp., *Lutzomyia* spp., *Culicoides* spp., *Chrysops* spp., *Hybomitra* spp., *Atylotus* spp., *Tabanus* spp., *Haematopota* spp.,
 5 *Philipomyia* spp., *Braula* spp., *Musca* spp., *Hydrotaea* spp., *Stomoxys* spp., *Haematobia* spp., *Morellia* spp., *Fannia* spp., *Glossina* spp., *Calliphora* spp., *Lucilia* spp., *Chrysomya* spp., *Wohlfahrtia* spp., *Sarcophaga* spp., *Oestrus* spp., *Hypoderma* spp., *Gasterophilus* spp., *Hippobosca* spp., *Lipoptena* spp. and *Melophagus* spp..

- 10 Of the order Siphonaptera, for example *Pulex* spp., *Ctenocephalides* spp., *Xenopsylla* spp., *Ceratophyllus* spp..

Of the order Heteroptera, for example *Cimex* spp., *Triatoma* spp., *Rhodnius* spp., *Panstrongylus* spp..

15

Of the order Blattellidae, for example *Blattella orientalis*, *Periplaneta americana*, *Blattella germanica* and *Supella* spp..

- 20 Of the subclass Acarina (Acarida) and the orders Meta- and Meso-stigmata, for example *Argas* spp., *Ornithodoros* spp., *Otobius* spp., *Ixodes* spp., *Amblyomma* spp., *Boophilus* spp., *Dermacentor* spp., *Haemophysalis* spp., *Hyalomma* spp., *Rhipicephalus* spp., *Dermanyssus* spp., *Raillietia* spp., *Pneumonyssus* spp., *Sternostoma* spp. and *Varroa* spp..

- 25 Of the orders Actinotrichida (Prostigmata) and Actinotrichida (Astigmata), for example *Acarapis* spp., *Cheyletiella* spp., *Ornithocheyletia* spp., *Myobia* spp., *Psorergates* spp., *Demodex* spp., *Trombicula* spp., *Listrophorus* spp., *Acarus* spp., *Tyrophagus* spp., *Caloglyphus* spp., *Hypodectes* spp., *Pterolichus* spp., *Psoroptes* spp., *Chorioptes* spp., *Otodectes* spp., *Sarcoptes* spp., *Notoedres* spp., *Knemidocoptes* spp., *Cytodites* spp. and *Laminosioptes* spp..

- 30 The compositions according to the invention are also suitable for protecting against insect infestation in the case of materials such as wood, textiles, plastics, adhesives, glues, paints, paper and card, leather, floor coverings and buildings.

- 35 The compositions according to the invention can be used, for example, against the following pests: beetles such as *Hylotrupes bajulus*, *Chlorophorus pilosus*, *Anobium punctatum*, *Xestobium rufovillosum*, *Ptilinuspecticornis*, *Dendrobium pertinax*, *Ernobius mollis*, *Priobium carpini*, *Lyctus brunneus*, *Lyctus africanus*, *Lyctus planicollis*, *Lyctus linearis*, *Lyctus pubescens*, *Trogosynon aequale*, *Minthesrugicollis*, *Xyleborus spec.*, *Tryptodendron spec.*, *Apate monachus*, *Bostrychus capucinus*, *Heterobostrychus brunneus*, *Sinoxylon spec.* and *Dinoderus minutus*, and also hymenoptera such

as *Sirex juvencus*, *Urocerus gigas*, *Urocerus gigas taignus* and *Urocerus augur*, and termites such as *Kaloterms flavicollis*, *Cryptoterms brevis*, *Heteroterms indicola*, *Reticulitermes flavipes*, *Reticulitermes santonensis*, *Reticulitermes lucifugus*, *Mastoterms darwiniensis*, *Zootermopsis nevadensis* and *Coptoterms formosanus*, and bristletails such as *Lepisma saccharina*. The

5 compounds of formulae I, and I'a, or salts thereof, are especially suitable for controlling one or more pests selected from the family: Noctuidae, Plutellidae, Chrysomelidae, Thripidae, Pentatomidae, Tortricidae, Delphacidae, Aphididae, Noctuidae, Crambidae, Meloidogynidae, and Heteroderidae. In a preferred embodiment of each aspect, a compound TX (where the abbreviation "TX" means "one compound selected from the compounds defined in the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27, and Table P") controls one or more of pests selected from the family: Noctuidae, Plutellidae, Chrysomelidae, Thripidae, Pentatomidae, Tortricidae, Delphacidae, Aphididae, Noctuidae, Crambidae, Meloidogynidae, and Heteroderidae.

15 The compounds of formulae I, and I'a, or salts thereof, are especially suitable for controlling one or more of pests selected from the genus: *Spodoptera spp*, *Plutella spp*, *Frankliniella spp*, *Thrips spp*, *Euschistus spp*, *Cydia spp*, *Nilaparvata spp*, *Myzus spp*, *Aphis spp*, *Diabrotica spp*, *Rhopalosiphum spp*, *Pseudoplusia spp* and *Chilo spp*. . In a preferred embodiment of each aspect, a compound TX (where the abbreviation "TX" means "one compound selected from the compounds defined in the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27, and Table P") controls one or more of pests selected from the genus: *Spodoptera spp*, *Plutella spp*, *Frankliniella spp*, *Thrips spp*, *Euschistus spp*, *Cydia spp*, *Nilaparvata spp*, *Myzus spp*, *Aphis spp*, *Diabrotica spp*, *Rhopalosiphum spp*, *Pseudoplusia spp* and *Chilo spp*.

25 The compounds of formulae I, and I'a, or salts thereof, are especially suitable for controlling one or more of *Spodoptera littoralis*, *Plutella xylostella*, *Frankliniella occidentalis*, *Thrips tabaci*, *Euschistus heros*, *Cydia pomonella*, *Nilaparvata lugens*, *Myzus persicae*, *Chrysodeixis includens*, *Aphis craccivora*, *Diabrotica balteata*, *Rhopalosiphum padi*, and *Chilo suppressalis*.

30 In a preferred embodiment of each aspect, a compound TX (where the abbreviation "TX" means "one compound selected from the compounds defined in the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27, and Table P") controls one or more of *Spodoptera littoralis*, *Plutella xylostella*, *Frankliniella occidentalis*, *Thrips tabaci*, *Euschistus heros*, *Cydia pomonella*, *Nilaparvata lugens*, *Myzus persicae*, *Chrysodeixis includens*, *Aphis craccivora*, *Diabrotica balteata*, *Rhopalosiphum Padi*, and *Chilo Suppressalis*, such as *Spodoptera littoralis* + TX, *Plutella xylostella* + TX; *Frankliniella occidentalis* + TX, *Thrips tabaci* + TX, *Euschistus heros* + TX, *Cydia pomonella* + TX, *Nilaparvata lugens* + TX, *Myzus persicae* + TX, *Chrysodeixis includens* + TX, *Aphis craccivora* + TX, *Diabrotica balteata* + TX, *Rhopalosiphum Padi* + TX, and *Chilo suppressalis* + TX.

In an embodiment, of each aspect, one compound selected from the compounds defined in the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27, and Table P, is suitable for controlling *Spodoptera littoralis*, *Plutella xylostella*, *Frankliniella occidentalis*, *Thrips tabaci*, *Euschistus heros*, *Cydia pomonella*, *Nilaparvata lugens*, *Myzus persicae*, *Chrysodeixis includens*, *Aphis craccivora*, *Diabrotica balteata*, *Rhopalosiphum Padia*, and *Chilo Suppressalis* in cotton, vegetable, maize, cereal, rice and soya crops.

In an embodiment, one compound from selected from the compounds defined in the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27, and Table P, is suitable for controlling Mamestra (preferably in vegetables), *Cydia pomonella* (preferably in apples), *Empoasca* (preferably in vegetables, vineyards), *Leptinotarsa* (preferably in potatoes) and *Chilo suppressalis* (preferably in rice).

Compounds according to the invention may possess any number of benefits including, inter alia, advantageous levels of biological activity for protecting plants against insects or superior properties for use as agrochemical active ingredients (for example, greater biological activity, an advantageous spectrum of activity, an increased safety profile (against non-target organisms above and below ground (such as fish, birds and bees), improved physico-chemical properties, or increased biodegradability). In particular, it has been surprisingly found that certain compounds of formula I may show an advantageous safety profile with respect to non-target arthropods, in particular pollinators such as honey bees, solitary bees, and bumble bees. Most particularly, *Apis mellifera*.

The compounds according to the invention can be used as pesticidal agents in unmodified form, but they are generally formulated into compositions in various ways using formulation adjuvants, such as carriers, solvents and surface-active substances. The formulations can be in various physical forms, e.g. in the form of dusting powders, gels, wettable powders, water-dispersible granules, water-dispersible tablets, effervescent pellets, emulsifiable concentrates, microemulsifiable concentrates, oil-in-water emulsions, oil-flowables, aqueous dispersions, oily dispersions, suspo-emulsions, capsule suspensions, emulsifiable granules, soluble liquids, water-soluble concentrates (with water or a water-miscible organic solvent as carrier), impregnated polymer films or in other forms known e.g. from the Manual on Development and Use of FAO and WHO Specifications for Pesticides, United Nations, First Edition, Second Revision (2010). Such formulations can either be used directly or diluted prior to use. The dilutions can be made, for example, with water, liquid fertilisers, micronutrients, biological organisms, oil or solvents.

The formulations can be prepared e.g. by mixing the active ingredient with the formulation adjuvants in order to obtain compositions in the form of finely divided solids, granules, solutions, dispersions or emulsions. The active ingredients can also be formulated with other adjuvants, such as finely divided solids, mineral oils, oils of vegetable or animal origin, modified oils of vegetable or animal origin, organic solvents, water, surface-active substances or combinations thereof.

The active ingredients can also be contained in very fine microcapsules. Microcapsules contain the active ingredients in a porous carrier. This enables the active ingredients to be released into the environment in controlled amounts (e.g. slow-release). Microcapsules usually have a diameter of from 0.1 to 500 microns. They contain active ingredients in an amount of about from 25 to 95 % by weight of the capsule weight. The active ingredients can be in the form of a monolithic solid, in the form of fine particles in solid or liquid dispersion or in the form of a suitable solution. The encapsulating membranes can comprise, for example, natural or synthetic rubbers, cellulose, styrene/butadiene copolymers, polyacrylonitrile, polyacrylate, polyesters, polyamides, polyureas, polyurethane or chemically modified polymers and starch xanthates or other polymers that are known to the person skilled in the art. Alternatively, very fine microcapsules can be formed in which the active ingredient is contained in the form of finely divided particles in a solid matrix of base substance, but the microcapsules are not themselves encapsulated.

The formulation adjuvants that are suitable for the preparation of the compositions according to the invention are known *per se*. As liquid carriers there may be used: water, toluene, xylene, petroleum ether, vegetable oils, acetone, methyl ethyl ketone, cyclohexanone, acid anhydrides, acetonitrile, acetophenone, amyl acetate, 2-butanone, butylene carbonate, chlorobenzene, cyclohexane, cyclohexanol, alkyl esters of acetic acid, diacetone alcohol, 1,2-dichloropropane, diethanolamine, p-diethylbenzene, diethylene glycol, diethylene glycol abietate, diethylene glycol butyl ether, diethylene glycol ethyl ether, diethylene glycol methyl ether, *N,N*-dimethylformamide, dimethyl sulfoxide, 1,4-dioxane, dipropylene glycol, dipropylene glycol methyl ether, dipropylene glycol dibenzoate, diproxitol, alkylpyrrolidone, ethyl acetate, 2-ethylhexanol, ethylene carbonate, 1,1,1-trichloroethane, 2-heptanone, alpha-pinene, d-limonene, ethyl lactate, ethylene glycol, ethylene glycol butyl ether, ethylene glycol methyl ether, gamma-butyrolactone, glycerol, glycerol acetate, glycerol diacetate, glycerol triacetate, hexadecane, hexylene glycol, isoamyl acetate, isobornyl acetate, isooctane, isophorone, isopropylbenzene, isopropyl myristate, lactic acid, laurylamine, mesityl oxide, methoxypropanol, methyl isoamyl ketone, methyl isobutyl ketone, methyl laurate, methyl octanoate, methyl oleate, methylene chloride, m-xylene, *n*-hexane, *n*-octylamine, octadecanoic acid, octylamine acetate, oleic acid, oleylamine, o-xylene, phenol, polyethylene glycol, propionic acid, propyl lactate, propylene carbonate, propylene glycol, propylene glycol methyl ether, p-xylene, toluene, triethyl phosphate, triethylene glycol, xylenesulfonic acid, paraffin, mineral oil, trichloroethylene, perchloroethylene, ethyl acetate, amyl acetate, butyl acetate, propylene glycol methyl ether, diethylene glycol methyl ether, methanol, ethanol, isopropanol, and alcohols of higher molecular weight, such as amyl alcohol, tetrahydrofurfuryl alcohol, hexanol, octanol, ethylene glycol, propylene glycol, glycerol, *N*-methyl-2-pyrrolidone and the like.

Suitable solid carriers are, for example, talc, titanium dioxide, pyrophyllite clay, silica, attapulgite clay, kieselguhr, limestone, calcium carbonate, bentonite, calcium montmorillonite, cottonseed husks, wheat flour, soybean flour, pumice, wood flour, ground walnut shells, lignin and similar substances.

- 5 A large number of surface-active substances can advantageously be used in both solid and liquid formulations, especially in those formulations which can be diluted with a carrier prior to use. Surface-active substances may be anionic, cationic, non-ionic or polymeric and they can be used as emulsifiers, wetting agents or suspending agents or for other purposes. Typical surface-active substances include, for example, salts of alkyl sulfates, such as diethanolammonium lauryl sulfate; salts of alkylarylsulfonates, such as calcium dodecylbenzenesulfonate; alkylphenol/alkylene oxide addition products, such as nonylphenol ethoxylate; alcohol/alkylene oxide addition products, such as tridecylalcohol ethoxylate; soaps, such as sodium stearate; salts of alkylnaphthalenesulfonates, such as sodium dibutylnaphthalenesulfonate; dialkyl esters of sulfosuccinate salts, such as sodium di(2-ethylhexyl)sulfosuccinate; sorbitol esters, such as sorbitol oleate; quaternary amines, such as lauryltrimethylammonium chloride, polyethylene glycol esters of fatty acids, such as polyethylene glycol stearate; block copolymers of ethylene oxide and propylene oxide; and salts of mono- and dialkylphosphate esters; and also further substances described e.g. in McCutcheon's Detergents and Emulsifiers Annual, MC Publishing Corp., Ridgewood New Jersey (1981).
- 10
- 15
- 20 Further adjuvants that can be used in pesticidal formulations include crystallisation inhibitors, viscosity modifiers, suspending agents, dyes, anti-oxidants, foaming agents, light absorbers, mixing auxiliaries, antifoams, complexing agents, neutralising or pH-modifying substances and buffers, corrosion inhibitors, fragrances, wetting agents, take-up enhancers, micronutrients, plasticisers, glidants, lubricants, dispersants, thickeners, antifreezes, microbicides, and liquid and solid fertilisers.
- 25
- The compositions according to the invention can include an additive comprising an oil of vegetable or animal origin, a mineral oil, alkyl esters of such oils or mixtures of such oils and oil derivatives. The amount of oil additive in the composition according to the invention is generally from 0.01 to 10 %, based on the mixture to be applied. For example, the oil additive can be added to a spray tank in the desired concentration after a spray mixture has been prepared. Preferred oil additives comprise mineral oils or an oil of vegetable origin, for example rapeseed oil, olive oil or sunflower oil, emulsified vegetable oil, alkyl esters of oils of vegetable origin, for example the methyl derivatives, or an oil of animal origin, such as fish oil or beef tallow. Preferred oil additives comprise alkyl esters of C₈-C₂₂ fatty acids, especially the methyl derivatives of C₁₂-C₁₈ fatty acids, for example the methyl esters of lauric acid, palmitic acid and oleic acid (methyl laurate, methyl palmitate and methyl oleate, respectively).
- 30
- 35 Many oil derivatives are known from the Compendium of Herbicide Adjuvants, 10th Edition, Southern Illinois University, 2010.

-67-

The inventive compositions generally comprise from 0.1 to 99 % by weight, especially from 0.1 to 95 % by weight, of compounds of the present invention and from 1 to 99.9 % by weight of a formulation adjuvant which preferably includes from 0 to 25 % by weight of a surface-active substance.

Whereas commercial products may preferably be formulated as concentrates, the end user will normally employ dilute formulations.

The rates of application vary within wide limits and depend on the nature of the soil, the method of application, the crop plant, the pest to be controlled, the prevailing climatic conditions, and other factors governed by the method of application, the time of application and the target crop. As a general guideline compounds may be applied at a rate of from 1 to 2000 l/ha, especially from 10 to 1000 l/ha.

Preferred formulations can have the following compositions (weight %):

Emulsifiable concentrates:

- active ingredient: 1 to 95 %, preferably 60 to 90 %
 surface-active agent: 1 to 30 %, preferably 5 to 20 %
 liquid carrier: 1 to 80 %, preferably 1 to 35 %

Dusts:

- active ingredient: 0.1 to 10 %, preferably 0.1 to 5 %
 solid carrier: 99.9 to 90 %, preferably 99.9 to 99 %

Suspension concentrates:

- active ingredient: 5 to 75 %, preferably 10 to 50 %
 water: 94 to 24 %, preferably 88 to 30 %
 surface-active agent: 1 to 40 %, preferably 2 to 30 %

Wettable powders:

- active ingredient: 0.5 to 90 %, preferably 1 to 80 %
 surface-active agent: 0.5 to 20 %, preferably 1 to 15 %
 solid carrier: 5 to 95 %, preferably 15 to 90 %

Granules:

- active ingredient: 0.1 to 30 %, preferably 0.1 to 15 %
 solid carrier: 99.5 to 70 %, preferably 97 to 85 %

The following Examples further illustrate, but do not limit, the invention.

<u>Wettable powders</u>	a)	b)	c)
-------------------------	----	----	----

-68-

active ingredients	25 %	50 %	75 %
sodium lignosulfonate	5 %	5 %	-
sodium lauryl sulfate	3 %	-	5 %
sodium diisobutylphenathalenesulfonate	-	6 %	10 %
phenol polyethylene glycol ether (7-8 mol of ethylene oxide)	-	2 %	-
highly dispersed silicic acid	5 %	10 %	10 %
Kaolin	62 %	27 %	-

The combination is thoroughly mixed with the adjuvants and the mixture is thoroughly ground in a suitable mill, affording wettable powders that can be diluted with water to give suspensions of the desired concentration.

<u>Powders for dry seed treatment</u>	a)	b)	c)
active ingredients	25 %	50 %	75 %
light mineral oil	5 %	5 %	5 %
highly dispersed silicic acid	5 %	5 %	-
Kaolin	65 %	40 %	-
Talcum	-		20 %

- 5 The combination is thoroughly mixed with the adjuvants and the mixture is thoroughly ground in a suitable mill, affording powders that can be used directly for seed treatment.

<u>Emulsifiable concentrate</u>	
active ingredients	10 %
octylphenol polyethylene glycol ether (4-5 mol of ethylene oxide)	3 %
calcium dodecylbenzenesulfonate	3 %
castor oil polyglycol ether (35 mol of ethylene oxide)	4 %
Cyclohexanone	30 %
xylene mixture	50 %

Emulsions of any required dilution, which can be used in plant protection, can be obtained from this concentrate by dilution with water.

-69-

<u>Dusts</u>	a)	b)	c)
Active ingredients	5 %	6 %	4 %
Talcum	95 %	-	-
Kaolin	-	94 %	-
mineral filler	-	-	96 %

Ready-for-use dusts are obtained by mixing the combination with the carrier and grinding the mixture in a suitable mill. Such powders can also be used for dry dressings for seed.

<u>Extruder granules</u>	
Active ingredients	15 %
sodium lignosulfonate	2 %
carboxymethylcellulose	1 %
Kaolin	82 %

The combination is mixed and ground with the adjuvants, and the mixture is moistened with water. The mixture is extruded and then dried in a stream of air.

<u>Coated granules</u>	
Active ingredients	8 %
polyethylene glycol (mol. wt. 200)	3 %
Kaolin	89 %

The finely ground combination is uniformly applied, in a mixer, to the kaolin moistened with polyethylene glycol. Non-dusty coated granules are obtained in this manner.

10 Suspension concentrate

active ingredients	40 %
propylene glycol	10 %
nonylphenol polyethylene glycol ether (15 mol of ethylene oxide)	6 %
Sodium lignosulfonate	10 %
carboxymethylcellulose	1 %
silicone oil (in the form of a 75 % emulsion in water)	1 %
Water	32 %

The finely ground combination is intimately mixed with the adjuvants, giving a suspension concentrate from which suspensions of any desired dilution can be obtained by dilution with water. Using such dilutions, living plants as well as plant propagation material can be treated and protected against infestation by microorganisms, by spraying, pouring or immersion.

15

Flowable concentrate for seed treatment

active ingredients	40 %
--------------------	------

-70-

propylene glycol	5 %
copolymer butanol PO/EO	2 %
Tristyrenephenole with 10-20 moles EO	2 %
1,2-benzisothiazolin-3-one (in the form of a 20% solution in water)	0.5 %
monoazo-pigment calcium salt	5 %
Silicone oil (in the form of a 75 % emulsion in water)	0.2 %
Water	45.3 %

The finely ground combination is intimately mixed with the adjuvants, giving a suspension concentrate from which suspensions of any desired dilution can be obtained by dilution with water. Using such dilutions, living plants as well as plant propagation material can be treated and protected against infestation by microorganisms, by spraying, pouring or immersion.

5

Slow Release Capsule Suspension

28 parts of the combination are mixed with 2 parts of an aromatic solvent and 7 parts of toluene diisocyanate/polymethylene-polyphenylisocyanate-mixture (8:1). This mixture is emulsified in a mixture of 1.2 parts of polyvinylalcohol, 0.05 parts of a defoamer and 51.6 parts of water until the desired particle size is achieved. To this emulsion a mixture of 2.8 parts 1,6-diaminohexane in 5.3 parts of water is added. The mixture is agitated until the polymerization reaction is completed. The obtained capsule suspension is stabilized by adding 0.25 parts of a thickener and 3 parts of a dispersing agent. The capsule suspension formulation contains 28% of the active ingredients. The medium capsule diameter is 8-15 microns. The resulting formulation is applied to seeds as an aqueous suspension in an apparatus suitable for that purpose.

Formulation types include an emulsion concentrate (EC), a suspension concentrate (SC), a suspo-emulsion (SE), a capsule suspension (CS), a water dispersible granule (WG), an emulsifiable granule (EG), an emulsion, water in oil (EO), an emulsion, oil in water (EW), a micro-emulsion (ME), an oil dispersion (OD), an oil miscible flowable (OF), an oil miscible liquid (OL), a soluble concentrate (SL), an ultra-low volume suspension (SU), an ultra-low volume liquid (UL), a technical concentrate (TK), a dispersible concentrate (DC), a wettable powder (WP), a soluble granule (SG) or any technically feasible formulation in combination with agriculturally acceptable adjuvants.

25 Preparatory Examples:

LCMS Methods:

Method 1:

Spectra were recorded on a Mass Spectrometer from Waters (SQD, SQDII Single quadrupole mass spectrometer) equipped with an electrospray source (Polarity: positive and negative ions, Capillary: 3.00 kV, Cone range: 30 V, Extractor: 2.00 V, Source Temperature: 150°C, Desolvation Temperature: 350°C, Cone Gas Flow: 50 l/h, Desolvation Gas Flow: 650 l/h, Mass range: 100 to 900 Da) and an Acquity UPLC from Waters: Binary pump, heated column compartment, diode-array

30

-71-

detector and ELSD detector. Column: Waters UPLC HSS T3, 1.8 μ m, 30 x 2.1 mm, Temp: 60 °C, DAD Wavelength range (nm): 210 to 500, Solvent Gradient: A = water + 5% MeOH + 0.05 % HCOOH, B= Acetonitrile + 0.05 % HCOOH, gradient: 10-100% B in 1.2 min; Flow (ml/min) 0.85

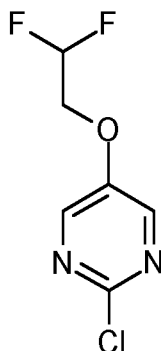
5 Method 2:

Spectra were recorded on a Mass Spectrometer from Waters (SQD, SQDII or ZQ Single quadrupole mass spectrometer) equipped with an electrospray source (Polarity: positive and negative ions), Capillary: 3.00 kV, Cone range: 30 V, Extractor: 2.00 V, Source Temperature: 150°C, Desolvation Temperature: 350°C, Cone Gas Flow: 50 l/h, Desolvation Gas Flow: 650 l/h, Mass range: 100 to 900 Da) and an Acquity UPLC from Waters: Binary pump, heated column compartment, diode-array detector and ELSD detector. Column: Waters UPLC HSS T3, 1.8 μ m, 30 x 2.1 mm, Temp: 60 °C, DAD Wavelength range (nm): 210 to 500, Solvent Gradient: A = water + 5% MeOH + 0.05 % HCOOH, B= Acetonitrile + 0.05 % HCOOH, gradient: 0-10% B in 2.5 min; Flow (ml/min) 0.85

- 15 Chiral SFC method 1: Spectra were recorded on a SFC from Waters (Waters Acquity UPC²/QDa) equipped with a PDA Detector Waters Acquity UPC². Column: Daicel SFC CHIRALPAK® IC, (3 μ m, 0.3cm x 10cm, 40 °C; Mobile phase: A: CO₂ B: MeOH isocratic: 10% B in 2.0 min; ABPR: 1800 psi; Flow rate: 2.0 ml/min; Detection: 220 nm; Sample concentration: 1 mg/mL in ACN; Injection: 1 μ L
- 20 Chiral SFC method 2: Spectra were recorded on a SFC from Waters (Waters Acquity UPC²/QDa) equipped with a PDA Detector Waters Acquity UPC². Column: Daicel SFC CHIRALPAK® IG, (3 μ m, 0.3cm x 10cm, 40 °C; Mobile phase: A: CO₂ B: MeOH isocratic: 15% B in 4.8 min; ABPR: 1800 psi; Flow rate: 2.0 ml/min; Detection: 270 nm; Sample concentration: 1 mg/mL in ACN/MeOH (1:1); Injection: 1 μ L

25

Preparation of 2-chloro-5-(2,2-difluoroethoxy)pyrimidine



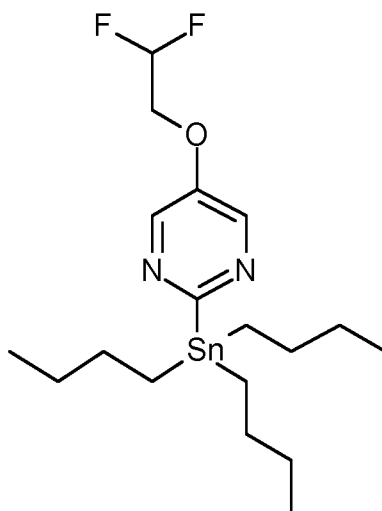
- 2-Chloropyrimidin-5-ol ([CAS: 4983-28-2] 5.00 g, 38.3 mmol) was dissolved in DMF (30.0 mL). Potassium carbonate (10.6 g, 76.6 mmol) and 1,1-difluoro-2-iodoethane (8.8 g, 46.0 mmol) were added.
- 30 The resulting reaction mixture was stirred at 80°C for 12 h before being cooled to room temperature. It was then poured into a mixture ice cold water and extracted twice with ethyl acetate (200 mL each). The combined organic layers were dried over sodium sulfate, filtered and concentrated in *vacuo*. The crude

-72-

material was purified by flash chromatography over silica gel (eluting with ethyl acetate in hexanes) to afford 2-chloro-5-(2,2-difluoroethoxy)pyrimidine.

LC-MS (method 1): m/z 195.1 $[M+H]^+$.

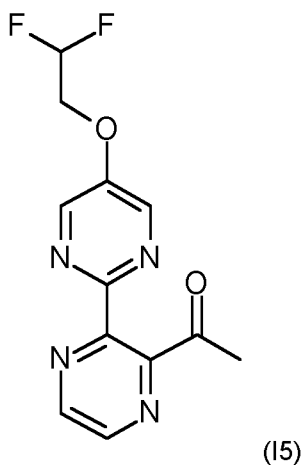
5 Preparation of tributyl-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]stannane



2-Chloro-5-(2,2-difluoroethoxy)pyrimidine (2.00 g, 9.251 mmol) was dissolved in toluene (40.0 mL) and hexa-n-butylditin (8.05g, 13.877 mmol) was added. The reaction mixture was purged with argon for 5 minutes, tetrakis(triphenylphosphine)palladium(0) (0.5345 g, 0.4626 mmol) was added, the reaction mixture was purged with argon for another 5 minutes and subsequently stirred at 100°C for 16 h. It was filtered through a pad of Celite and the filtrate was concentrated in *vacuo*. The crude material was purified by neutral alumina chromatography (eluting with ethyl acetate in hexanes) to afford tributyl-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]stannane.

^1H NMR (400 MHz, chloroform- d) δ ppm: 0.87 (t, 9 H) 1.15 (t, 6 H) 1.30 – 1.35 (q, 6 H) 1.54 – 1.62 (m, 6 H) 4.21 – 4.30 (dt, 2 H) 5.95 – 6.26 (br tt, 1 H) 8.47 (s, 2 H)

Preparation of 1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone (15)

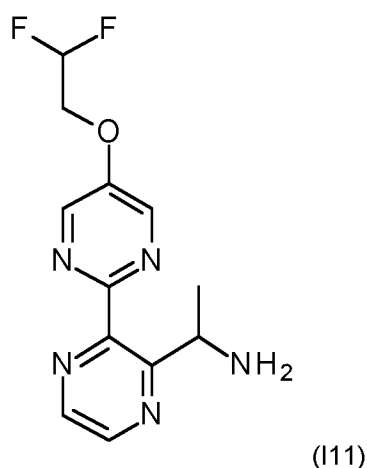


-73-

Tributyl-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]stannane (1.00 g, 2.00 mmol) was dissolved in toluene (15 mL), then 1-(3-chloropyrazin-2-yl)ethanone ([CAS: 121246-90-0] 0.439 g, 2.52 mmol) was added. The mixture was purged with argon for 5 minutes, then copper(I) iodide (0.0763 g, 0.401 mmol) and tetrakis(triphenylphosphine)palladium(0) (0.232 g, 0.200 mmol) were added and the resulting reaction mixture was stirred at 100°C for 4h. After cooling to room temperature, it was filtered through a pad of Celite and the filtrate was concentrated in *vacuo*. The crude product was purified by flash chromatography (silica gel, 0 – 100% ethyl acetate in hexanes) to afford 1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone.

LC-MS (method 1): m/z 281.1 $[M+H]^+$; ^1H NMR (400 MHz, DMSO- d_6) δ ppm: 8.90 (s, 1H), 8.70-8.90 (m, 3H), 6.48 (t, 1H), 4.63 (td, 2H), 2.62 (s, 3H).

Preparation of 1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine (I11)

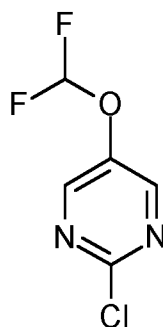


1-[3-[5-(2,2-Difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone (0.80 g, 0.285 mmol) was dissolved in a saturated solution of ammonium acetate in ethanol (10 mL). Ammonia solution 30% in water (5.0 mL) and sodium cyanoborohydride (0.0538 g, 0.856 mmol) were added and the reaction mixture was stirred at reflux for 18h. The reaction mixture was cooled to room temperature and washed with dichloromethane (50mL). The aqueous layer was concentrated in *vacuo* to afford a crude product which was purified by reverse phase chromatography (C18 column, 0 to 50% acetonitrile in water) to afford 1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine.

LC-MS (method 1): m/z 282.1 $[M+H]^+$; ^1H NMR (400 MHz, DMSO- d_6) δ ppm: 8.80-9.00 (m, 4H), 6.50 (tt, 1H), 4.90 (m, 1H), 4.78 (td, 2H), 1.45 (d, 3H)

Preparation of 2-chloro-5-(difluoromethoxy)pyrimidine

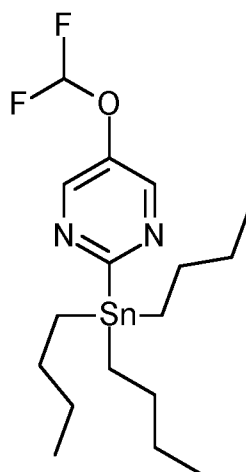
-74-



2-Chloropyrimidin-5-ol ([CAS: 4983-28-2] 5.00 g, 38.3 mmol) was dissolved in DMF (20.0 mL). Potassium carbonate (10.6 g, 76.6 mmol) and sodium 2-chloro-2,2-difluoroacetate (8.76 g, 57.5 mmol) were added. The resulting reaction mixture was stirred at 80°C for 4 h before being cooled to room temperature and diluted with ethyl acetate. This organic layer was washed twice with cold water (100 mL each), dried over sodium sulfate, filtered and concentrated in *vacuo*. The crude material was purified by flash chromatography over silica gel (eluting with ethyl acetate in hexanes) to afford 2-chloro-5-(difluoromethoxy)pyrimidine.

¹H NMR (400 MHz, chloroform-d) δ ppm: 6.45 – 6.82 (br t, 1 H) 8.55 (s, 2 H)

Preparation of tributyl-[5-(difluoromethoxy)pyrimidin-2-yl]stannane



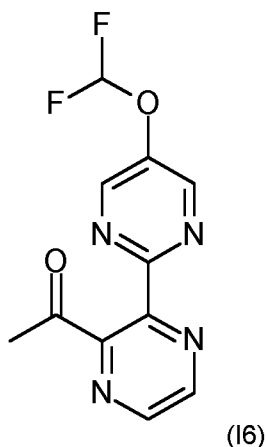
To a solution of 2-chloro-5-(difluoromethoxy)pyrimidine (2.70 g, 13.5 mmol) in toluene (50 mL) was added bis(tributyltin) (10.2 mL, 20.2 mmol). The reaction mixture was purged with argon for 5 minutes then tetrakis(triphenylphosphine)palladium(0) (778 mg, 0.673 mmol) was added and the reaction mixture was purged again with argon for another 2 minutes. The resulting reaction mixture was stirred at 100 °C for 16h. The reaction mixture was cooled to 0°C, diluted with water and extracted twice with ethyl acetate. The combined organic layers were dried over sodium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in n-hexane) afforded tributyl-[5-(difluoromethoxy)pyrimidin-2-yl]stannane.

¹H-NMR (400 MHz, CDCl₃): δ ppm: 8.60 (s, 1H), 7.26 (s, 1H), 6.57 (t, 1H), 1.50-1.70 (m, 6H), 1.25-1.40

-75-

(m, 6H), 1.10-1.20 (m, 6H), 0.88 (m, 9H).

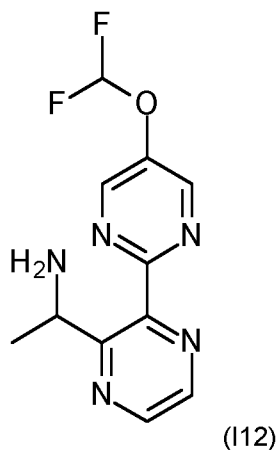
Preparation of 1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone (I6)



To a mixture of tributyl-[5-(difluoromethoxy)pyrimidin-2-yl]stannane (3.20 g, 6.62 mmol) in toluene (50.0 mL) was added 1-(4-chloropyrimidin-5-yl)ethanone (1267 mg, 7.28 mmol). The reaction mixture was purged with argon for 5 minutes. Tetrakis(triphenylphosphine)palladium(0) (382 mg, 0.331 mmol) and copper iodide (252 mg, 1.32 mmol) were added to the reaction mixture and purged again with argon for another 2 minutes. The resulting reaction mixture was stirred at 100 °C for 16 h. The reaction mixture was cooled to 0°C, diluted with water (100 mL) and extracted with ethyl acetate (2 x 100 mL). The combined organic layers were dried over sodium sulfate and concentrated *in vacuo*. The crude material was purified by flash chromatography over silica gel (gradient of ethyl acetate in hexanes) to afford 1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone as a light brown solid.

¹H NMR (400 MHz, DMSO-d₆) δ ppm: 8.80-9.00 (m, 4H), 7.50 (t, 1H), 2.65 (s, 3H)

Preparation of 1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine (I12)



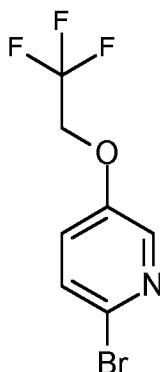
To a solution of 1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone (1.70 g, 6.39 mmol) in a solution of saturated ammonium acetate in ethanol (130 mL) were added sodium cyanoborohydride

-76-

(1.19 g, 19.2 mmol) and 30% aqueous ammonia (50 mL). The mixture was stirred at reflux for 16 h, cooled to room temperature, and concentrated in vacuo. The crude material was purified by reverse phase chromatography (C18 column, gradient of acetonitrile in water) to give 1-[3-5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine as a light brown gum.

5 ^1H NMR (400 MHz, DMSO- d_6) δ ppm: 8.80-9.10 (m, 4H), 7.51 (t, 1H), 4.88 (m, 1H), 1.50 (d, 3H)

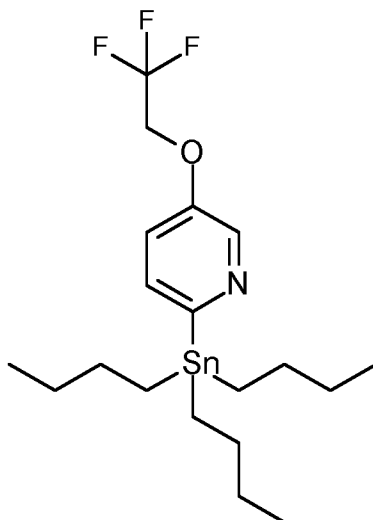
Preparation of 2-bromo-5-(2,2,2-trifluoroethoxy)pyridine



To a solution of 6-bromopyridin-3-ol (20.0 g, 115 mmol) and potassium carbonate (31.8 g, 230 mmol) in acetonitrile (200 mL), stirred at room temperature for 5 min, was added 2,2,2-Trifluoroethyl trifluoromethanesulfonate (29.3 g, 126 mmol). The reaction mixture was stirred at room temperature for 16 h. The reaction mixture was poured in ice cold water and extracted with ethyl acetate. The combined organic layers were washed with brine, dried over sodium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in n-hexane) afforded 2-bromo-5-(2,2,2-trifluoroethoxy)pyridine.

^1H -NMR (400 MHz, CDCl_3): δ ppm: 8.15 (d, 1H), 7.45 (d, 1H), 7.2 (dd, 1H), 4.4 (q, 2H).

Preparation of tributyl-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]stannane

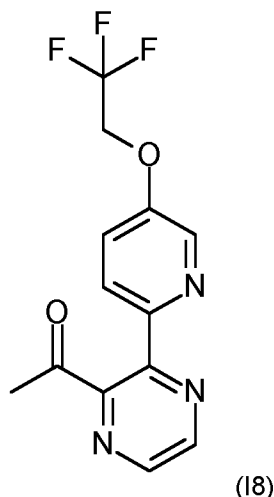


-77-

To a solution of 2-bromo-5-(2,2,2-trifluoroethoxy)pyridine (9.00 g, 31.6 mmol) in toluene (300 mL) was added bis(tributyltin) (20.7 mL, 41.1 mmol). The reaction mixture was purged with argon for 20 minutes then tetrakis(triphenylphosphine)palladium(0) (2.74 g, 2.37 mmol) was added and the reaction mixture was purged again with argon for another 2 minutes. The resulting reaction mixture was stirred at 100 °C for 48h. The reaction mixture was cooled to 0°C, diluted with water and extracted with ethyl acetate. Combined organic layers were dried over sodium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in n-hexane) afforded tributyl-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]stannane.

¹H-NMR (400 MHz, CDCl₃): δ ppm: 8.55 (d, 1H), 7.4 (dd, 1H), 7.15 (m, 1H), 4.4 (q, 2H), 1.55 (m, 6H), 1.35 (m, 6H), 1.15 (m, 6H), 0.95 (m, 9H).

Preparation of 1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethanone (18)

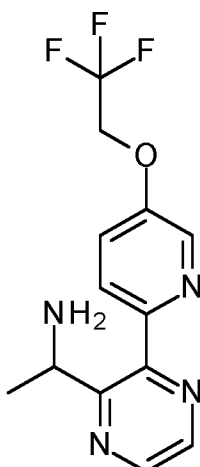


To a solution of tributyl-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]stannane (550 mg, 1.06 mmol) in toluene (20mL) were added 1-(3-chloropyrazin-2-yl)ethanone (203 mg, 1.17 mmol) and copper(I) iodide (40.4 mg, 0.212 mmol). The reaction mixture was purged with argon for 10 min and tetrakis(triphenylphosphine)palladium(0) (61.4 mg, 0.0531 mmol) was added. The reaction was stirred at 100°C for 2h. The reaction mixture was cooled to 0°C, diluted with water and extracted with ethyl acetate. Organic phase was dried over sodium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in n-hexane) afforded 1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethanone.

¹H-NMR (400 MHz, DMSO-d₆): δ ppm: 8.85 (d, 1H), 8.7 (m, 1H), 8.45 (s, 1H), 8.25 (d, 1H), 7.75 (d, 1H), 5 (q, 2H), 2.6 (s, 3H).

Preparation of 1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethanamine (113)

-78-

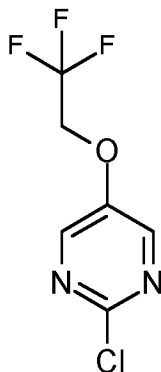


(I13)

To a solution of 1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethanone (1.80 g, 5.45 mmol) in a saturated solution of ammonium acetate in ethanol (120 mL) were added at room temperature sodium cyanoborohydride (1.01 g, 16.4 mmol) and ammonia (30% in water, 50 mL). The reaction mixture was stirred at reflux for 18 hours. After cooling down to room temperature, it was concentrated under reduced pressure. Purification of the crude material by reverse-phase chromatography (C18 column, eluting acetonitrile in water) afforded 1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethanamine.

¹H-NMR (400 MHz, DMSO-d₆): δ ppm: 8.8 (s, 2H), 8.65 (d, 1H), 8.15 (d, 1H), 7.8 (m, 1H), 7.45 (br s, 2H), 7.25 (m, 1H), 7.15 (m, 1H), 5.2 (br s, 1H), 5 (q, 2H), 1.5 (m, 3H).

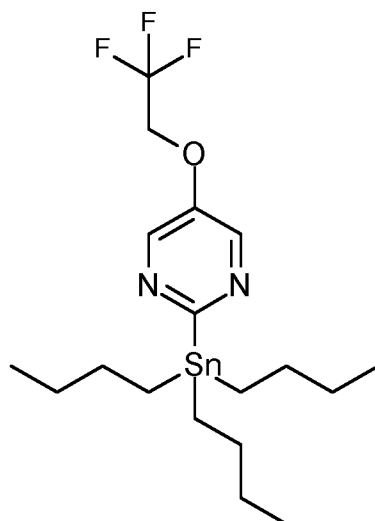
Preparation of 2-chloro-5-(2,2,2-trifluoroethoxy)pyrimidine



To a solution of 2-chloropyrimidin-5-ol (1.0 g, 7.7 mmol) in N,N-dimethylformamide (7.7 mL), was added cesium carbonate (3.2 g, 10 mmol). 2,2,2-trifluoroethyl trifluoromethanesulfonate (2.2 g, 9.2 mmol) was added dropwise to the mixture. The mixture was stirred at room temperature for 18 hours. The reaction mixture was poured into ice-water then extracted with ethyl acetate. The combined organic layers were washed four times with water, then with brine. They were dried over sodium sulfate and concentrated in vacuo. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in n-hexane) afforded 2-chloro-5-(2,2,2-trifluoroethoxy)pyrimidine as a light yellow oil.

¹H NMR (400 MHz, Chloroform) δ ppm: 4.46 - 4.52 (m, 2 H) 8.40 (s, 2 H); ¹⁹F NMR (377 MHz, Chloroform) δ ppm: -73.74 (s)

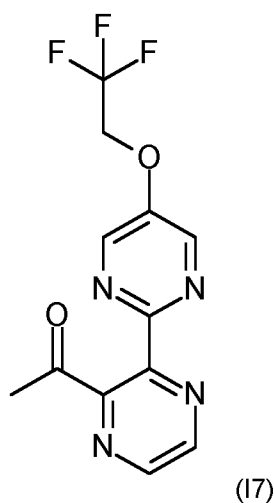
-79-

Preparation of tributyl-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]stannane

To a solution of 2-chloro-5-(2,2,2-trifluoroethoxy)pyrimidine (1.00 g, 4.70 mmol) in toluene (10 mL) was added bis(tributyltin) (4.09 g, 7.06 mmol). The reaction mixture was purged with argon for 5 minutes then tetrakis(triphenylphosphine)palladium(0) (0.544 g, 0.470 mmol) was added and the reaction mixture was purged again with argon for another 5 minutes. The resulting reaction mixture was stirred at 100 °C for 16 hours. The reaction mixture was filtered over Celite and the filtrate was concentrated under reduced pressure. Purification of the crude material by flash chromatography over neutral alumina (eluting with ethyl acetate in n-hexane) afforded tributyl-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]stannane.

¹H-NMR (400 MHz, CDCl₃): δ ppm: 8.50 (s, 1H), 8.40 (s, 1H), 4.43 (m, 2H), 4.12 (q, 2H), 1.50-1.70 (m, 6H), 1.20-1.45 (m, 6H), 1.10-1.20 (m, 6H), 0.83-0.98 (m, 9H).

Preparation of 1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone (17)



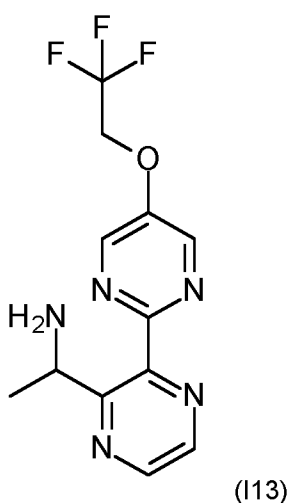
To a solution of tributyl-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]stannane (1.00 g, 2.14 mmol) in toluene (20

-80-

mL) were added 1-(3-chloropyrazin-2-yl)ethanone (0.402 g, 2.57 mmol). The reaction mixture was purged with argon for 5 min then copper(I) iodide (0.0815 g, 0.428 mmol) and [1,1'-bis(diphenylphosphino)ferrocene] dichloropalladium (II) (1.57 g, 2.14 mmol) were added. The reaction was stirred at 100°C for 4 hours. The reaction mixture was cooled to 0°C, filtered through celite and the filtrate was concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in n-hexane) afforded 1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone.

$m/z = 299.1$ $[M+H]^+$

10 Preparation of 1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine (I13)

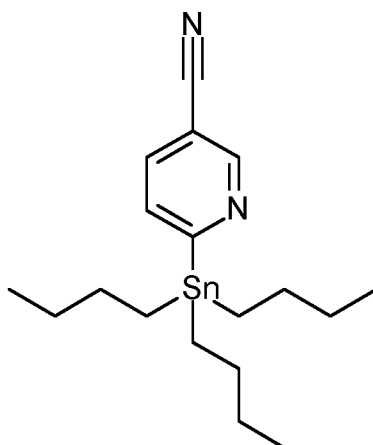


To a solution of 1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone (700 mg, 1.88 mmol) in a saturated solution of ammonium acetate in ethanol (100 mL) were added at room temperature sodium cyanoborohydride (0.354 g, 5.63 mmol) and ammonia (30% in water, 30 mL). The reaction mixture was stirred at reflux for 18 hours. After cooling down to room temperature, the mixture was washed with dichloromethane. The aqueous layer was concentrated in vacuo. Purification of the crude material by reverse-phase chromatography (C18 column, eluting acetonitrile in water) afforded 1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine.

$^1\text{H-NMR}$ (400 MHz, DMSO- d_6): δ ppm: 8.95 (s, 2H), 8.60-9.00 (m, 2H), 7.8-8.30 (br s, 2H), 5.08-5.20 (m, 2H), 4.95-5.05 (m, 1H), 1.5 (m, 3H).

Preparation of 6-tributylstannylpyridine-3-carbonitrile

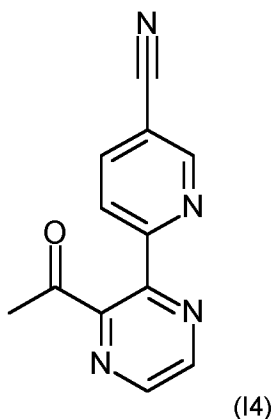
-81-



To a solution of 6-chloropyridine-3-carbonitrile (250 mg, 1.80 mmol) in toluene (10 mL) was added hexa-
n-butylditin (1.00 mL, 1.98 mmol). The reaction mixture was purged with argon for 2 minutes. Then
tetrakis(triphenylphosphine)palladium(0) (146 mg, 0.126 mmol) was added and it was purged again with
5 argon for additional 2 minutes. The resulting reaction mixture was heated up to 130 °C and stirred for
16 hours. After cooling down to room temperature, the reaction mixture was filtered through Celite. The
filtrate was concentrated under reduced pressure. Purification of the crude material by neutral alumina
flash chromatography (eluting with ethyl acetate in hexane) afforded 6-tributylstannylpyridine-3-
carbonitrile.

10 ¹H-NMR (400 MHz, CDCl₃): δ ppm: 8.90-9.00 (m, 1H), 7.65-7.75 (m, 1H), 7.50-7.60 (m, 1H), 7.25-7.40
(m, 2H), 1.45-1.65 (m, 4H), 1.25-1.40 (m, 7H), 1.10-1.20 (m, 5H), 0.80-0.95 (m, 9H).

Preparation of 6-(3-acetylpyrazin-2-yl)pyridine-3-carbonitrile (I4)



15 To a solution of 6-tributylstannylpyridine-3-carbonitrile (24.0 g, 48.8 mmol) in toluene (600 mL) were
added 1-(4-chloropyrimidin-5-yl)ethanone (9.18 g, 52.7 mmol) and copper(I) iodide (1.86 g, 9.77 mmol).
The reaction mixture was purged with argon for 10 minutes. Then
tetrakis(triphenylphosphine)palladium(0) (2.82 g, 2.44 mmol) was added. The reaction mixture was
heated up to 95 °C and stirred for 5 hours. After cooling down to room temperature, it was filtered through
20 Celite and the filtrate was concentrated under reduced pressure. Purification of the crude material by

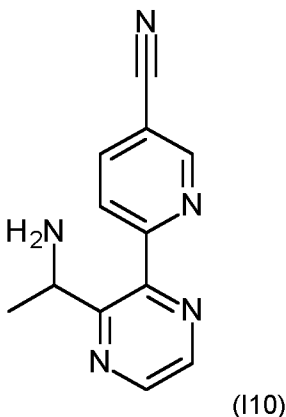
-82-

flash chromatography over silica gel (eluting with ethyl acetate in hexane) afforded 6-(3-acetylpyrazin-2-yl)pyridine-3-carbonitrile.

$^1\text{H-NMR}$ (400 MHz, $\text{d}_6\text{-DMSO}$): δ ppm: 9.1 (m, 2H), 8.92 (m, 1H), 8.83 (m, 1H) 8.53 (d, 1H) 8.32 (d, 1H) 2.65 (s, 3H).

5

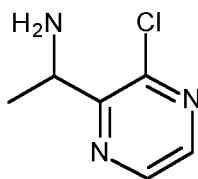
Preparation of 6-[3-(1-aminoethyl)pyrazin-2-yl]pyridine-3-carbonitrile (I10)



To a solution of 6-(3-acetylpyrazin-2-yl)pyridine-3-carbonitrile (0.200 g, 0.803 mmol) in a saturated solution of ammonium acetate in ethanol (30 mL) were added at room temperature aqueous ammonia (20 mL) and sodium cyanoborohydride (154 mg, 2.41 mmol). The reaction mixture was heated up to reflux and stirred for 12 hours. After cooling down to room temperature, it was concentrated under reduced pressure. Purification of the crude material by reverse-phase chromatography (eluting acetonitrile in water) afforded 6-[3-(1-aminoethyl)pyrazin-2-yl]pyridine-3-carbonitrile.

$^1\text{H-NMR}$ (400 MHz, $\text{d}_6\text{-DMSO}$): δ ppm: 9.22 (s, 1H), 8.85-8.95 (m, 2H), 8.50-8.60 (m, 1H), 8.30-8.40 (m, 1H) 7.80-8.10 (br. s, 2H), 5.25-5.35 (m, 1 H), 1.52 (d, 3H).

Preparation of 1-(3-chloropyrazin-2-yl)ethanamine



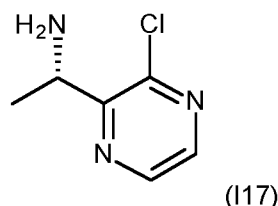
To a of 1-(3-chloropyrazin-2-yl)ethanone (0.200 g, 1.28 mmol) in methanol (4.5 mL) were added at room temperature ammonium acetate (0.995 g, 12.8 mmol) and sodium cyanoborohydride (0.0591 g, 0.894 mmol). The resulting suspension was stirred at room temperature for 18 hours, then concentrated in vacuo. The crude material was purified by reverse phase chromatography (C18 column, gradient of acetonitrile in water) to afford 1-(3-chloropyrazin-2-yl)ethanamine.

$^1\text{H NMR}$ (400 MHz, CDCl_3) δ ppm: 8.49 (d, 1H), 8.26 (d, 1H), 4.56 (q, 1H), 1.95 (br s, 2H), 1.44 (d, 3H)

Preparation of (1S)-1-(3-chloropyrazin-2-yl)ethanamine (I17)

25

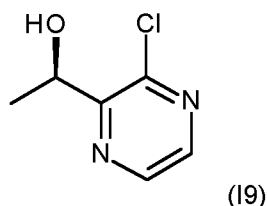
-83-



To a solution of 1-(3-chloropyrazin-2-yl)ethanamine (202.2 mg, 1.20 mmol) in tert-butyl methyl ether (11 mL) was added Novozym® 435 (240 mg), followed by ethyl methoxyacetate (1.44 mL, 12.0 mmol) at room temperature. The mixture was stirred at 40 °C for 5.5 hours. The reaction mixture was diluted with dichloromethane and filtered. The filtrate was concentrated in vacuo. The crude material was purified by flash chromatography over silica gel (eluting with a gradient of methanol in dichloromethane) to afford (1S)-1-(3-chloropyrazin-2-yl)ethanamine.

¹H NMR (400 MHz, CDCl₃) δ ppm: 8.49 (d, 1H), 8.27 (d, 1H), 4.56 (q, 1H), 1.73 (br s, 2H), 1.44 (d, 3H) ppm; [α]_D²⁰: -32.3° (c: 1.157, CHCl₃)

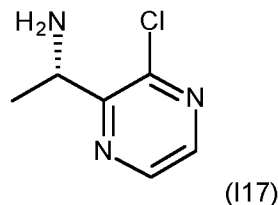
Preparation of (1R)-1-(3-chloropyrazin-2-yl)ethanol (I9)



1-(3-chloropyrazin-2-yl)ethanone (157 mg, 1.00 mmol) was dissolved in dichloromethane (10.0 mL) and the flask was evacuated and backfilled with argon three times. Then RuBF₄[(*R,R*)-TsDPEN](*p*-cymene) (0.0362 g, 0.0526 mmol) was added. A cooled solution of triethylamine (0.348 mL, 2.50 mmol) and formic acid (0.160 mL, 4.29 mmol) was added dropwise to the reaction mixture, which was stirred at room temperature for 4 hours. The reaction mixture was concentrated in vacuo. The crude material was purified by flash chromatography over silica gel (eluting with a gradient of ethyl acetate in cyclohexane) to afford (1R)-1-(3-chloropyrazin-2-yl)ethanol.

¹H-NMR (400 MHz, CDCl₃) δ ppm: 8.49 (d, 1H), 8.34 (d, 1H), 5.18 (m, 1H), 3.81 (d, 1H), 1.52 (d, 3H); Chiral SFC (method 2): 1.98 min (minor enantiomer), 2.55 min (major enantiomer); ee = 85%

Preparation of (1S)-1-(3-chloropyrazin-2-yl)ethanamine (I17)



(1R)-1-(3-chloropyrazin-2-yl)ethanol (87.8 mg, 0.554 mmol) was dissolved in tetrahydrofuran (1.9 mL). Then, 1,8-diazabicyclo[5.4.0]undec-7-ene (0.10 mL, 0.66 mmol) was added dropwise to the reaction mixture followed by diphenylphosphine azide (0.130 mL, 0.585 mmol). The reaction mixture was stirred at rt for 19 hours.

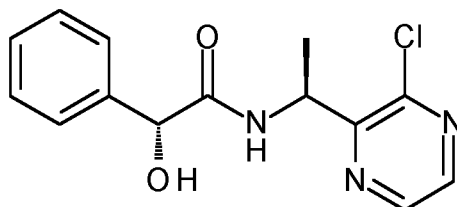
-84-

Tetrahydrofuran (1.4 mL) was added, followed by triphenylphosphine (179.4 mg, 0.677 mmol). The reaction mixture was stirred at room temperature for 2 hours. Water (0.15 mL) was added, and the reaction mixture was stirred at room temperature for 46 hours.

The reaction mixture was concentrated to a volume of 1 mL then diluted with dichloromethane. 1M hydrochloric acid was added, then the aqueous layer was washed with dichloromethane. The aqueous layer was basified to pH = 14 with 4 M sodium hydroxide solution and extracted with dichloromethane. The combined organic layers were dried over magnesium sulfate and concentrated in vacuo. The crude material was purified by flash chromatography over silica gel (eluting with a gradient of methanol in dichloromethane) to afford (1S)-1-(3-chloropyrazin-2-yl)ethanamine.

¹H NMR (400 MHz, CDCl₃) δ ppm: 8.49 (d, 1H), 8.27 (d, 1H), 4.56 (q, 1H), 1.84 (s, 2H), 1.44 (d, 3H); [α]_D²⁰: -26.0° (c: 0.960, CHCl₃)

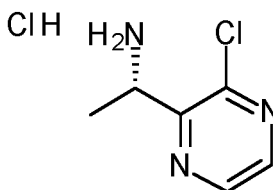
Preparation of (2R)-N-[(1S)-1-(3-chloropyrazin-2-yl)ethyl]-2-hydroxy-2-phenyl-acetamide



To a solution of 1-(3-chloropyrazin-2-yl)ethanamine;hydrochloride (700 mg, 3.61 mmol) in dichloromethane (18 mL) were added (R)-(-)-mandelic acid (610 mg, 3.97 mmol), N-ethyl-diisopropylamine (1.26 mL, 7.21 mmol), 1-hydroxybenzotriazole (50.8 mg, 0.361 mmol) and N,N'-dicyclohexylcarbodiimide (844 mg, 3.97 mmol). The reaction mixture was stirred at room temperature for 18 hours. The reaction mixture was diluted with saturated aqueous sodium carbonate solution and extracted with dichloromethane. The organic layers were dried over magnesium sulfate and concentrated in vacuo. Purification of the crude material by flash chromatography over silica gel (eluting with methanol in dichloromethane) afforded (2R)-N-[(1R)-1-(3-chloropyrazin-2-yl)ethyl]-2-hydroxy-2-phenyl-acetamide and (2R)-N-[(1S)-1-(3-chloropyrazin-2-yl)ethyl]-2-hydroxy-2-phenyl-acetamide. The relative stereochemistry of (2R)-N-[(1S)-1-(3-chloropyrazin-2-yl)ethyl]-2-hydroxy-2-phenyl-acetamide was determined by X-ray crystallography (crystallized from acetonitrile/water).

LCMS: Rt 0.74, m/z = 291 (M+H⁺)

Preparation of (1S)-1-(3-chloropyrazin-2-yl)ethanamine;hydrochloride



A solution of (2R)-N-[(1S)-1-(3-chloropyrazin-2-yl)ethyl]-2-hydroxy-2-phenyl-acetamide (0.93 g, 3.2 mmol) in hydrochloric acid (32% in water, 13 mL) was heated up to reflux and stirred for 2 hours. After

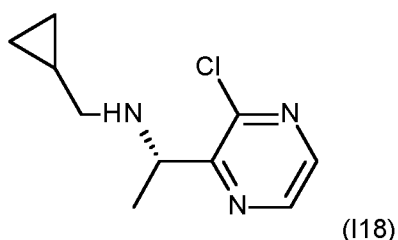
-85-

cooling down to room temperature, the reaction mixture was basified with 3 N sodium hydroxide and diluted and extracted with ethyl acetate. The aqueous layer was freeze-dried overnight and the resulting solid was suspended in acetone. The suspension was filtered and the filtrate was concentrated under reduced pressure. The resulting oil was dissolved in ethyl acetate and 1 N hydrochloric acid was added.

5 A precipitate appeared, it was filtered and dried under reduced pressure to afford the desired product.

LCMS: Rt 0.19, m/z = 158 (M+H+).

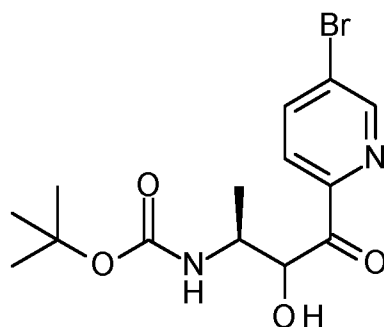
Preparation of (1S)-1-(3-chloropyrazin-2-yl)-N-(cyclopropylmethyl)ethanamine (I18)



Sodium triacetoxyborohydride (59.4 mg, 0.267 mmol) was added to a stirred solution of (1S)-1-(3-chloropyrazin-2-yl)ethanamine (30.0 mg, 0.190 mmol), cyclopropanecarboxylaldehyde (15.0 mg, 0.209 mmol) and acetic acid (0.0109 mL, 0.190 mmol) in 1,2-dichloroethane (0.95 mL). The mixture was stirred at room temperature for 4 hours. Saturated aqueous sodium carbonate solution was added, the aqueous layer was extracted with dichloromethane. The organic layer was dried over magnesium sulfate

15 and concentrated in vacuo. The crude material was purified by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) to afford (1S)-1-(3-chloropyrazin-2-yl)-N-(cyclopropylmethyl)ethanamine. ¹H NMR (400 MHz, Solvent) δ ppm: -0.03 - 0.10 (m, 2 H) 0.38 - 0.52 (m, 2 H) 0.83 - 1.00 (m, 1 H) 1.40 (d, 3 H) 2.07 (dd, 1 H) 2.15 - 2.29 (m, 1 H) 2.53 (dd, 1 H) 4.39 (q, 1 H) 8.26 (d, 1 H) 8.51 (d, 1 H); $[\alpha]_D^{20} = -54^\circ$ (c 0.327, CHCl₃)

Preparation of tert-butyl N-[(1S)-3-(5-bromo-2-pyridyl)-2-hydroxy-1-methyl-3-oxo-propyl]carbamate



In a round-bottomed flask was prepared a solution of tert-butyl N-[(1S)-1-methyl-2-oxo-ethyl]carbamate (CAS 79069-50-4, 1.07 g, 6.18 mmol) in dichloromethane (12 mL). The flask was evacuated and refilled

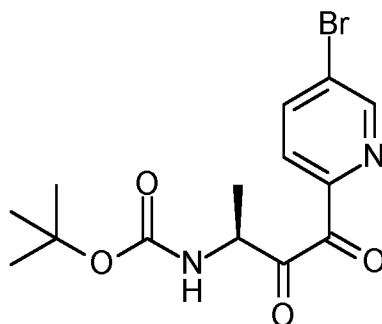
25 with argon three times. Then, 2-(3-benzyl-4-methyl-thiazol-3-ium-5-yl)ethanol;bromide (0.388 g, 1.24 mmol), 5-bromopyridine-2-carbaldehyde (CAS 31181-90-5, 1.81 g, 9.27 mmol) and dichloromethane (6 mL) were added successively, followed by N,N-diisopropylethylamine (2.16 mL, 12.4 mmol). The reaction mixture was stirred for 1 hour at room temperature. It was quenched with ammonium chloride

-86-

sat. aq. and extracted three times with dichloromethane. The combined organic layers were dried over magnesium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded tert-butyl N-[(1S)-3-(5-bromo-2-pyridyl)-2-hydroxy-1-methyl-3-oxo-propyl]carbamate as an orange gum.

- 5 LCMS: Rt 0.98, m/z = 359-361 (M+H⁺) (Bromo pattern); ¹H-NMR (400 MHz, CDCl₃) δ ppm: 1.37 - 1.40 (m, 3 H) 1.43 - 1.44 (m, 9 H) 4.34 - 4.69 (m, 2 H) 5.22 - 5.36 (m, 1 H) 7.86 - 8.08 (m, 2 H) 8.73 (d, J = 2.20 Hz, 1 H).

Preparation of tert-butyl N-[(1S)-3-(5-bromo-2-pyridyl)-1-methyl-2,3-dioxo-propyl]carbamate



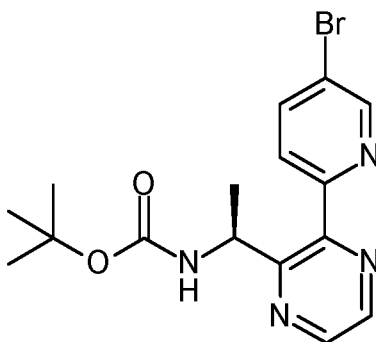
10

To a solution of tert-butyl N-[(1S)-3-(5-bromo-2-pyridyl)-2-hydroxy-1-methyl-3-oxo-propyl]carbamate (15.2 g, 42.3 mmol) in dichloromethane (100 mL) and dimethyl sulfoxide (20 mL) were added at 0 °C N,N-diisopropylethylamine (21.8 mL, 127 mmol, 3.00 equiv.) and in two portions sulfur trioxide pyridine complex (13.9 g, 84.6 mmol, 2.00 equiv.). The reaction mixture was stirred at 0 °C for 1 hour. It was quenched with water and diluted with dichloromethane and 1 N hydrochloric acid. The aqueous layer was extracted twice with dichloromethane. The combined organic layers were dried over magnesium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded tert-butyl N-[(1S)-3-(5-bromo-2-pyridyl)-1-methyl-2,3-dioxo-propyl]carbamate as an orange oil.

15

- 20 ¹H-NMR (400 MHz, CDCl₃) δ ppm: 1.36 - 1.41 (m, 9 H) 1.45 - 1.48 (m, 3 H) 4.82 - 4.96 (m, 1 H) 5.10 (br s, 1 H) 7.91 - 8.00 (m, 1 H) 8.01 - 8.11 (m, 1 H) 8.79 (d, J = 1.83 Hz, 1 H).

Preparation of tert-butyl N-[(1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]carbamate

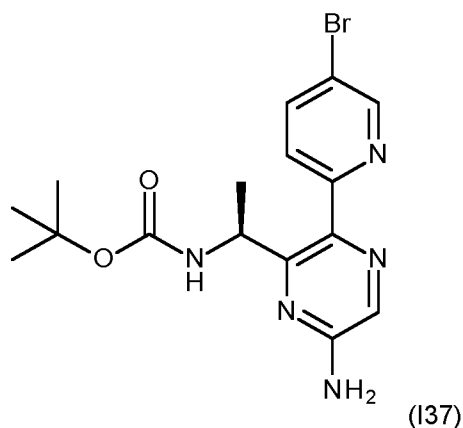


-87-

To a solution of tert-butyl N-[(1S)-3-(5-bromo-2-pyridyl)-1-methyl-2,3-dioxo-propyl]carbamate (375 mg, 1.05 mmol) in ethanol (22 mL) was added ethylenediamine (0.36 mL, 5.24 mmol). The reaction mixture was stirred at room temperature for 60 h in the presence of air. It was concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded tert-butyl N-[(1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]carbamate as a colorless gum.

LCMS: Rt 1.09, m/z = 379-381 (M+H⁺) (Bromo pattern); ¹H-NMR (400 MHz, CDCl₃) δ ppm: 1.33 - 1.45 (m, 9 H) 1.52 - 1.56 (m, 3 H) 5.65 - 5.83 (m, 2 H) 7.96 - 8.02 (m, 2 H) 8.53 - 8.60 (m, 2 H) 8.79 (dd, J = 2.20, 1.10 Hz, 1 H); Chiral SFC (method 1): 1.80 min (major enantiomer), 1.11 min (minor enantiomer); ee = 92%

Preparation of tert-butyl N-[(1S)-1-[6-amino-3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]carbamate (I37)

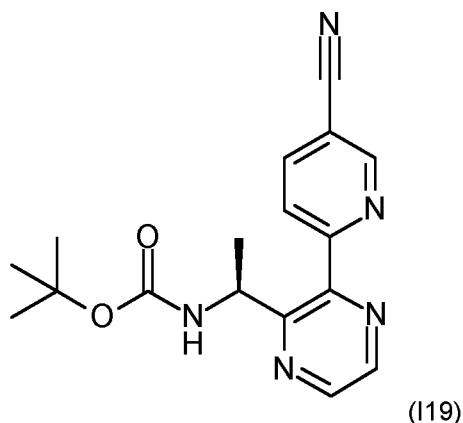


To a solution of tert-butyl N-[(1S)-3-(5-bromo-2-pyridyl)-1-methyl-2,3-dioxo-propyl]carbamate (500 mg, 0.894 mmol) in isopropanol (13.4 mL) was added 2-aminoacetamidine dihydrobromide (1.21 g, 4.11 mmol). Potassium acetate (266 mg, 2.68 mmol) was added. The reaction mixture was stirred at room temperature for 2.5 hours in the presence of air. The reaction was quenched with water and the aqueous layer was extracted with ethyl acetate. The combined organic layers were washed with brine, dried over magnesium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by reverse phase chromatography (C18, eluting with ACN in water) afforded tert-butyl N-[(1S)-1-[6-amino-3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]carbamate.

LCMS: Rt 1.00, m/z = 394-396 (M+H⁺) (Bromo pattern); ¹H-NMR (600 MHz, CDCl₃) δ ppm: 1.45 (br s, 9H) 1.47 (d, J=6.7 Hz, 3H) 4.84 (br s, 2H) 5.66 - 5.74 (m, 1H) 5.89 (br s, 1H) 7.86 - 7.88 (m, 1H) 7.89 (br d, J=2.0 Hz, 1H) 7.90 (s, 1H) 8.72 (s, 1H).

Preparation of tert-butyl N-[(1S)-1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]carbamate (I19)

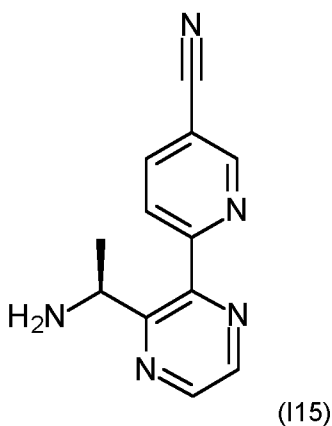
-88-



Degassed 1,4-dioxane (9.20 mL) was added to a mixture of tert-butyl N-[(1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]carbamate (1.396 g, 3.681 mmol), potassium ferricyanide (1.224 g, 3.681 mmol), tBuXPhos Pd-G3 (0.151 g, 0.184 mmol) and tBuXPhos (0.082 g, 0.18 mmol) at room temperature under argon. A degassed solution of potassium acetate (0.05 M in water, 9.20 mL, 0.500 mmol) was added and the mixture was stirred at 100 °C for 5 hours. The reaction mixture was diluted with water, then extracted three times with ethyl acetate. The combined organic layers were concentrated in vacuo. The crude material was purified by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) to afford tert-butyl N-[(1S)-1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]carbamate as a white solid.

¹H-NMR (400 MHz, CDCl₃) δ ppm: 1.38 (br s, 9 H), 1.55 (d, 3 H), 5.66 - 5.78 (m, 2 H), 8.11 (dd, 1 H), 8.30 (d, 1 H), 8.59 (d, 1 H), 8.63 (d, 1 H), 8.93 - 9.04 (m, 1 H).

Preparation of 6-[3-[(1S)-1-aminoethyl]pyrazin-2-yl]pyridine-3-carbonitrile (I15)



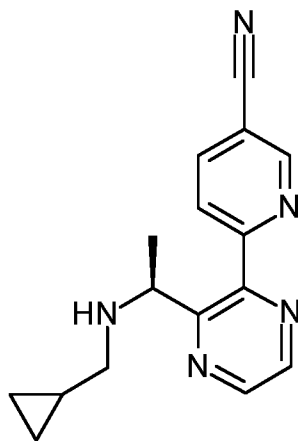
Trifluoroacetic acid (0.69 mL, 8.7 mmol) was added to a solution of tert-butyl N-[(1S)-1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]carbamate (0.520 g, 1.60 mmol) in dichloromethane (3.5 mL). The mixture was stirred at room temperature for 18 hours. The reaction mixture was concentrated in vacuo. The residue was dissolved in dichloromethane, then washed with saturated aqueous sodium carbonate, dried over magnesium sulfate and concentrated in vacuo to afford 6-[3-[(1S)-1-aminoethyl]pyrazin-2-yl]pyridine-3-carbonitrile.

-89-

LCMS (method 1): Rt 0.28, m/z = 226 [M+H]⁺; ¹H-NMR (400 MHz, CDCl₃) δ ppm: 1.48 (d, J=6.60 Hz, 3H) 1.96 (s, 2H) 4.74 (q, J=6.60 Hz, 1H) 8.09 - 8.14 (m, 1H) 8.19 - 8.23 (m, 1H) 8.55 (d, J=2.57 Hz, 1H) 8.65 (d, J=2.20 Hz, 1H) 8.97 (dd, J=2.20, 0.73 Hz, 1H)

5

Preparation of 6-[3-[(1S)-1-(cyclopropylmethylamino)ethyl]pyrazin-2-yl]pyridine-3-carbonitrile (I16)



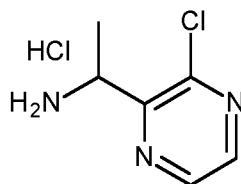
(I16)

To a solution of 6-[3-[(1S)-1-aminoethyl]pyrazin-2-yl]pyridine-3-carbonitrile (0.200 g, 0.888 mmol) in 1,2-dichloroethane (4.4 mL) were added cyclopropanecarboxaldehyde (0.0745 mL, 0.977 mmol), acetic acid (0.051 mL, 0.89 mmol) and sodium triacetoxyborohydride (0.277 g, 1.24 mmol). The mixture was stirred at room temperature for 1.5 hour. Saturated aqueous sodium carbonate was added, the aqueous layer was extracted with ethyl acetate. The combined organic layers were dried over magnesium sulfate and concentrated in vacuo. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded 6-[3-[(1S)-1-(cyclopropylmethylamino)ethyl]pyrazin-2-yl]pyridine-3-carbonitrile as a yellow oil.

¹H NMR (400 MHz, CDCl₃) δ ppm: -0.05 (dd, 2H) 0.39 (td, 2H) 0.81 - 0.94 (m, 1H) 1.50 (d, 3H) 1.98 (dd, 1H) 2.44 (dd, 1H) 2.49 - 2.89 (m, 1H) 4.73 (q, 1H) 8.11 - 8.18 (m, 1H) 8.20 - 8.27 (m, 1H) 8.57 (d, 1H) 8.69 (d, 1H) 9.00 (dd, 1H)

20

Preparation of 1-(3-chloropyrazin-2-yl)ethanamine hydrochloride



At room temperature, to a solution of 1-(3-chloropyrazin-2-yl)ethanone [CAS 121246-90-0] (5.00 g, 31.9 mmol) in methanol (80 mL) were added portionwise ammonium acetate (49.7 g, 639 mmol) and sodium cyanoborohydride (2.11 g, 31.9 mmol). The resulting suspension was stirred at room temp. overnight before being concentrated in vacuo. The residual was picked up in ethyl acetate and 2M

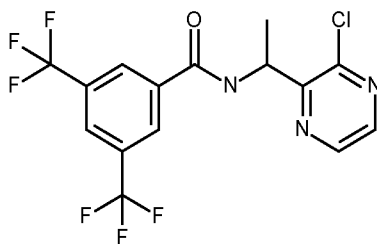
25

-90-

NaOH. The organic layer was dried (MgSO_4), filtered and evaporated. The residue was dissolved in diethyl ether and HCl in ethyl acetate (25 mL) was added dropwise. The formed precipitate was filtered off and dried to afford 1-(3-chloropyrazin-2-yl)ethanamine hydrochloride as a beige solid.

^1H NMR (400 MHz, DMSO-d_6) δ ppm: 1.52 (s, 3 H) 4.77 (br s, 1 H) 8.61 (d, $J = 2.57$ Hz, 1 H) 8.78 (d, $J = 2.2$ Hz, 1 H); LC-MS (method 1): Rt 0.17 min, m/z 158 $[\text{M}+\text{H}^+]$.

Preparation of N-[1-(3-chloropyrazin-2-yl)ethyl]-3,5-bis(trifluoromethyl)benzamide (I20)



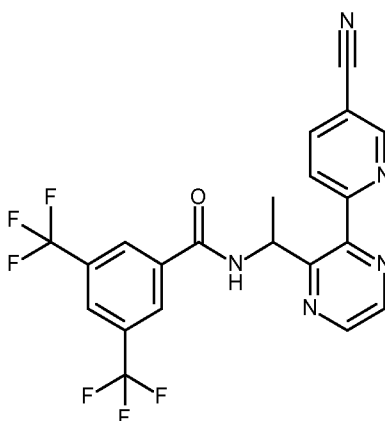
(I20)

1-(3-chloropyrazin-2-yl)ethanamine hydrochloride (2.50 g, 12.9 mmol) was suspended in 2-methyl-tetrahydrofuran (51 mL). N,N-diisopropylethylamine (6.68 mL, 38.6 mmol) was added followed by 3,5-bis(trifluoromethyl)benzoyl chloride [CAS 785-56-8] (2.41 mL, 12.9 mmol). The resulting suspension was stirred at room temp. overnight. The reaction mixture was diluted with ethyl acetate and water was added. Organic layer was isolated, dried (MgSO_4), filtered and evaporated. Purification by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded N-[1-(3-chloropyrazin-2-yl)ethyl]-3,5-bis(trifluoromethyl)benzamide as colorless solid.

^1H NMR (400 MHz, Chloroform-d) δ ppm: 1.65 (d, $J = 6.97$ Hz, 3 H) 5.77 – 5.83 (m, 1 H) 7.63 (br d, $J = 6.97$ Hz, 1 H) 8.08 (s, 1 H) 8.30 (s, 2 H) 8.41 (d, $J = 2.57$ Hz, 1 H) 8.54 (d, $J = 2.57$ Hz, 1 H)

LC-MS (method 1): Rt 1.09 min, m/z 398 $[\text{M}+\text{H}^+]$.

Preparation of N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide (P31)



(P31)

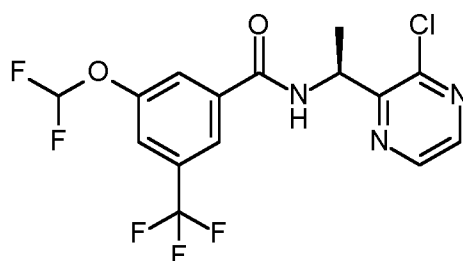
A solution of N-[1-(3-chloropyrazin-2-yl)ethyl]-3,5-bis(trifluoromethyl)benzamide (step 2, 300 mg, 0.754 mmol) in 8 mL DMF was purged with Ar, followed by the addition of (5-cyano-2-pyridyl)boronic acid [CAS 910547-29-4] (223 mg, 1.51 mmol), diacetoxypalladium (25 mg, 0.0377 mmol, 5 mol%) and dppf (42 mg, 0.0754 mmol, 10 mol%), copper(II) chloride (101 mg, 0.754 mmol) and cesium carbonate (492

-91-

mg, 1.51 mmol). After purging again with Ar, the reaction mixture was heated overnight at 100°C. After cooling, the reaction mixture was poured into a saturated ammonium chloride solution and extracted twice with ethyl acetate, the combined organic phases were washed with water, dried over sodium sulfate, filtered and concentrated to get a brown gum which was purified by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) to afford N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide.

¹H NMR (400 MHz, CDCl₃) δ ppm: 1.71 (d, 3 H) 6.46 (m, 1 H) 7.69 (b, d, 1 H) 8.03 (m, 1 H) 8.19 (d, 1 H) 8.25 (s, 2 H) 8.39 (d, 1 H) 8.70 (s, 2 H) 9.10 (s, 1 H); LC-MS (method 1): R_t 1.11 min, m/z 466 [M+H⁺].

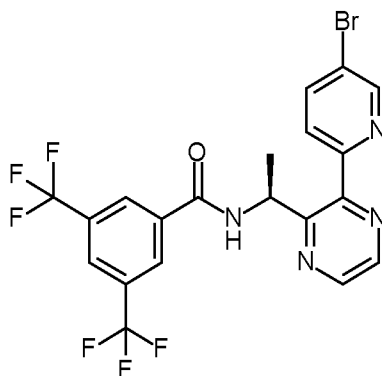
Preparation of N-[(1S)-1-(3-chloropyrazin-2-yl)ethyl]-3-(difluoromethoxy)-5-(trifluoromethyl)benzamide (intermediate I3)



(I3)

To a solution of 3-(difluoromethoxy)-5-(trifluoromethyl)benzoic acid (0.140 g, 0.547 mmol) in dichloromethane (1.4 mL) were added triethylamine (0.230 mL, 1.60 mmol) and one drop of N,N-dimethylformamide. Then oxalyl chloride (0.0950 mL, 1.10 mmol) was added and the resulting mixture was stirred at room temperature for 20 minutes. It was concentrated under reduced pressure. The resulting residue was dissolved in dichloromethane (0.7 mL) and a suspension of (1S)-1-(3-chloropyrazin-2-yl)ethanamine (80 mg, 0.41 mmol) and triethylamine (0.23 mL, 1.6 mmol) in dichloromethane (0.7 mL) was added dropwise at 0 °C to the previous solution of acid chloride. The reaction mixture was stirred at room temperature for 1.5 hours. It was quenched dropwise at 0 °C by addition of sodium bicarbonate sat. aq. and diluted with dichloromethane. The aqueous layer was extracted three times with dichloromethane. The combined organic layers were dried over magnesium sulfate, filtered and concentrated under reduced pressure. A first purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) failed. Thus the recovered fractions containing the desired product were dissolved in ethyl acetate and washed several times with sodium bicarbonate sat. aq., then brine, to remove the carboxylic acid as the impurity. The organic layer was dried over magnesium sulfate, filtered and concentrated under reduced pressure to afford N-[(1S)-1-(3-chloropyrazin-2-yl)ethyl]-3-(difluoromethoxy)-5-(trifluoromethyl)benzamide as an orange gum.

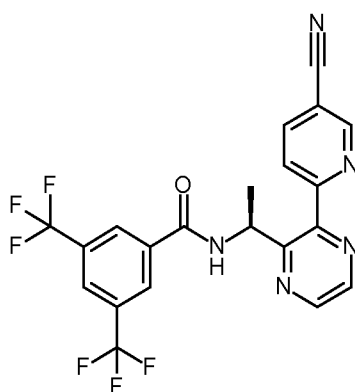
LCMS (method 1): R_t 1.04, m/z = 396 [M+H⁺]; ¹H-NMR (400 MHz, CDCl₃) δ ppm: 1.63 (d, J = 6.60 Hz, 3 H) 5.77 (quin, J = 6.97 Hz, 1H) 6.41-6.84 (m, 1H) 7.55 (s, 1H) 7.58 (br s, 1H) 7.81 (s, 1H) 7.92 (s, 1H) 8.40 (d, J = 2.57 Hz, 1H) 8.52 (d, J = 2.57 Hz, 1H).

Preparation of N-[(1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide

To a solution of (1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethanamine (0.539 g, 1.93 mmol) in ethyl acetate (8 mL) were added sodium bicarbonate (1 N solution in water, 7.7 mL, 7.7 mmol) and 3,5-bis(trifluoromethyl)benzoyl chloride (0.385 mL, 2.12 mmol). The biphasic reaction mixture was stirred vigorously at room temperature for 1.5 hours. The layers were separated and the aqueous layer was extracted twice with ethyl acetate. The combined organic layers were dried over magnesium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded N-[(1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide as a beige solid.

LC-MS (method 1): Rt 1.21 min, m/z 519/521 $[M+H]^+$, bromo pattern; ^1H -NMR (400 MHz, $CDCl_3$) δ ppm: 1.66 (d, J = 6.60 Hz, 3 H) 6.27-6.34 (m, 1 H) 7.86 (br d, J = 7.70 Hz, 1 H) 8.00-8.09 (m, 3 H) 8.25 (s, 2 H) 8.63 (q, J = 2.20 Hz, 2 H) 8.86 (dd, J = 2.20, 0.73 Hz, 1 H).

Example P34: Preparation of N-[(1S)-1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide (P34)



(P34)

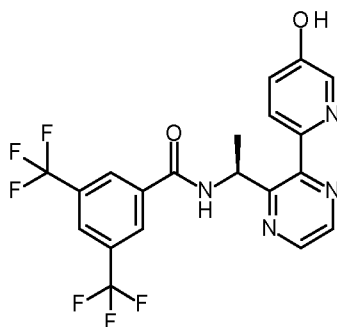
In a microwave vial under argon, were charged N-[(1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide (intermediate I-5 prepared as described above, 0.200 g, 0.385 mmol), potassium ferricyanide (0.064 g, 0.19 mmol, 0.50 equiv.), [(2-di-tert-butylphosphino-2',4',6'-triisopropyl-1,1'-biphenyl)-2-(2'-amino-1,1'-biphenyl)] palladium(II) methanesulfonate (tBuXPhos Pd G3) (7.9 mg, 9.6 μ mol, 0.025 equiv.) and 2-di-tert-butylphosphino-2',4',6'-triisopropylbiphenyl (tBuXPhos) (4.3 mg, 9.6 mmol, 0.025 equiv.). The vial was sealed and degassed three times with argon. Then previously

-93-

degassed 1,4-dioxane (0.96 mL) and previously prepared and degassed potassium acetate (0.05 M solution in water, 0.96 mL, 0.10 equiv.) were added to the reaction mixture. It was heated up to 100 °C and stirred overnight. After cooling down to room temperature, the reaction mixture was diluted with water and extracted three times with ethyl acetate. The combined organic layers were dried over magnesium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded the desired product as a white solid (36 mg, 77 µmol).

LC-MS (method 1): Rt 1.10, m/z = 465 [M+H⁺]; ¹H-NMR (400 MHz, CDCl₃) δ ppm: 1.72 (d, J = 6.60 Hz, 3 H) 6.29-6.40 (m, 1 H) 7.68 (br d, J = 7.70 Hz, 1 H) 8.03 (s, 1 H) 8.19 (dd, J = 8.44, 2.20 Hz, 1 H) 8.24 (s, 2 H) 8.39 (dd, J = 8.44, 0.73 Hz, 1 H) 8.67-8.73 (m, 2 H) 9.09 (dd, J = 2.02, 0.92 Hz, 1 H). Chiral SFC (method 2): 13.31 min (minor enantiomer), 15.02 min (major enantiomer); ee = 88%

Preparation of N-[(1S)-1-[3-[5-hydroxy-2-pyridyl]pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide

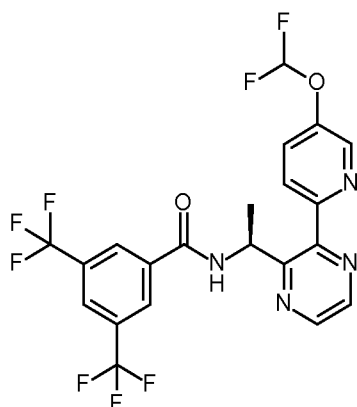


In a flask under argon, were charged N-[(1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide (0.150 g, 0.289 mmol), 1,4-dioxane (2.9 mL), water (1.2 mL), potassium carbonate (56.5 mg, 0.867 mmol), 2-di-tert-butylphosphino-3,4,5,6-tetramethyl-2',4',6'-triisopropyl-1,1'-biphenyl (1.46 mg, 2.89 µmol) and tris(dibenzylideneacetone)dipalladium(0) (5.51 mg, 5.78 µmol). The reaction mixture was heated up to 80 °C and stirred overnight. After cooling down to room temperature, it was diluted with ethyl acetate and water. The aqueous layer was extracted with ethyl acetate. The combined organic layers were dried over magnesium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded N-[(1S)-1-[3-[5-hydroxy-2-pyridyl]pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide.

LC-MS (method 1): Rt 1.01, m/z = 457 [M+H⁺]; ¹H-NMR (400 MHz, CDCl₃) δ ppm: 1.69 (d, J = 6.60 Hz, 3 H) 6.38-6.53 (m, 1 H) 7.30 (dd, J = 8.80, 2.93 Hz, 1 H) 7.99 (d, J = 8.44 Hz, 1 H) 8.05 (s, 1 H) 8.26 (br d, J = 8.07 Hz, 1 H) 8.30 (s, 2 H) 8.35 (d, J = 2.93 Hz, 1 H) 8.61 (dd, J = 14.12, 2.38 Hz, 2 H).

Example P24: Preparation of N-[(1S)-1-[3-[5-(difluoromethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide

-94-

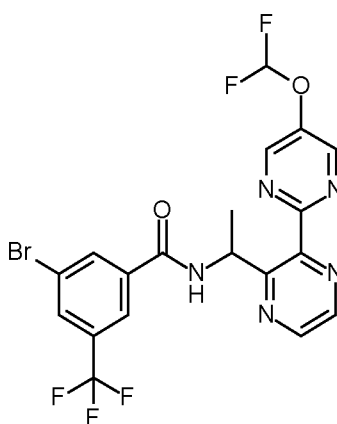


(P24)

In a flask, were charged N-[(1S)-1-[3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide (91.0 mg, 0.199 mmol), N,N-dimethylformamide (2 mL), potassium carbonate (0.281 g, 1.99 mmol) and chlorodifluoroacetic acid (88.9 μ L, 0.997 mmol). The reaction mixture was heated up to 80 $^{\circ}$ C and stirred for 4 hours. After cooling down to room temperature, it was diluted with ethyl acetate and water. The aqueous layer was extracted with ethyl acetate. The combined organic layers were dried over magnesium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with ethyl acetate in cyclohexane) afforded N-[(1S)-1-[3-[5-(difluoromethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide as a yellow solid.

LC-MS (method 1): R_t 1.16, m/z = 507 $[M+H]^+$; ^1H -NMR (400 MHz, $CDCl_3$) δ ppm: 1.68 (d, J = 6.60 Hz, 3 H) 6.27-6.40 (m, 1 H) 6.45-6.91 (m, 1 H) 7.70 (dd, J = 8.8, 2.6 Hz, 1 H) 7.94 (br d, J = 7.7 Hz, 1 H) 8.02 (s, 1 H) 8.23 (d, J = 8.8 Hz, 1 H) 8.27 (s, 2 H) 8.58 - 8.73 (m, 3 H); ^{19}F -NMR (377 MHz, $CDCl_3$) δ ppm: -81.58 (s, 2 F, -CHF₂) -62.88 (s, 6 F, -CF₃).

Example P23: Preparation of 3-bromo-N-[1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide



(P23)

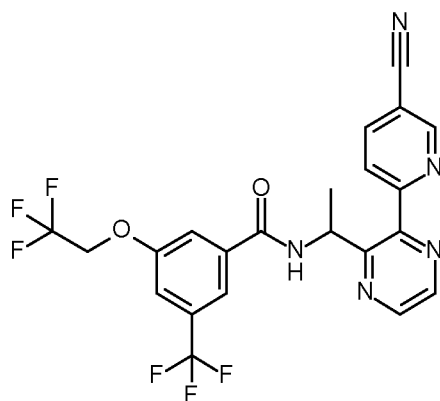
To a solution of 3-bromo-5-(trifluoromethyl)benzoic acid (150 mg, 0.558 mmol) in toluene (3.0 mL) was added thionyl chloride (0.122 mL, 1.67 mmol) dropwise. The reaction mixture was stirred at reflux for 2 hours, cooled to room temperature, and concentrated under reduced pressure to afford the

-95-

crude acyl chloride. The acyl chloride was dissolved in dichloromethane (3.0 mL) and a solution of 1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine (182 mg, 0.669 mmol) in dichloromethane (3.0 mL) and triethylamine (0.235 mL, 1.67 mmol) was added dropwise over 5 minutes. The reaction mixture was stirred at room temperature for 2 hours. Water and ethyl acetate were added to the reaction mixture. The aqueous layer was extracted with ethyl acetate. The combined organic layers were dried over sodium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by flash chromatography over silica gel (eluting with methanol in dichloromethane) afforded 3-bromo-N-[1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide as a pale yellow solid.

¹H-NMR (400 MHz, DMSO-d₆) δ ppm: 1.59 (d, 3 H) 5.50-5.65 (m, 1 H) 7.48 (t, 1H) 8.03 (s, 1 H) 8.12 (s, 1H) 8.20 (s, 1H) 8.68 (s, 1H) 8.77 (s, 1H) 8.94 (s, 1H) 9.19 (m, 1H)

Example P25: Preparation of N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-3-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)benzamide



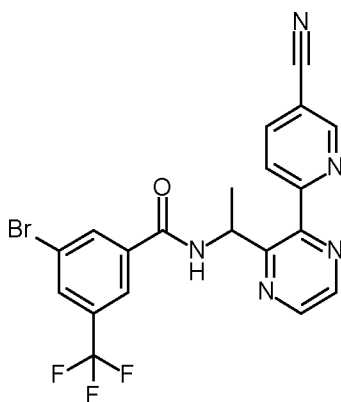
(P25)

Thionyl chloride (0.0721 mL, 0.989 mmol) was added dropwise to a stirred solution of 3-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)benzoic acid (150 mg, 0.495 mmol) in toluene (8.00 mL) at 0 °C. The reaction mixture was heated to 90 °C for 2.5 hours, then cooled down to room temperature and concentrated under reduced pressure. The crude acyl chloride was dissolved in dichloromethane (10 mL) and added to a stirred solution of 6-[3-(1-aminoethyl)pyrazin-2-yl]pyridine-3-carbonitrile (129 mg, 0.544 mmol) and triethylamine (0.139 mL, 0.989 mmol) in dichloromethane (10 mL) at 0 °C. the reaction mixture was stirred for 5 hours at room temperature. Dichloromethane and water were added. The organic layer was dried over sodium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by reverse-phase chromatography (C18 column, eluting with acetonitrile in water) afforded N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-3-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)benzamide as an off-white solid.

¹H-NMR (400 MHz, DMSO-d₆) δ ppm: 1.65 (d, 3 H) 4.85-5.00 (m, 2H) 5.70-5.80 (m, 1H) 7.55 (s, 1H) 7.70-7.80 (m, 2H) 8.12-8.21 (m, 1H) 8.48-8.53 (m, 1H) 8.68-8.80 (m, 2H) 9.10-9.20 (m, 2H); ¹⁹F-NMR (377 MHz, DMSO-d₆) δ ppm: -72.53 (s, 3H) -61.11 (s, 3H)

-96-

Example P10: Preparation of 3-bromo-N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide

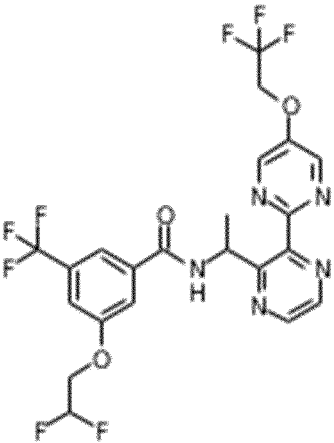
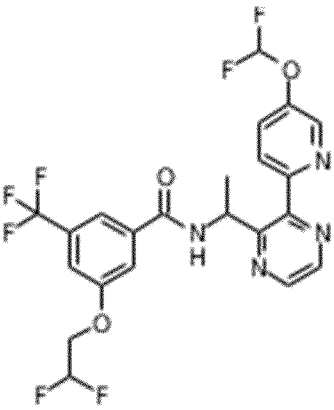
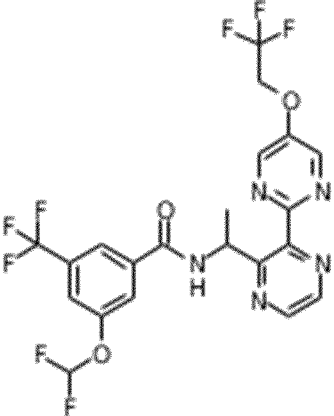


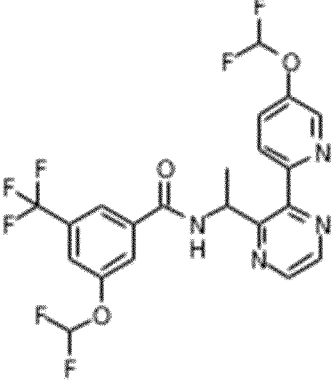
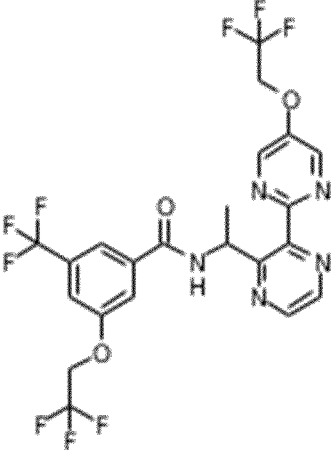
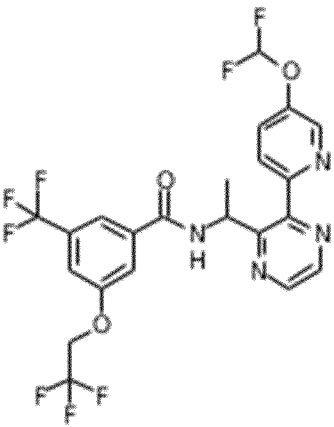
(P10)

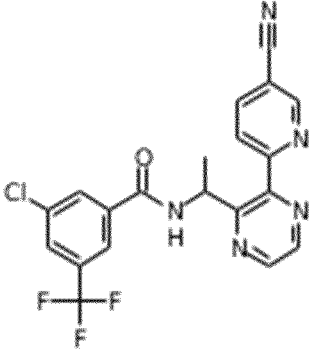
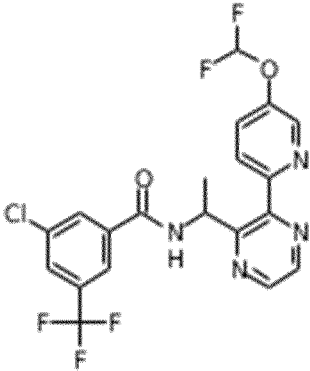
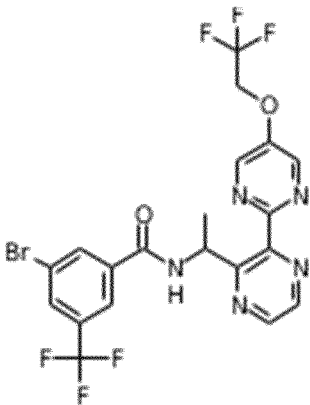
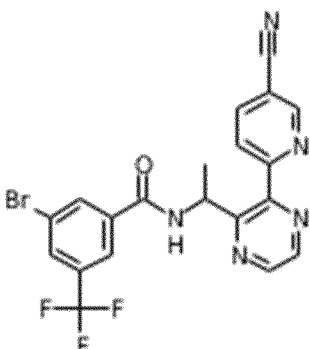
- 5 Thionyl chloride (0.140 mL, 1.91 mmol) was added dropwise to a stirred solution of 3-bromo-5-(trifluoromethyl)benzoic acid (130 mg, 0.478 mmol) in toluene (20.00 mL) at 0 °C. The reaction mixture was heated to 90 °C for 2 hours, then cooled down to room temperature and concentrated under reduced pressure. The crude acyl chloride was dissolved in dichloromethane (10 mL) and added to a stirred solution of 6-[3-(1-aminoethyl)pyrazin-2-yl]pyridine-3-carbonitrile (148 mg, 0.593 mmol) and triethylamine (0.269 mL, 1.91 mmol) in dichloromethane (10 mL) at 0 °C. The reaction mixture was stirred for 2 hours at room temperature. Dichloromethane and water were added. The organic layer was dried over sodium sulfate, filtered and concentrated under reduced pressure. Purification of the crude material by reverse-phase chromatography (C18 column, eluting with acetonitrile in water) afforded 3-bromo-N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide as an off-white solid.

¹H-NMR (400 MHz, DMSO-d₆) δ ppm: 1.63 (d, 3 H) 5.70-5.81 (m, 1H) 8.08 (s, 1H) 8.12 (s, 1H) 8.15-8.21 (m, 1H), 8.23 (s, 1H) 8.48-8.52 (m, 1H) 8.68-8.72 (m, 1H) 8.72-8.80 (m, 1H) 9.15-9.19 (m, 1H) 9.22-9.30 (m, 1H); ¹⁹F-NMR (377 MHz, DMSO-d₆) δ ppm: -61.25 (s, 3H)

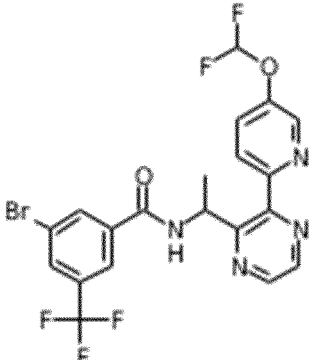
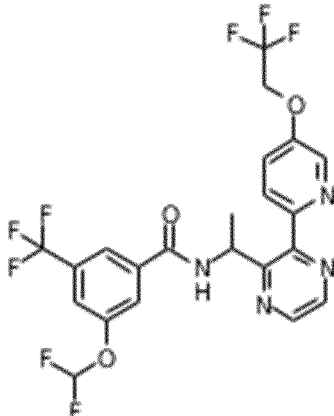
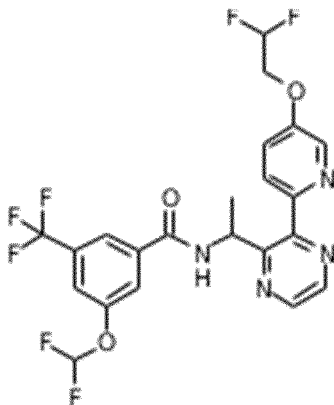
Table P: Examples of compounds of formula I

	IUPAC name	STRUCTURE	RT (min)	[M+H] (measured)	Method	MP °C
P1	3-(2,2-difluoroethoxy)-N-[1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					65 - 70
P2	3-(2,2-difluoroethoxy)-N-[1-[3-[5-(difluoromethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					130 - 140
P3	3-(difluoromethoxy)-N-[1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					122 - 125

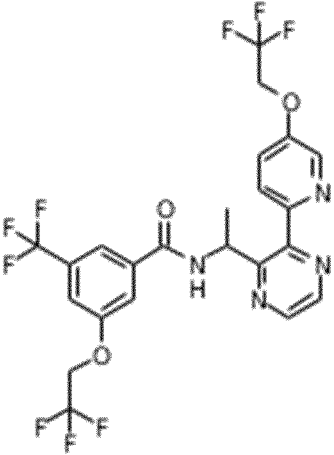
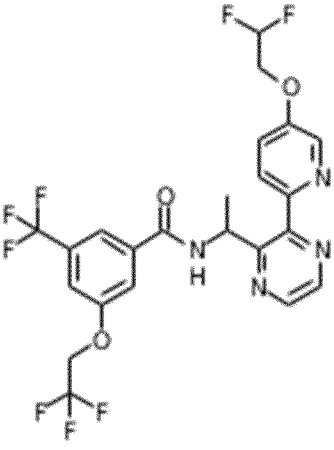
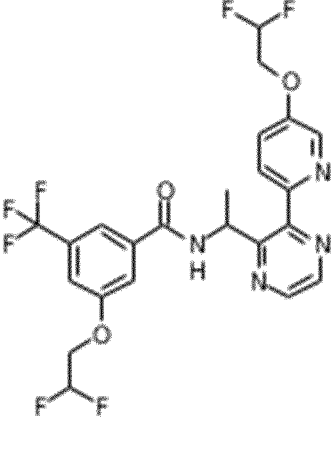
P4	3-(difluoromethoxy)-N-[1-[3-[5-(difluoromethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					140 - 150
P5	3-(2,2,2-trifluoroethoxy)-N-[1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					105 - 110
P6	N-[1-[3-[5-(difluoromethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-3-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)benzamide					130 - 140

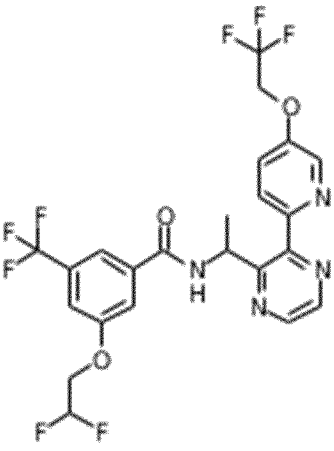
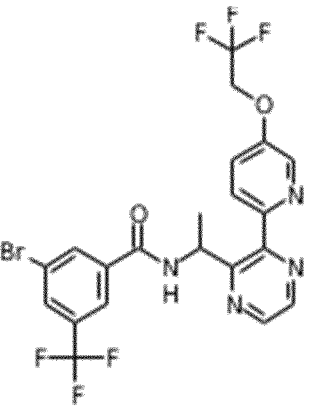
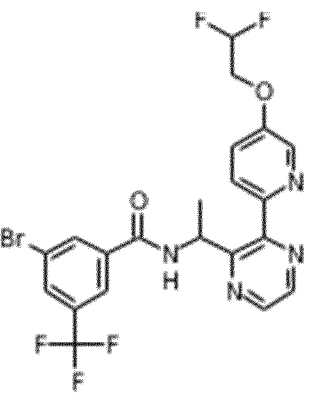
P7	3-chloro-N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					170 - 175
P8	3-chloro-N-[1-[3-[5-(difluoromethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					130 - 140
P9	3-bromo-N-[1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					85 - 90
P10	3-bromo-N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					195 - 200

-100-

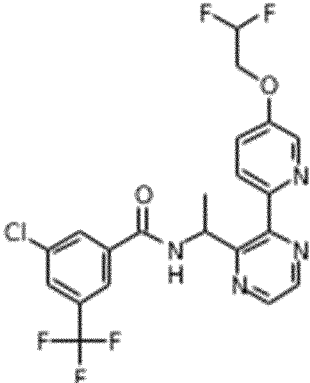
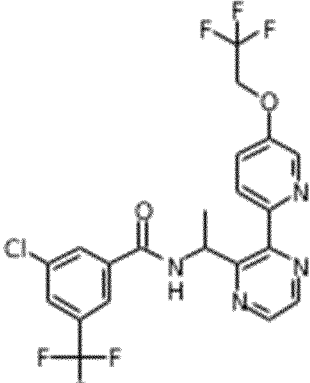
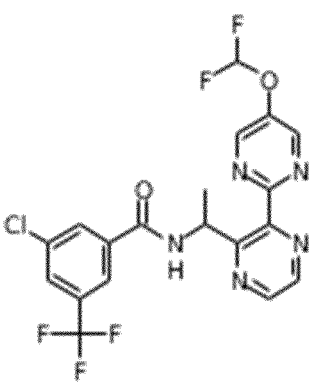
P11	3-bromo-N-[1-[3-[5-(difluoromethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					140 - 150
P12	3-(difluoromethoxy)-N-[1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					130 - 135
P13	N-[1-[3-[5-(2,2-difluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-3-(difluoromethoxy)-5-(trifluoromethyl)benzamide					115 - 120

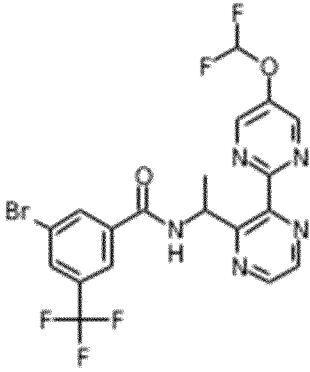
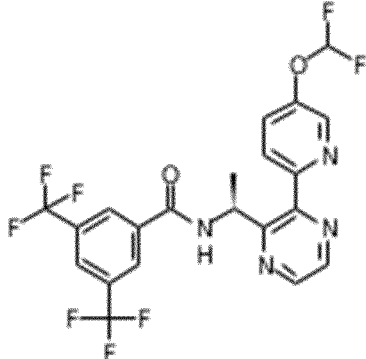
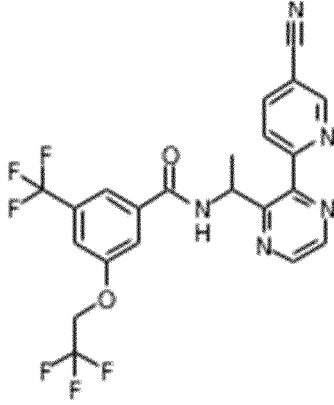
-101-

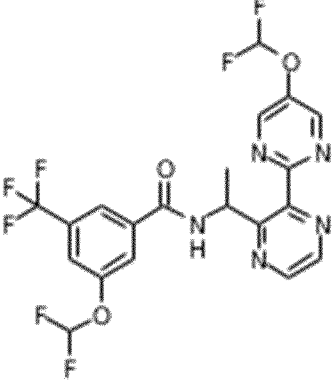
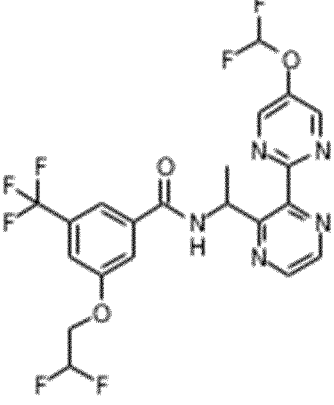
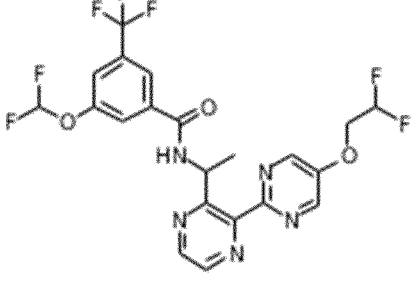
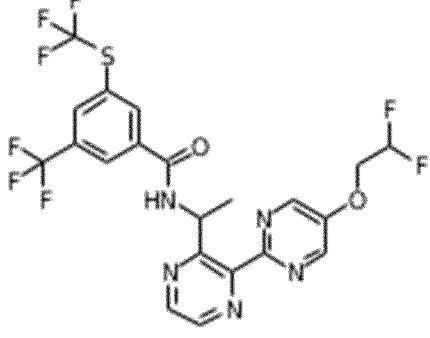
P14	3-(2,2,2-trifluoroethoxy)-N-[1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					130 - 135
P15	N-[1-[3-[5-(2,2-difluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-3-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)benzamide					130 - 135
P16	3-(2,2-difluoroethoxy)-N-[1-[3-[5-(2,2-difluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					120 - 125

P17	3-(2,2-difluoroethoxy)-N-[1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					130 - 135
P18	3-bromo-N-[1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					140 - 150
P19	3-bromo-N-[1-[3-[5-(2,2-difluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					140 - 150

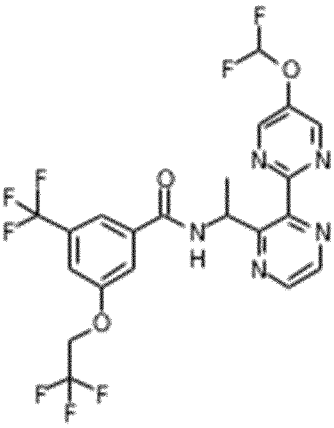
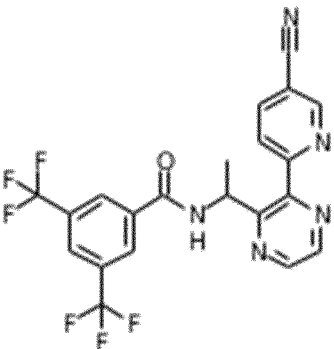
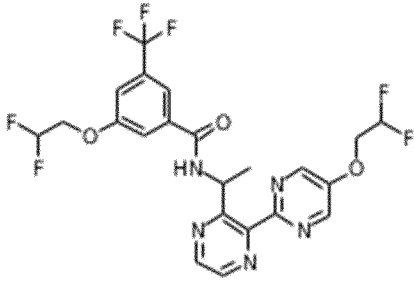
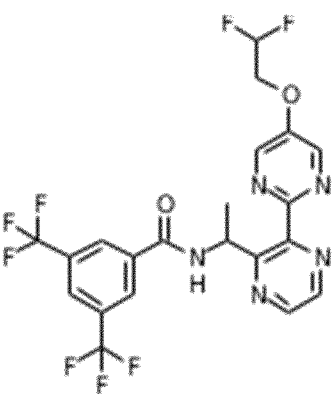
-103-

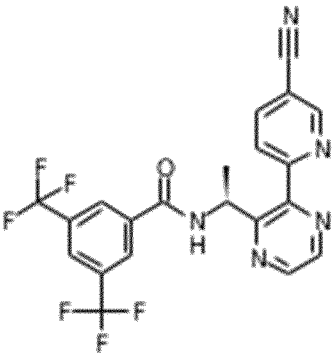
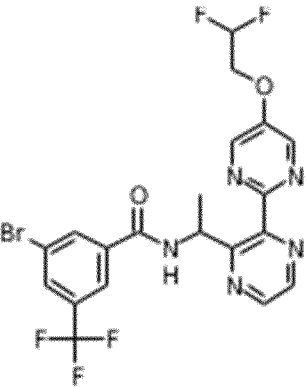
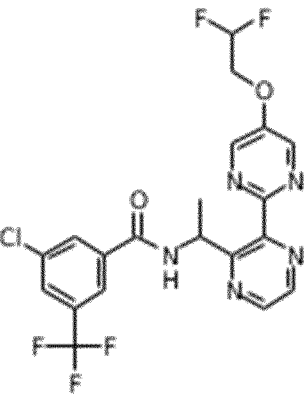
P20	3-chloro-N-[1-[3-[5-(2,2-difluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					120 - 130
P21	3-chloro-N-[1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					140 - 150
P22	3-chloro-N-[1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					100 - 110

P23	3-bromo-N-[1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					100 - 110
P24	N-[(1S)-1-[3-[5-(difluoromethoxy)-2-pyridyl]pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide					153 - 159
P25	N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-3-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)benzamide					165 - 175

P26	3-(difluoromethoxy)-N-[1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					120 - 125
P27	3-(2,2-difluoroethoxy)-N-[1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					130 - 135
P28	N-[1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-3-(difluoromethoxy)-5-(trifluoromethyl)benzamide					82 - 86
P29	N-[1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-3-(trifluoromethyl)-5-(trifluoromethylsulfanyl)benzamide					92 - 95

-106-

P30	N-[1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-3-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)benzamide					110 - 115
P31	N-[1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide		1.11	466.3	1	
P32	3-(2,2-difluoroethoxy)-N-[1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					130 - 132
P33	N-[1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide					115 - 120

P34	N-[(1S)-1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]-3,5-bis(trifluoromethyl)benzamide		1.11	466.3	1	
P35	3-bromo-N-[1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					60 - 65
P36	3-chloro-N-[1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					105 - 110

-108-

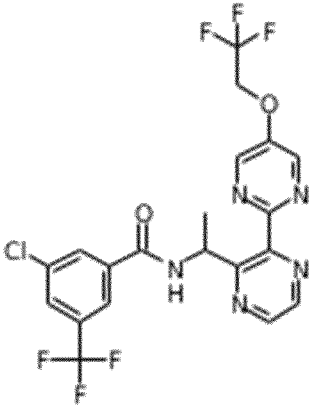
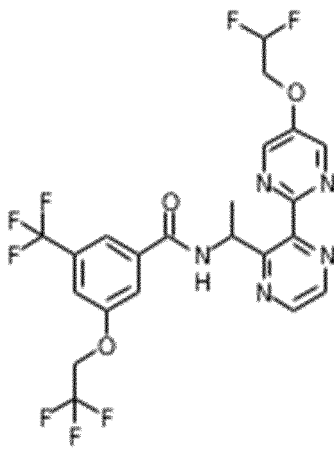
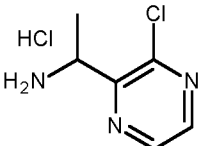
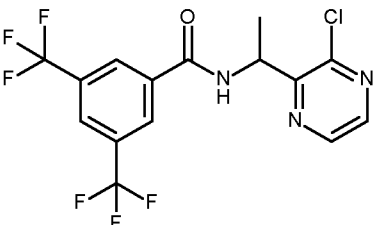
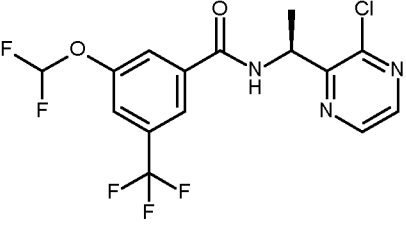
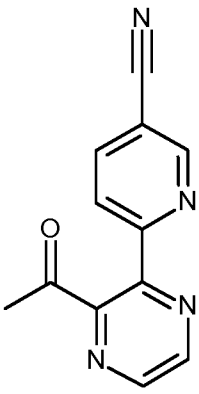
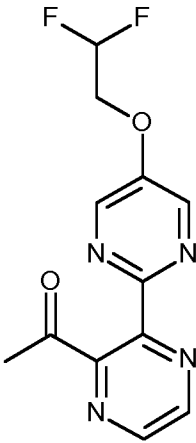
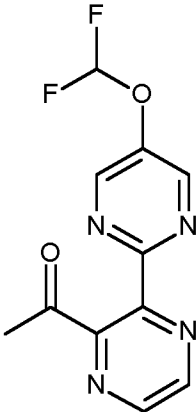
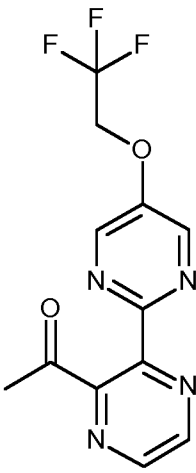
P37	3-chloro-N-[1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-5-(trifluoromethyl)benzamide					74 - 78
P38	N-[1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethyl]-3-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)benzamide					130 - 135

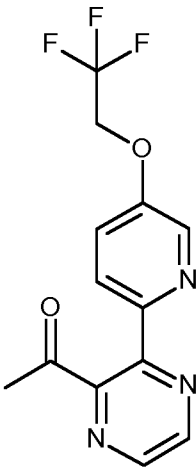
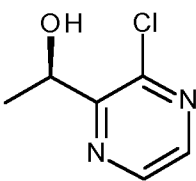
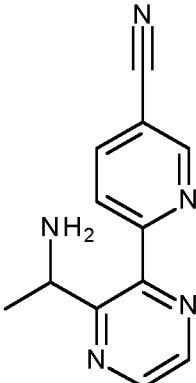
Table I: Table of Intermediates

Index	IUPAC name	STRUCTURE	RT (min)	[M+H] (m/Z)	Method	NMR
I1	methyl 2-chloro-6-(trifluoromethyl)pyridine-4-carboxylate		0.17	158	1	
I2	N-[1-(3-chloropyrazin-2-yl)ethyl]-3,5-bis(trifluoromethyl)benzamide		1.09	398	1	
I3	N-[(1S)-1-(3-chloropyrazin-2-yl)ethyl]-3-(difluoromethoxy)-5-(trifluoromethyl)benzamide		1.04	396.3	1	
I4	6-(3-acetylpyrazin-2-yl)pyridine-3-carbonitrile		0.73	225.1	1	

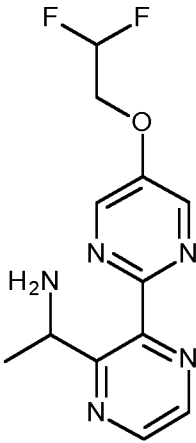
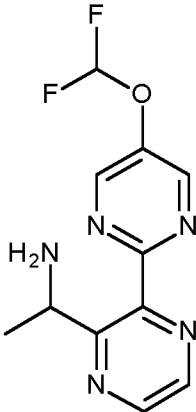
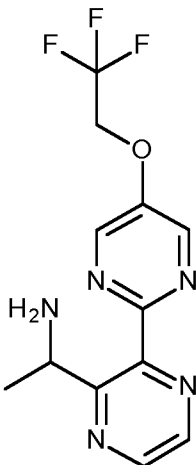
-110-

Index	IUPAC name	STRUCTURE	RT (min)	[M+H] (m/Z)	Method	NMR
I5	1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone					1)
I6	1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone					2)
I7	1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanone			299.1		

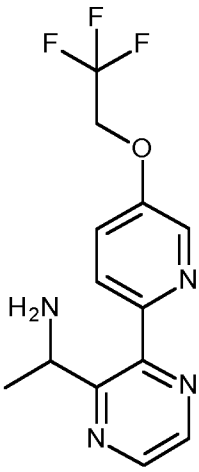
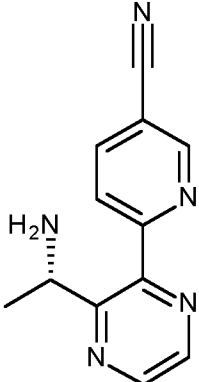
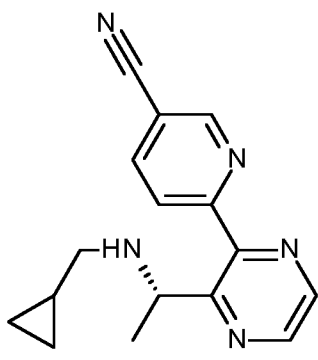
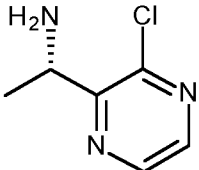
-111-

Index	IUPAC name	STRUCTURE	RT (min)	[M+H] (m/Z)	Method	NMR
I8	<u>1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethanone</u>					3)
I9	(1R)-1-(3-chloropyrazin-2-yl)ethanol		0.40	159/160	1	
I10	6-[3-(1-aminoethyl)pyrazin-2-yl]pyridine-3-carbonitrile					4)

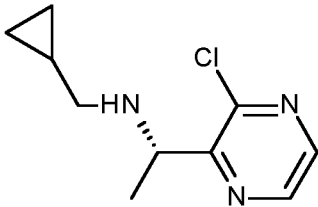
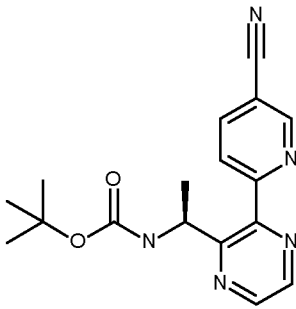
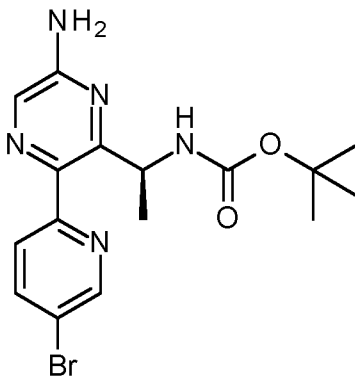
-112-

Index	IUPAC name	STRUCTURE	RT (min)	[M+H] (m/Z)	Method	NMR
I11	1-[3-[5-(2,2-difluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine					5)
I12	1-[3-[5-(difluoromethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine					6)
I13	1-[3-[5-(2,2,2-trifluoroethoxy)pyrimidin-2-yl]pyrazin-2-yl]ethanamine					7)

-113-

Index	IUPAC name	STRUCTURE	RT (min)	[M+H] (m/z)	Method	NMR
I14	1-[3-[5-(2,2,2-trifluoroethoxy)-2-pyridyl]pyrazin-2-yl]ethanamine					8)
I15	6-[3-[(1S)-1-aminoethyl]pyrazin-2-yl]pyridine-3-carbonitrile		0.28	226	1	
I16	6-[3-[(1S)-1-(cyclopropylmethylamino)ethyl]pyrazin-2-yl]pyridine-3-carbonitrile					9)
I17	(1S)-1-(3-chloropyrazin-2-yl)ethanamine		0.17	158	1	

-114-

Index	IUPAC name	STRUCTURE	RT (min)	[M+H] (m/Z)	Method	NMR
I18	(1S)-1-(3-chloropyrazin-2-yl)-N-(cyclopropylmethyl)ethanamine		0.26	212	1	
I19	tert-butyl N-[(1S)-1-[3-(5-cyano-2-pyridyl)pyrazin-2-yl]ethyl]carbamate		0.91	326.4	1	
I20	tert-butyl N-[(1S)-1-[6-amino-3-(5-bromo-2-pyridyl)pyrazin-2-yl]ethyl]carbamate		0.99	394/396 [M+H] ⁺ (bromo pattern)	1	

¹) ¹H NMR (400 MHz, DMSO-d₆) δ ppm: 8.90 (s, 1H), 8.70-8.90 (m, 3H), 6.48 (t, 1H), 4.63 (td, 2H), 2.62 (s, 3H)

²) ¹H NMR (400 MHz, DMSO-d₆) δ ppm: 8.80-9.00 (m, 4H), 7.50 (t, 1H), 2.65 (s, 3H)

5 ³) ¹H-NMR (400 MHz, DMSO-d₆): δ ppm 8.85 (d, 1H), 8.70 (m, 1H), 8.45 (s, 1H), 8.25 (d, 1H), 7.75 (d, 1H), 5 (q, 2H), 2.6 (s, 3H)

⁴) ¹H-NMR (400 MHz, DMSO-d₆): δ ppm 9.22 (s, 1H), 8.85-8.95 (m, 2H), 8.50-8.60 (m, 1H), 8.30-8.40 (m, 1H) 7.80-8.10 (br. s, 2H), 5.25-5.35 (m, 1 H), 1.52 (d, 3H)

10 ⁵) ¹H NMR (400 MHz, DMSO-d₆) δ ppm: 8.80-9.00 (m, 4H), 6.50 (tt, 1H), 4.90 (m, 1H), 4.78 (td, 2H), 1.45 (d, 3H)

⁶) ¹H NMR (400 MHz, DMSO-d₆) δ ppm: 8.80-9.10 (m, 4H), 7.51 (t, 1H), 4.88 (m, 1H), 1.50 (d, 3H)

⁷) ¹H-NMR (400 MHz, DMSO-d₆): δ ppm: 8.95 (s, 2H), 8.60-9.00 (m, 2H), 7.8-8.30 (br s, 2H), 5.08-5.20 (m, 2H), 4.95-5.05 (m, 1H), 1.5 (m, 3H)

-115-

⁸⁾ ¹H-NMR (400 MHz, DMSO-d₆): δ ppm: 8.8 (s, 2H), 8.65 (d, 1H), 8.15 (d, 1H), 7.8 (m, 1H), 7.45 (br s, 2H), 7.25 (m, 1H), 7.15 (m, 1H), 5.2 (br s, 1H), 5 (q, 2H), 1.5 (m, 3H)

⁹⁾ ¹H NMR (400 MHz, chloroform-d) δ ppm -0.05 (dd, 2H) 0.39 (td, 2H) 0.81 - 0.94 (m, 1H) 1.50 (d, 3H) 1.98 (dd, 1H) 2.44 (dd, 1H) 2.49 - 2.89 (m, 1H) 4.73 (q, 1H) 8.11 - 8.18 (m, 1H) 8.20 - 8.27 (m, 1H) 8.57 (d, 1H) 8.69 (d, 1H) 9.00 (dd, 1H)

The activity of the compositions according to the invention can be broadened considerably, and adapted to prevailing circumstances, by adding other insecticidally, acaricidally and/or fungicidally active ingredients. The mixtures of the compounds of formula I with other insecticidally, acaricidally and/or fungicidally active ingredients may also have further surprising advantages which can also be described, in a wider sense, as synergistic activity. For example, better tolerance by plants, reduced phytotoxicity, insects can be controlled in their different development stages or better behaviour during their production, for example during grinding or mixing, during their storage or during their use.

Suitable additions to active ingredients here are, for example, representatives of the following classes of active ingredients: organophosphorus compounds, nitrophenol derivatives, thioureas, juvenile hormones, formamidines, benzophenone derivatives, ureas, pyrrole derivatives, carbamates, pyrethroids, chlorinated hydrocarbons, acylureas, pyridylmethyleamino derivatives, macrolides, neonicotinoids and *Bacillus thuringiensis* preparations.

The following mixtures of the compounds of formula I with active ingredients are preferred (where the abbreviation "TX" means "one compound selected from the compounds defined in the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27, and Table P"):

an adjuvant selected from the group of substances consisting of petroleum oils (alternative name) (628) + TX,

an insect control active substance selected from Abamectin + TX, Acequinocyl + TX, Acetamiprid + TX, Acetoprole + TX, Acrinathrin + TX, Acynonapyr + TX, Afidopyropen + TX, Afoxalaner + TX, Alanycarb + TX, Allethrin + TX, Alpha-Cypermethrin + TX, Alphamethrin + TX, Amidoflumet + TX, Aminocarb + TX, Azocyclotin + TX, Bensultap + TX, Benzoximate + TX, Benzpyrimoxan + TX, Betacyfluthrin + TX, Beta-cypermethrin + TX, Bifenazate + TX, Bifenthrin + TX, Binapacryl + TX, Bioallethrin + TX, Bioallethrin S)-cyclopentylisomer + TX, Bioresmethrin + TX, Bistrifluron + TX, Broflanilide + TX, Brofluthrin + TX, Bromophos-ethyl + TX, Buprofezine + TX, Butocarboxim + TX, Cadusafos + TX, Carbaryl + TX, Carbosulfan + TX, Cartap + TX, CAS number: 1472050-04-6 + TX, CAS number: 1632218-00-8 + TX, CAS number: 1808115-49-2 + TX, CAS number: 2032403-97-5 + TX, CAS number: 2044701-44-0 + TX, CAS number: 2128706-05-6 + TX, CAS number: 2249718-27-0 + TX, Chlorantraniliprole + TX, Chlordane + TX, Chlorfenapyr + TX, Chloroprallethrin + TX, Chromafenozide + TX, Clenpirin + TX, Cloethocarb + TX, Clothianidin + TX, 2-chlorophenyl N-methylcarbamate (CPMC) + TX, Cyanofenphos + TX, Cyantraniliprole + TX, Cyclaniliprole + TX, Cyclobutirfluram + TX, Cycloprothrin + TX, Cycloxaprid + TX, Cyenopyrafen + TX,

- Cyetyprafen (or Etpyrafen) + TX, Cyflumetofen + TX, Cyfluthrin + TX, Cyhalodiamide + TX, Cyhalothrin + TX, Cypermethrin + TX, Cyphenothrin + TX, Cyromazine + TX, Deltamethrin + TX, Diafenthiuron + TX, Dialifos + TX, Dibrom + TX, Dicloromezotiaz + TX, Diflovidazine + TX, Diflubenzuron + TX, dimpropyridaz + TX, Dinactin + TX, Dinocap + TX, Dinotefuran + TX,
- 5 Dioxabenzofos + TX, Emamectin + TX, Empenthrin + TX, Epsilon - momfluorothrin + TX, Epsilon-metofluthrin + TX, Esfenvalerate + TX, Ethion + TX, Ethiprole + TX, Etofenprox + TX, Etoazole + TX, Famphur + TX, Fenazaquin + TX, Fenfluthrin + TX, Fenitrothion + TX, Fenobucarb + TX, Fenothiocarb + TX, Fenoxycarb + TX, Fenpropathrin + TX, Fenpyroxymate + TX, Fensulfothion + TX, Fenthion + TX, Fentinacetate + TX, Fenvalerate + TX, Fipronil + TX, Flometoquin + TX, Flonicamid + TX,
- 10 Fluacrypyrim + TX, Fluazaindolizine + TX, Fluazuron + TX, Flubendiamide + TX, Flubenzimine + TX, Flucitrate + TX, Flucycloxuron + TX, Flucythrinate + TX, Fluensulfone + TX, Flufenerim + TX, Flufenprox + TX, Flufiprole + TX, Fluhexafon + TX, Flumethrin + TX, Fluopyram + TX, Flupentiofenox + TX, Flupyradifurone + TX, Flupyrimin + TX, Fluralaner + TX, Fluvalinate + TX, Fluxametamide + TX, Fosthiazate + TX, Gamma-Cyhalothrin + TX, Gossypure™ + TX, Guadipyr + TX, Halofenozide + TX,
- 15 Halofenozide + TX, Halofenprox + TX, Heptafluthrin + TX, Hexythiazox + TX, Hydramethylnon + TX, Imicyafos + TX, Imidacloprid + TX, Imiprothrin + TX, Indoxacarb + TX, Iodomethane + TX, Iprodione + TX, Isocycloseram + TX, Isothioate + TX, Ivermectin + TX, Kappa-bifenthrin + TX, Kappa-tefluthrin + TX, Lambda-Cyhalothrin + TX, Lepimectin + TX, Lufenuron + TX, Metaflumizone + TX, Metaldehyde + TX, Metam + TX, Methomyl + TX, Methoxyfenozide + TX, Metofluthrin + TX, Metolcarb + TX,
- 20 Mexacarbate + TX, Milbemectin + TX, Momfluorothrin + TX, Niclosamide + TX, Nitenpyram + TX, Nithiazine + TX, Omethoate + TX, Oxamyl + TX, Oxazosufyl + TX, Parathion-ethyl + TX, Permethrin + TX, Phenothrin + TX, Phosphocarb + TX, Piperonylbutoxide + TX, Pirimicarb + TX, Pirimiphos-ethyl + TX, Polyhedrosis virus + TX, Prallethrin + TX, Profenofos + TX, Profenofos + TX, Profluthrin + TX, Propargite + TX, Propetamphos + TX, Propoxur + TX, Prothiophos + TX, Protrifenbute + TX,
- 25 Pyflubumide + TX, Pymetrozine + TX, Pyraclofos + TX, Pyrafluprole + TX, Pyridaben + TX, Pyridalyl + TX, Pyrfluquinazon + TX, Pyrimidifen + TX, Pyrimostrobin + TX, Pyriprole + TX, Pyriproxyfen + TX, Resmethrin + TX, Sarolaner + TX, Selamectin + TX, Silafluofen + TX, Spinetoram + TX, Spinosad + TX, Spirodiclofen + TX, Spiromesifen + TX, Spiropidion + TX, Spirotetramat + TX, Sulfoxaflor + TX, Tebufenozide + TX, Tebufenpyrad + TX, Tebupirimiphos + TX, Tefluthrin + TX, Temephos + TX,
- 30 Tetrachloraniliprole + TX, Tetradiaphon + TX, Tetramethrin + TX, Tetramethylfluthrin + TX, Tetranactin + TX, Tetraniliprole + TX, Theta-cypermethrin + TX, Thiacloprid + TX, Thiamethoxam + TX, Thiocyclam + TX, Thiodicarb + TX, Thiofanox + TX, Thiometon + TX, Thiosultap + TX, Tioxazafen + TX, Tolfenpyrad + TX, Toxaphene + TX, Tralomethrin + TX, Transfluthrin + TX, Triazamate + TX, Triazophos + TX, Trichlorfon + TX, Trichloronate + TX, Trichlorphon + TX, Triflumezopyrim + TX,
- 35 Tyclopyrazoflor + TX, Zeta-Cypermethrin + TX, Extract of seaweed and fermentation product derived from melasse + TX, Extract of seaweed and fermentation product derived from melasse comprising urea + TX, amino acids + TX, potassium and molybdenum and EDTA-chelated manganese + TX, Extract of seaweed and fermented plant products + TX, Extract of seaweed and fermented plant products comprising phytohormones + TX, vitamins + TX, EDTA-chelated copper + TX, zinc + TX, and

-117-

- iron + TX, Azadirachtin + TX, *Bacillus aizawai* + TX, *Bacillus chitosporus* AQ746 (NRRL Accession No B-21 618) + TX, *Bacillus firmus* + TX, *Bacillus kurstaki* + TX, *Bacillus mycoides* AQ726 (NRRL Accession No. B-21664) + TX, *Bacillus pumilus* (NRRL Accession No B-30087) + TX, *Bacillus pumilus* AQ717 (NRRL Accession No. B-21662) + TX, *Bacillus* sp. AQ178 (ATCC Accession No. 53522) + TX,
- 5 *Bacillus* sp. AQ175 (ATCC Accession No. 55608) + TX, *Bacillus* sp. AQ177 (ATCC Accession No. 55609) + TX, *Bacillus subtilis* unspecified + TX, *Bacillus subtilis* AQ153 (ATCC Accession No. 55614) + TX, *Bacillus subtilis* AQ30002 (NRRL Accession No. B-50421) + TX, *Bacillus subtilis* AQ30004 (NRRL Accession No. B- 50455) + TX, *Bacillus subtilis* AQ713 (NRRL Accession No. B-21661) + TX, *Bacillus subtilis* AQ743 (NRRL Accession No. B-21665) + TX, *Bacillus thuringiensis* AQ52 (NRRL
- 10 Accession No. B-21619) + TX, *Bacillus thuringiensis* BD#32 (NRRL Accession No B-21530) + TX, *Bacillus thuringiensis* subspec. *kurstaki* BMP 123 + TX, *Beauveria bassiana* + TX, D-limonene + TX, Granulovirus + TX, Harpin + TX, *Helicoverpa armigera* Nucleopolyhedrovirus + TX, *Helicoverpa zea* Nucleopolyhedrovirus + TX, *Heliothis virescens* Nucleopolyhedrovirus + TX, *Heliothis punctigera* Nucleopolyhedrovirus + TX, *Metarhizium* spp. + TX, *Muscodor albus* 620 (NRRL Accession No.
- 15 30547) + TX, *Muscodor roseus* A3-5 (NRRL Accession No. 30548) + TX, Neem tree based products + TX, *Paecilomyces fumosoroseus* + TX, *Paecilomyces lilacinus* + TX, *Pasteuria nishizawae* + TX, *Pasteuria penetrans* + TX, *Pasteuria ramosa* + TX, *Pasteuria thornei* + TX, *Pasteuria usgae* + TX, P-cymene + TX, *Plutella xylostella* Granulosis virus + TX, *Plutella xylostella* Nucleopolyhedrovirus + TX, Polyhedrosis virus + TX, pyrethrum + TX, QRD 420 (a terpenoid blend) + TX, QRD 452 (a terpenoid
- 20 blend) + TX, QRD 460 (a terpenoid blend) + TX, *Quillaja saponaria* + TX, *Rhodococcus globerulus* AQ719 (NRRL Accession No B-21663) + TX, *Spodoptera frugiperda* Nucleopolyhedrovirus + TX, *Streptomyces galbus* (NRRL Accession No. 30232) + TX, *Streptomyces* sp. (NRRL Accession No. B-30145) + TX, Terpenoid blend + TX, and *Verticillium* spp.,
- an algicide selected from the group of substances consisting of bethoxazin [CCN] + TX, copper
- 25 dioctanoate (IUPAC name) (170) + TX, copper sulfate (172) + TX, cybutryne [CCN] + TX, dichlone (1052) + TX, dichlorophen (232) + TX, endothal (295) + TX, fentin (347) + TX, hydrated lime [CCN] + TX, nabam (566) + TX, quinoxamine (714) + TX, quinonamid (1379) + TX, simazine (730) + TX, triphenyltin acetate (IUPAC name) (347) and triphenyltin hydroxide (IUPAC name) (347) + TX,
- 30 an anthelmintic selected from the group of substances consisting of abamectin (1) + TX, crufomate (1011) + TX, Cyclobutrifluram + TX, doramectin (alternative name) [CCN] + TX, emamectin (291) + TX, emamectin benzoate (291) + TX, eprinomectin (alternative name) [CCN] + TX, ivermectin (alternative name) [CCN] + TX, milbemycin oxime (alternative name) [CCN] + TX, moxidectin (alternative name) [CCN] + TX, piperazine [CCN] + TX, selamectin (alternative name) [CCN] + TX,
- 35 spinosad (737) and thiophanate (1435) + TX,
- an avicide selected from the group of substances consisting of chloralose (127) + TX, endrin (1122) + TX, fenthion (346) + TX, pyridin-4-amine (IUPAC name) (23) and strychnine (745) + TX,
- a bactericide selected from the group of substances consisting of 1-hydroxy-1*H*-pyridine-2-thione (IUPAC name) (1222) + TX, 4-(quinoxalin-2-ylamino)benzenesulfonamide (IUPAC name) (748) + TX,

- 8-hydroxyquinoline sulfate (446) + TX, bronopol (97) + TX, copper dioctanoate (IUPAC name) (170) + TX, copper hydroxide (IUPAC name) (169) + TX, cresol [CCN] + TX, dichlorophen (232) + TX, dipyrithione (1105) + TX, dodicin (1112) + TX, fenaminosulf (1144) + TX, formaldehyde (404) + TX, hydrargaphen (alternative name) [CCN] + TX, kasugamycin (483) + TX, kasugamycin
- 5 hydrochloride hydrate (483) + TX, nickel bis(dimethyldithiocarbamate) (IUPAC name) (1308) + TX, nitrapyrin (580) + TX, octhilineone (590) + TX, oxolinic acid (606) + TX, oxytetracycline (611) + TX, potassium hydroxyquinoline sulfate (446) + TX, probenazole (658) + TX, streptomycin (744) + TX, streptomycin sesquisulfate (744) + TX, tecloftalam (766) + TX, and thiomersal (alternative name) [CCN] + TX,
- 10 a biological agent selected from the group of substances consisting of *Adoxophyes orana* GV (alternative name) (12) + TX, *Agrobacterium radiobacter* (alternative name) (13) + TX, *Amblyseius* spp. (alternative name) (19) + TX, *Anagrapha falcifera* NPV (alternative name) (28) + TX, *Anagrus atomus* (alternative name) (29) + TX, *Aphelinus abdominalis* (alternative name) (33) + TX, *Aphidius colemani* (alternative name) (34) + TX, *Aphidoletes aphidimyza* (alternative name) (35) + TX,
- 15 *Autographa californica* NPV (alternative name) (38) + TX, *Bacillus firmus* (alternative name) (48) + TX, *Bacillus sphaericus* Neide (scientific name) (49) + TX, *Bacillus thuringiensis* Berliner (scientific name) (51) + TX, *Bacillus thuringiensis* subsp. *aizawai* (scientific name) (51) + TX, *Bacillus thuringiensis* subsp. *israelensis* (scientific name) (51) + TX, *Bacillus thuringiensis* subsp. *japonensis* (scientific name) (51) + TX, *Bacillus thuringiensis* subsp. *kurstaki* (scientific name) (51) + TX,
- 20 *Bacillus thuringiensis* subsp. *tenebrionis* (scientific name) (51) + TX, *Beauveria bassiana* (alternative name) (53) + TX, *Beauveria brongniartii* (alternative name) (54) + TX, *Chrysoperla carnea* (alternative name) (151) + TX, *Cryptolaemus montrouzieri* (alternative name) (178) + TX, *Cydia pomonella* GV (alternative name) (191) + TX, *Dacnusa sibirica* (alternative name) (212) + TX, *Diglyphus isaea* (alternative name) (254) + TX, *Encarsia formosa* (scientific name) (293) + TX,
- 25 *Eretmocerus eremicus* (alternative name) (300) + TX, *Helicoverpa zea* NPV (alternative name) (431) + TX, *Heterorhabditis bacteriophora* and *H. megidis* (alternative name) (433) + TX, *Hippodamia convergens* (alternative name) (442) + TX, *Leptomastix dactylopii* (alternative name) (488) + TX, *Macrolophus caliginosus* (alternative name) (491) + TX, *Mamestra brassicae* NPV (alternative name) (494) + TX, *Metaphycus helvolus* (alternative name) (522) + TX, *Metarhizium anisopliae* var.
- 30 *acridum* (scientific name) (523) + TX, *Metarhizium anisopliae* var. *anisopliae* (scientific name) (523) + TX, *Neodiprion sertifer* NPV and *N. lecontei* NPV (alternative name) (575) + TX, *Orius* spp. (alternative name) (596) + TX, *Paecilomyces fumosoroseus* (alternative name) (613) + TX, *Phytoseiulus persimilis* (alternative name) (644) + TX, *Spodoptera exigua* multicapsid nuclear polyhedrosis virus (scientific name) (741) + TX, *Steinernema bibionis* (alternative name) (742) + TX,
- 35 *Steinernema carpocapsae* (alternative name) (742) + TX, *Steinernema feltiae* (alternative name) (742) + TX, *Steinernema glaseri* (alternative name) (742) + TX, *Steinernema riobrave* (alternative name) (742) + TX, *Steinernema riobrave* (alternative name) (742) + TX, *Steinernema scapterisci* (alternative name) (742) + TX, *Steinernema* spp. (alternative name) (742) + TX, *Trichogramma* spp.

(alternative name) (826) + TX, *Typhlodromus occidentalis* (alternative name) (844) and *Verticillium lecanii* (alternative name) (848) + TX,

a soil sterilant selected from the group of substances consisting of iodomethane (IUPAC name) (542) and methyl bromide (537) + TX,

- 5 a chemosterilant selected from the group of substances consisting of apholate [CCN] + TX, bisazir (alternative name) [CCN] + TX, busulfan (alternative name) [CCN] + TX, diflubenzuron (250) + TX, dimatif (alternative name) [CCN] + TX, hemel [CCN] + TX, hempa [CCN] + TX, metepa [CCN] + TX, methiotepa [CCN] + TX, methyl apholate [CCN] + TX, morzid [CCN] + TX, penfluron (alternative name) [CCN] + TX, tepa [CCN] + TX, thiohempa (alternative name) [CCN] + TX, thiotepa (alternative name) [CCN] + TX,
- 10 tretamine (alternative name) [CCN] and uredepa (alternative name) [CCN] + TX, an insect pheromone selected from the group of substances consisting of (*E*)-dec-5-en-1-yl acetate with (*E*)-dec-5-en-1-ol (IUPAC name) (222) + TX, (*E*)-tridec-4-en-1-yl acetate (IUPAC name) (829) + TX, (*E*)-6-methylhept-2-en-4-ol (IUPAC name) (541) + TX, (*E,Z*)-tetradeca-4,10-dien-1-yl acetate (IUPAC name) (779) + TX, (*Z*)-dodec-7-en-1-yl acetate (IUPAC name) (285) + TX, (*Z*)-hexadec-11-
- 15 enal (IUPAC name) (436) + TX, (*Z*)-hexadec-11-en-1-yl acetate (IUPAC name) (437) + TX, (*Z*)-hexadec-13-en-11-yn-1-yl acetate (IUPAC name) (438) + TX, (*Z*)-icos-13-en-10-one (IUPAC name) (448) + TX, (*Z*)-tetradec-7-en-1-ol (IUPAC name) (782) + TX, (*Z*)-tetradec-9-en-1-ol (IUPAC name) (783) + TX, (*Z*)-tetradec-9-en-1-yl acetate (IUPAC name) (784) + TX, (*7E,9Z*)-dodeca-7,9-dien-1-yl acetate (IUPAC name) (283) + TX, (*9Z,11E*)-tetradeca-9,11-dien-1-yl acetate (IUPAC name) (780) +
- 20 TX, (*9Z,12E*)-tetradeca-9,12-dien-1-yl acetate (IUPAC name) (781) + TX, 14-methyloctadec-1-ene (IUPAC name) (545) + TX, 4-methylnonan-5-ol with 4-methylnonan-5-one (IUPAC name) (544) + TX, alpha-multistriatin (alternative name) [CCN] + TX, brevicomin (alternative name) [CCN] + TX, codlelure (alternative name) [CCN] + TX, codlemone (alternative name) (167) + TX, cuelure (alternative name) (179) + TX, disparlure (277) + TX, dodec-8-en-1-yl acetate (IUPAC name) (286)
- 25 + TX, dodec-9-en-1-yl acetate (IUPAC name) (287) + TX, dodeca-8 + TX, 10-dien-1-yl acetate (IUPAC name) (284) + TX, dominicalure (alternative name) [CCN] + TX, ethyl 4-methyloctanoate (IUPAC name) (317) + TX, eugenol (alternative name) [CCN] + TX, frontalinal (alternative name) [CCN] + TX, gossyplure (alternative name) (420) + TX, grandlure (421) + TX, grandlure I (alternative name) (421) + TX, grandlure II (alternative name) (421) + TX, grandlure III (alternative name) (421) + TX, grandlure IV (alternative name) (421) + TX, hexalure [CCN] + TX, ipsdienol (alternative name) [CCN] + TX, ipsenol (alternative name) [CCN] + TX, japonilure (alternative name) (481) + TX, lineatin (alternative name) [CCN] + TX, litlure (alternative name) [CCN] + TX, looplure (alternative name) [CCN] + TX, medlure [CCN] + TX, megatomoic acid (alternative name) [CCN] +
- 30 TX, methyl eugenol (alternative name) (540) + TX, muscalure (563) + TX, octadeca-2,13-dien-1-yl acetate (IUPAC name) (588) + TX, octadeca-3,13-dien-1-yl acetate (IUPAC name) (589) + TX, orfralure (alternative name) [CCN] + TX, oryctalure (alternative name) (317) + TX, ostramone (alternative name) [CCN] + TX, siglure [CCN] + TX, sordidin (alternative name) (736) + TX, sulcatol (alternative name) [CCN] + TX, tetradec-11-en-1-yl acetate (IUPAC name) (785) + TX, trimedlure (839) + TX, trimedlure A (alternative name) (839) + TX, trimedlure B₁ (alternative name) (839) + TX,

-120-

trimedlure B₂ (alternative name) (839) + TX, trimedlure C (alternative name) (839) and trunc-call (alternative name) [CCN] + TX,

an insect repellent selected from the group of substances consisting of 2-(octylthio)ethanol (IUPAC name) (591) + TX, butopyronoxyl (933) + TX, butoxy(polypropylene glycol) (936) + TX, dibutyl adipate (IUPAC name) (1046) + TX, dibutyl phthalate (1047) + TX, dibutyl succinate (IUPAC name) (1048) + TX, diethyltoluamide [CCN] + TX, dimethyl carbate [CCN] + TX, dimethyl phthalate [CCN] + TX, ethyl hexanediol (1137) + TX, hexamide [CCN] + TX, methoquin-butyl (1276) + TX, methylneodecanamide [CCN] + TX, oxamate [CCN] and picaridin [CCN] + TX,

a molluscicide selected from the group of substances consisting of bis(tributyltin) oxide (IUPAC name) (913) + TX, bromoacetamide [CCN] + TX, calcium arsenate [CCN] + TX, cloethocarb (999) + TX, copper acetoarsenite [CCN] + TX, copper sulfate (172) + TX, fentin (347) + TX, ferric phosphate (IUPAC name) (352) + TX, metaldehyde (518) + TX, methiocarb (530) + TX, niclosamide (576) + TX, niclosamide-olamine (576) + TX, pentachlorophenol (623) + TX, sodium pentachlorophenoxide (623) + TX, tazimcarb (1412) + TX, thiodicarb (799) + TX, tributyltin oxide (913) + TX, trifenmorph (1454) + TX, trimethacarb (840) + TX, triphenyltin acetate (IUPAC name) (347) and triphenyltin hydroxide (IUPAC name) (347) + TX, pyriprole [394730-71-3] + TX,

a nematicide selected from the group of substances consisting of AKD-3088 (compound code) + TX, 1,2-dibromo-3-chloropropane (IUPAC/Chemical Abstracts name) (1045) + TX, 1,2-dichloropropane (IUPAC/ Chemical Abstracts name) (1062) + TX, 1,2-dichloropropane with 1,3-dichloropropene (IUPAC name) (1063) + TX, 1,3-dichloropropene (233) + TX, 3,4-dichlorotetrahydrothiophene 1,1-dioxide (IUPAC/Chemical Abstracts name) (1065) + TX, 3-(4-chlorophenyl)-5-methylrhodanine (IUPAC name) (980) + TX, 5-methyl-6-thioxo-1,3,5-thiadiazinan-3-ylacetic acid (IUPAC name) (1286) + TX, 6-isopentenylaminopurine (alternative name) (210) + TX, abamectin (1) + TX, acetoprole [CCN] + TX, alanycarb (15) + TX, aldicarb (16) + TX, aldoxycarb (863) + TX, AZ 60541

(compound code) + TX, benclothiaz [CCN] + TX, benomyl (62) + TX, butylpyridaben (alternative name) + TX, cadusafos (109) + TX, carbofuran (118) + TX, carbon disulfide (945) + TX, carbosulfan (119) + TX, chloropicrin (141) + TX, chlorpyrifos (145) + TX, cloethocarb (999) + TX, Cyclobutrifluram + TX, cytokinins (alternative name) (210) + TX, dazomet (216) + TX, DBCP (1045) + TX, DCIP (218) + TX, diamidafos (1044) + TX, dichlofenthion (1051) + TX, dicliphos (alternative name) + TX, dimethoate (262) + TX, doramectin (alternative name) [CCN] + TX, emamectin (291) + TX, emamectin benzoate (291) + TX, eprinomectin (alternative name) [CCN] + TX, ethoprophos (312) + TX, ethylene dibromide (316) + TX, fenamiphos (326) + TX, fenpyrad (alternative name) + TX, fensulfthion (1158) + TX, fosthiazate (408) + TX, fosthietan (1196) + TX, furfural (alternative name) [CCN] + TX, GY-81 (development code) (423) + TX, heterophos [CCN] + TX, iodomethane (IUPAC name) (542) + TX, isamidofos (1230) + TX, isazofos (1231) + TX, ivermectin (alternative name) [CCN] + TX, kinetin (alternative name) (210) + TX, mecarphon (1258) + TX, metam (519) + TX, metam-potassium (alternative name) (519) + TX, metam-sodium (519) + TX, methyl bromide (537) + TX, methyl isothiocyanate (543) + TX, milbemycin oxime (alternative name) [CCN] + TX, moxidectin (alternative name) [CCN] + TX, *Myrothecium verrucaria* composition (alternative name)

-121-

- (565) + TX, NC-184 (compound code) + TX, oxamyl (602) + TX, phorate (636) + TX, phosphamidon (639) + TX, phosphocarb [CCN] + TX, sebufos (alternative name) + TX, selamectin (alternative name) [CCN] + TX, spinosad (737) + TX, terbam (alternative name) + TX, terbufos (773) + TX, tetrachlorothiophene (IUPAC/ Chemical Abstracts name) (1422) + TX, thiafenox (alternative name) + TX, thionazin (1434) + TX, triazophos (820) + TX, triazuron (alternative name) + TX, xlenols [CCN] + TX, YI-5302 (compound code) and zeatin (alternative name) (210) + TX, fluensulfone [318290-98-1] + TX, fluopyram + TX,
- a nitrification inhibitor selected from the group of substances consisting of potassium ethylxanthate [CCN] and nitrapyrin (580) + TX,
- 10 a plant activator selected from the group of substances consisting of acibenzolar (6) + TX, acibenzolar-S-methyl (6) + TX, probenazole (658) and *Reynoutria sachalinensis* extract (alternative name) (720) + TX,
- a rodenticide selected from the group of substances consisting of 2-isovalerylindan-1,3-dione (IUPAC name) (1246) + TX, 4-(quinoxalin-2-ylamino)benzenesulfonamide (IUPAC name) (748) + TX, alpha-chlorohydrin [CCN] + TX, aluminium phosphide (640) + TX, antu (880) + TX, arsenous oxide (882) + TX, barium carbonate (891) + TX, bisthiosemi (912) + TX, brodifacoum (89) + TX, bromadiolone (91) + TX, bromethalin (92) + TX, calcium cyanide (444) + TX, chloralose (127) + TX, chlorophacinone (140) + TX, cholecalciferol (alternative name) (850) + TX, coumachlor (1004) + TX, coumafuryl (1005) + TX, coumatetralyl (175) + TX, crimidine (1009) + TX, difenacoum
- 15 (246) + TX, difethialone (249) + TX, diphacinone (273) + TX, ergocalciferol (301) + TX, flocoumafen (357) + TX, fluoroacetamide (379) + TX, flupropadine (1183) + TX, flupropadine hydrochloride (1183) + TX, gamma-HCH (430) + TX, HCH (430) + TX, hydrogen cyanide (444) + TX, iodomethane (IUPAC name) (542) + TX, lindane (430) + TX, magnesium phosphide (IUPAC name) (640) + TX, methyl bromide (537) + TX, norbormide (1318) + TX, phosacetim (1336) + TX,
- 20 phosphine (IUPAC name) (640) + TX, phosphorus [CCN] + TX, pindone (1341) + TX, potassium arsenite [CCN] + TX, pyrinuron (1371) + TX, scilliroside (1390) + TX, sodium arsenite [CCN] + TX, sodium cyanide (444) + TX, sodium fluoroacetate (735) + TX, strychnine (745) + TX, thallium sulfate [CCN] + TX, warfarin (851) and zinc phosphide (640) + TX,
- a synergist selected from the group of substances consisting of 2-(2-butoxyethoxy)ethyl piperonylate (IUPAC name) (934) + TX, 5-(1,3-benzodioxol-5-yl)-3-hexylcyclohex-2-enone (IUPAC name) (903) + TX, farnesol with nerolidol (alternative name) (324) + TX, MB-599 (development code) (498) + TX, MGK 264 (development code) (296) + TX, piperonyl butoxide (649) + TX, piprotal (1343) + TX, propyl isomer (1358) + TX, S421 (development code) (724) + TX, sesamex (1393) + TX, sesasmolin (1394) and sulfoxide (1406) + TX,
- 30 an animal repellent selected from the group of substances consisting of anthraquinone (32) + TX, chloralose (127) + TX, copper naphthenate [CCN] + TX, copper oxychloride (171) + TX, diazinon (227) + TX, dicyclopentadiene (chemical name) (1069) + TX, guazatine (422) + TX, guazatine acetates (422) + TX, methiocarb (530) + TX, pyridin-4-amine (IUPAC name) (23) + TX, thiram (804) + TX, trimethacarb (840) + TX, zinc naphthenate [CCN] and ziram (856) + TX,

-122-

a virucide selected from the group of substances consisting of imanin (alternative name) [CCN] and ribavirin (alternative name) [CCN] + TX,

a wound protectant selected from the group of substances consisting of mercuric oxide (512) + TX, octhilinone (590) and thiophanate-methyl (802) + TX,

- 5 a biologically active substance selected from 1,1-bis(4-chloro-phenyl)-2-ethoxyethanol + TX, 2,4-dichlorophenyl benzenesulfonate + TX, 2-fluoro-N-methyl-N-1-naphthylacetamide + TX, 4-chlorophenyl phenyl sulfone + TX, acetoprole + TX, aldoxycarb + TX, amidithion + TX, amidothioate + TX, amiton + TX, amiton hydrogen oxalate + TX, amitraz + TX, aramite + TX, arsenous oxide + TX, azobenzene + TX, azothoate + TX, benomyl + TX, benoxa-fos + TX, benzyl benzoate + TX, bixafen + TX, brofenvalerate + TX, bromo-cyclen + TX, bromophos + TX, bromopropylate + TX, buprofezin + TX, butocarboxim + TX, butoxycarboxim + TX, butylpyridaben + TX, calcium polysulfide + TX, camphechlor + TX, carbanolate + TX, carbophenothion + TX, cymiazole + TX, chino-methionat + TX, chlorbenside + TX, chlordimeform + TX, chlordimeform hydrochloride + TX, chlorfenethol + TX, chlorfenson + TX, chlorfensulfide + TX, chlorobenzilate + TX, chloromebuform + TX, chloromethiuron + TX, chloropropylate + TX, chlorthiophos + TX, cinerin I + TX, cinerin II + TX, cinerins + TX, closantel + TX, coumaphos + TX, crotamiton + TX, crotoxyphos + TX, cufraneb + TX, cyanthoate + TX, DCPM + TX, DDT + TX, demephion + TX, demephion-O + TX, demephion-S + TX, demeton-methyl + TX, demeton-O + TX, demeton-O-methyl + TX, demeton-S + TX, demeton-S-methyl + TX, demeton-S-methylsulfon + TX, dichlofluanid + TX, dichlorvos + TX, dicliphos + TX, dienochlor + TX, dimefox + TX, dinex + TX, dinex-diclexine + TX, dinocap-4 + TX, dinocap-6 + TX, dinocton + TX, dino-penton + TX, dinosulfon + TX, dinoterbon + TX, dioxathion + TX, diphenyl sulfone + TX, disulfiram + TX, DNOC + TX, dofenapyn + TX, doramectin + TX, endothion + TX, eprinomectin + TX, ethoate-methyl + TX, etrimfos + TX, fenazaflor + TX, fenbutatin oxide + TX, fenothiocarb + TX, fenpyrad + TX, fen-pyroximate + TX, fenpyrazamine + TX, fenson + TX, fentripanil + TX, flubenzimine + TX, flucycloxuron + TX, fluenetil + TX, fluorbenside + TX, FMC 1137 + TX, formetanate + TX, formetanate hydrochloride + TX, formparanate + TX, gamma-HCH + TX, glyodin + TX, halfenprox + TX, hexadecyl cyclopropanecarboxylate + TX, isocarbophos + TX, jasmodin I + TX, jasmodin II + TX, jodfenphos + TX, lindane + TX, malonoben + TX, mecarbam + TX, mephosfolan + TX, mesulfen + TX, methacrifos + TX, methyl bromide + TX, metolcarb + TX, mexacarbate + TX, milbemycin oxime + TX, mipafox + TX, monocrotophos + TX, morphothion + TX, moxidectin + TX, naled + TX, 4-chloro-2-(2-chloro-2-methyl-propyl)-5-[(6-iodo-3-pyridyl)methoxy]pyridazin-3-one + TX, nifluridide + TX, nikkomycins + TX, nitrilacarb + TX, nitrilacarb 1:1 zinc chloride complex + TX, omethoate + TX, oxydeprofos + TX, oxydisulfoton + TX, pp'-DDT + TX, parathion + TX, permethrin + TX, phenkapton + TX, phosalone + TX, phosfolan + TX, phosphamidon + TX, polychloroterpenes + TX, polynactins + TX, proclonol + TX, promacyl + TX, propoxur + TX, prothidathion + TX, prothoate + TX, pyrethrin I + TX, pyrethrin II + TX, pyrethrins + TX, pyridaphenthion + TX, pyrimitate + TX, quinalphos + TX, quintiofos + TX, R-1492 + TX, phosglicin + TX, rotenone + TX, schradan + TX, sebufos + TX, selamectin + TX, sophamide + TX, SSI-121 + TX, sulfiram + TX, sulfluramid + TX, sulfotep + TX, sulfur + TX, diflovidazin + TX, tau-fluvalinate + TX, TEPP + TX, terbam + TX, tetradifon + TX, tetrasul + TX, thiafenox + TX,

- thiocarboxime + TX, thiofanox + TX, thiometon + TX, thioquinox + TX, thuringiensin + TX, triamphos + TX, triarathene + TX, triazophos + TX, triazuron + TX, trifenofos + TX, trinactin + TX, vamidothion + TX, vaniliprole + TX, bethoxazin + TX, copper dioctanoate + TX, copper sulfate + TX, cybutryne + TX, dichlone + TX, dichlorophen + TX, endothal + TX, fentin + TX, hydrated lime + TX, nabam + TX,
- 5 quinoclamine + TX, quinonamid + TX, simazine + TX, triphenyltin acetate + TX, triphenyltin hydroxide + TX, crufomate + TX, piperazine + TX, thiophanate + TX, chloralose + TX, fenthion + TX, pyridin-4-amine + TX, strychnine + TX, 1-hydroxy-1H-pyridine-2-thione + TX, 4-(quinoxalin-2-ylamino)benzenesulfonamide + TX, 8-hydroxyquinoline sulfate + TX, bronopol + TX, copper hydroxide + TX, cresol + TX, dipyrithione + TX, dodicin + TX, fenaminosulf + TX, formaldehyde + TX,
- 10 hydrargaphen + TX, kasugamycin + TX, kasugamycin hydrochloride hydrate + TX, nickel bis(dimethyldithiocarbamate) + TX, nitrapyrin + TX, octhilinone + TX, oxolinic acid + TX, oxytetracycline + TX, potassium hydroxyquinoline sulfate + TX, probenazole + TX, streptomycin + TX, streptomycin sesquisulfate + TX, tecloftalam + TX, thiomersal + TX, Adoxophyes orana GV + TX, Agrobacterium radiobacter + TX, Amblyseius spp. + TX, Anagrapta falcifera NPV + TX, Anagrus
- 15 atomus + TX, Aphelinus abdominalis + TX, Aphidius colemani + TX, Aphidoletes aphidimyza + TX, Autographa californica NPV + TX, Bacillus sphaericus Neide + TX, Beauveria brongniartii + TX, Chrysoperla carnea + TX, Cryptolaemus montrouzieri + TX, Cydia pomonella GV + TX, Dacnusa sibirica + TX, Diglyphus isaea + TX, Encarsia formosa + TX, Eretmocerus eremicus + TX, Heterorhabditis bacteriophora and H. megidis + TX, Hippodamia convergens + TX, Leptomastix
- 20 dactylopii + TX, Macrolophus caliginosus + TX, Mamestra brassicae NPV + TX, Metaphycus helvolus + TX, Metarhizium anisopliae var. acridum + TX, Metarhizium anisopliae var. anisopliae + TX, Neodiprion sertifer NPV and N. lecontei NPV + TX, Orius spp. + TX, Paecilomyces fumosoroseus + TX, Phytoseiulus persimilis + TX, Steinernema bibionis + TX, Steinernema carpocapsae + TX, Steinernema feltiae + TX, Steinernema glaseri + TX, Steinernema riobrave + TX, Steinernema
- 25 riobravis + TX, Steinernema scapterisci + TX, Steinernema spp. + TX, Trichogramma spp. + TX, Typhlodromus occidentalis + TX, Verticillium lecanii + TX, apholate + TX, bisazir + TX, busulfan + TX, dimatif + TX, hemel + TX, hempa + TX, metepa + TX, methiotepa + TX, methyl apholate + TX, morzid + TX, penfluron + TX, tepa + TX, thiohempa + TX, thiotepa + TX, tretamine + TX, uredepa + TX, (E)-dec-5-en-1-yl acetate with (E)-dec-5-en-1-ol + TX, (E)-tridec-4-en-1-yl acetate + TX, (E)-6-methylhept-
- 30 2-en-4-ol + TX, (E,Z)-tetradeca-4,10-dien-1-yl acetate + TX, (Z)-dodec-7-en-1-yl acetate + TX, (Z)-hexadec-11-enal + TX, (Z)-hexadec-11-en-1-yl acetate + TX, (Z)-hexadec-13-en-11-yn-1-yl acetate + TX, (Z)-icos-13-en-10-one + TX, (Z)-tetradec-7-en-1-al + TX, (Z)-tetradec-9-en-1-ol + TX, (Z)-tetradec-9-en-1-yl acetate + TX, (7E,9Z)-dodeca-7,9-dien-1-yl acetate + TX, (9Z,11E)-tetradeca-9,11-dien-1-yl acetate + TX, (9Z,12E)-tetradeca-9,12-dien-1-yl acetate + TX, 14-methyloctadec-1-ene + TX, 4-
- 35 methylnonan-5-ol with 4-methylnonan-5-one + TX, alpha-multistriatin + TX, brevicomin + TX, codlelure + TX, codlemone + TX, cuelure + TX, disparlure + TX, dodec-8-en-1-yl acetate + TX, dodec-9-en-1-yl acetate + TX, dodeca-8 + TX, 10-dien-1-yl acetate + TX, dominicalure + TX, ethyl 4-methyloctanoate + TX, eugenol + TX, frontaline + TX, grandlure + TX, grandlure I + TX, grandlure II + TX, grandlure III + TX, grandlure IV + TX, hexalure + TX, ipsdienol + TX, ipsenol + TX, japonilure + TX, lineatin + TX,

litlure + TX, looplure + TX, medlure + TX, megatomoic acid + TX, methyl eugenol + TX, muscalure + TX, octadeca-2,13-dien-1-yl acetate + TX, octadeca-3,13-dien-1-yl acetate + TX, orfralure + TX, oryctalure + TX, ostramone + TX, siglure + TX, sordidin + TX, sulcatol + TX, tetradec-11-en-1-yl acetate + TX, trimedlure + TX, trimedlure A + TX, trimedlure B₁ + TX, trimedlure B₂ + TX, trimedlure C + TX, trunc-call + TX, 2-(octylthio)-ethanol + TX, butopyronoxyl + TX, butoxy(polypropylene glycol) + TX, dibutyl adipate + TX, dibutyl phthalate + TX, dibutyl succinate + TX, diethyltoluamide + TX, dimethyl carbate + TX, dimethyl phthalate + TX, ethyl hexanediol + TX, hexamide + TX, methoquin-butyl + TX, methylneodecanamide + TX, oxamate + TX, picaridin + TX, 1-dichloro-1-nitroethane + TX, 1,1-dichloro-2,2-bis(4-ethylphenyl)-ethane + TX, 1,2-dichloropropane with 1,3-dichloropropene + TX, 1-bromo-2-chloroethane + TX, 2,2,2-trichloro-1-(3,4-dichloro-phenyl)ethyl acetate + TX, 2,2-dichlorovinyl 2-ethylsulfinyethyl methyl phosphate + TX, 2-(1,3-dithiolan-2-yl)phenyl dimethylcarbamate + TX, 2-(2-butoxyethoxy)ethyl thiocyanate + TX, 2-(4,5-dimethyl-1,3-dioxolan-2-yl)phenyl methylcarbamate + TX, 2-(4-chloro-3,5-xylyloxy)ethanol + TX, 2-chlorovinyl diethyl phosphate + TX, 2-imidazolidone + TX, 2-isovalerylindan-1,3-dione + TX, 2-methyl(prop-2-ynyl)aminophenyl methylcarbamate + TX, 2-thiocyanatoethyl laurate + TX, 3-bromo-1-chloroprop-1-ene + TX, 3-methyl-1-phenylpyrazol-5-yl dimethyl-carbamate + TX, 4-methyl(prop-2-ynyl)amino-3,5-xylyl methylcarbamate + TX, 5,5-dimethyl-3-oxocyclohex-1-enyl dimethylcarbamate + TX, acethion + TX, acrylonitrile + TX, aldrin + TX, allosamidin + TX, allyxycarb + TX, alpha-ecdysone + TX, aluminium phosphide + TX, aminocarb + TX, anabasine + TX, athidathion + TX, azamethiphos + TX, *Bacillus thuringiensis* delta endotoxins + TX, barium hexafluorosilicate + TX, barium polysulfide + TX, barthrin + TX, Bayer 22/190 + TX, Bayer 22408 + TX, beta-cyfluthrin + TX, beta-cypermethrin + TX, bioethanomethrin + TX, biopermethrin + TX, bis(2-chloroethyl) ether + TX, borax + TX, bromfenvinfos + TX, bromo-DDT + TX, bufencarb + TX, butacarb + TX, butathiofos + TX, butonate + TX, calcium arsenate + TX, calcium cyanide + TX, carbon disulfide + TX, carbon tetrachloride + TX, cartap hydrochloride + TX, cevadine + TX, chlorbicyclen + TX, chlordane + TX, chlordecone + TX, chloroform + TX, chloropicrin + TX, chlorphoxim + TX, chlorprazophos + TX, cis-resmethrin + TX, cismethrin + TX, clocythrin + TX, copper acetoarsenite + TX, copper arsenate + TX, copper oleate + TX, coumithoate + TX, cryolite + TX, CS 708 + TX, cyanofenphos + TX, cyanophos + TX, cyclethrin + TX, cythioate + TX, d-tetramethrin + TX, DAEP + TX, dazomet + TX, decarbofuran + TX, diamidafos + TX, dicapthion + TX, dichlofenthion + TX, dicresyl + TX, dicyclanil + TX, dieldrin + TX, diethyl 5-methylpyrazol-3-yl phosphate + TX, dilor + TX, dimefluthrin + TX, dimetan + TX, dimethrin + TX, dimethylvinphos + TX, dimetilan + TX, dinoprop + TX, dinosam + TX, dinoseb + TX, diofenolan + TX, dioxabenzofos + TX, dithicrofos + TX, DSP + TX, ecdysterone + TX, EI 1642 + TX, EMPC + TX, EPBP + TX, etaphos + TX, ethiofencarb + TX, ethyl formate + TX, ethylene dibromide + TX, ethylene dichloride + TX, ethylene oxide + TX, EXD + TX, fenchlorphos + TX, fenethacarb + TX, fenitrothion + TX, fenoxacrim + TX, fenpirithrin + TX, fensulfothion + TX, fenthion-ethyl + TX, flucufuron + TX, fosmethilan + TX, fospirate + TX, fosthietan + TX, furathiocarb + TX, furethrin + TX, guazatine + TX, guazatine acetates + TX, sodium tetrathiocarbonate + TX, halfenprox + TX, HCH + TX, HEOD + TX, heptachlor + TX, heterophos + TX, HHDN + TX, hydrogen cyanide + TX, hyquincarb + TX, IPSP + TX,

isazofos + TX, isobenzan + TX, isodrin + TX, isofenphos + TX, isolane + TX, isoprothiolane + TX, isoxathion + TX, juvenile hormone I + TX, juvenile hormone II + TX, juvenile hormone III + TX, kelevan + TX, kinoprene + TX, lead arsenate + TX, leptophos + TX, lirimfos + TX, lythidathion + TX, m-cumenyl methylcarbamate + TX, magnesium phosphide + TX, mazidox + TX, mecarphon + TX, menazon + TX, mercurous chloride + TX, mesulfenfos + TX, metam + TX, metam-potassium + TX, metam-sodium + TX, methanesulfonyl fluoride + TX, methocrotophos + TX, methoprene + TX, methothrin + TX, methoxychlor + TX, methyl isothiocyanate + TX, methylchloroform + TX, methylene chloride + TX, metoxadiazone + TX, mirex + TX, naftalofos + TX, naphthalene + TX, NC-170 + TX, nicotine + TX, nicotine sulfate + TX, nithiazine + TX, normicotine + TX, O-5-dichloro-4-iodophenyl O-ethyl ethylphosphonothioate + TX, O,O-diethyl O-4-methyl-2-oxo-2H-chromen-7-yl phosphorothioate + TX, O,O-diethyl O-6-methyl-2-propylpyrimidin-4-yl phosphorothioate + TX, O,O,O',O'-tetrapropyl dithiopyrophosphate + TX, oleic acid + TX, para-dichlorobenzene + TX, parathion-methyl + TX, pentachlorophenol + TX, pentachlorophenyl laurate + TX, PH 60-38 + TX, phenkapton + TX, phosnichlor + TX, phosphine + TX, phoxim-methyl + TX, pirimetaphos + TX, polychlorodicyclopentadiene isomers + TX, potassium arsenite + TX, potassium thiocyanate + TX, precocene I + TX, precocene II + TX, precocene III + TX, primidophos + TX, profluthrin + TX, promecarb + TX, prothiofos + TX, pyrazophos + TX, pyresmethrin + TX, quassia + TX, quinalphos-methyl + TX, quinothion + TX, rafoxanide + TX, resmethrin + TX, rotenone + TX, kadethrin + TX, ryania + TX, ryanodine + TX, sabadilla) + TX, schradan + TX, sebufos + TX, SI-0009 + TX, thiapronil + TX, sodium arsenite + TX, sodium cyanide + TX, sodium fluoride + TX, sodium hexafluorosilicate + TX, sodium pentachlorophenoxide + TX, sodium selenate + TX, sodium thiocyanate + TX, sulcofuron + TX, sulcofuron-sodium + TX, sulfuryl fluoride + TX, sulprofos + TX, tar oils + TX, tazimcarb + TX, TDE + TX, tebupirimfos + TX, temephos + TX, terallethrin + TX, tetrachloroethane + TX, thicrofos + TX, thiocyclam + TX, thiocyclam hydrogen oxalate + TX, thionazin + TX, thiosultap + TX, thiosultap-sodium + TX, tralomethrin + TX, transpermethrin + TX, triazamate + TX, trichlormetaphos-3 + TX, trichloronat + TX, trimethacarb + TX, tolprocarb + TX, triclopyricarb + TX, triprene + TX, veratridine + TX, veratrine + TX, XMC + TX, zetamethrin + TX, zinc phosphide + TX, zolaprofos + TX, and meperfluthrin + TX, tetramethylfluthrin + TX, bis(tributyltin) oxide + TX, bromoacetamide + TX, ferric phosphate + TX, niclosamide-olamine + TX, tributyltin oxide + TX, pyrimorph + TX, trifenmorph + TX, 1,2-dibromo-3-chloropropane + TX, 1,3-dichloropropene + TX, 3,4-dichlorotetrahydrothio-phene 1,1-dioxide + TX, 3-(4-chlorophenyl)-5-methylrhodanine + TX, 5-methyl-6-thioxo-1,3,5-thiadiazinan-3-ylacetic acid + TX, 6-isopentenylaminopurine + TX, 2-fluoro-N-(3-methoxyphenyl)-9H-purin-6-amine + TX, benclotiaz + TX, cytokinins + TX, DCIP + TX, furfural + TX, isamidofos + TX, kinetin + TX, Myrothecium verrucaria composition + TX, tetrachlorothiophene + TX, xlenols + TX, zeatin + TX, potassium ethylxanthate + TX, acibenzolar + TX, acibenzolar-S-methyl + TX, Reynoutria sachalinensis extract + TX, alpha-chlorohydrin + TX, antu + TX, barium carbonate + TX, bithiosemi + TX, brodifacoum + TX, bromadiolone + TX, bromethalin + TX, chlorophacinone + TX, cholecalciferol + TX, coumachlor + TX, coumafuryl + TX, coumatetralyl + TX, crimidine + TX, difenacoum + TX, difethialone + TX, diphacinone + TX, ergocalciferol + TX, flocoumafen + TX, fluoroacetamide + TX, flupropadine +

TX, flupropadine hydrochloride + TX, norbormide + TX, phosacetim + TX, phosphorus + TX, pindone + TX, pyrinuron + TX, scilliroside + TX, -sodium fluoroacetate + TX, thallium sulfate + TX, warfarin + TX, -2-(2-butoxyethoxy)ethyl piperonylate + TX, 5-(1,3-benzodioxol-5-yl)-3-hexylcyclohex-2-enone + TX, farnesol with nerolidol + TX, verbutin + TX, MGK 264 + TX, piperonyl butoxide + TX, piprotal + TX,

5 propyl isomer + TX, S421 + TX, sesamex + TX, sesasmolin + TX, sulfoxide + TX, anthraquinone + TX, copper naphthenate + TX, copper oxychloride + TX, dicyclopentadiene + TX, thiram + TX, zinc naphthenate + TX, ziram + TX, imanin + TX, ribavirin + TX, mercuric oxide + TX, thiophanate-methyl + TX, azaconazole + TX, bitertanol + TX, bromuconazole + TX, cyproconazole + TX, difenoconazole + TX, diniconazole -+ TX, epoxiconazole + TX, fenbuconazole + TX, fluquinconazole + TX, flusilazole +

10 TX, flutriafol + TX, furametpyr + TX, hexaconazole + TX, imazalil- + TX, imiben-conazole + TX, ipconazole + TX, metconazole + TX, myclobutanil + TX, paclobutrazole + TX, pefurazoate + TX, penconazole + TX, prothioconazole + TX, pyrifenoxy + TX, prochloraz + TX, propiconazole + TX, pyrisoxazole + TX, -simeconazole + TX, tebuconazole + TX, tetraconazole + TX, triadimefon + TX, triadimenol + TX, triflumizole + TX, triticonazole + TX, ancymidol + TX, fenarimol + TX, nuarimol + TX,

15 bupirimate + TX, dimethirimol + TX, ethirimol + TX, dodemorph + TX, fenpropidine + TX, fenpropimorph + TX, spiroxamine + TX, tridemorph + TX, cyprodinil + TX, mepanipyrim + TX, pyrimethanil + TX, fenpiclonil + TX, fludioxonil + TX, benalaxyl + TX, furalaxyl + TX, -metalaxyl -+ TX, Rmetalaxyl + TX, ofurace + TX, oxadixyl + TX, carbendazim + TX, debacarb + TX, fuberidazole -+ TX, thiabendazole + TX, chlozolate + TX, dichlozoline + TX, myclozoline- + TX, procymidone + TX,

20 vinclozoline + TX, boscalid + TX, carboxin + TX, fenfuram + TX, flutolanil + TX, mepronil + TX, oxycarboxin + TX, penthiopyrad + TX, thifluzamide + TX, dodine + TX, iminoctadine + TX, azoxystrobin + TX, dimoxystrobin + TX, enestroburin + TX, fenaminostrobin + TX, flufenoxystrobin + TX, fluoxastrobin + TX, kresoxim--methyl + TX, metominostrobin + TX, trifloxystrobin + TX, orysastrobin + TX, picoxystrobin + TX, pyraclostrobin + TX, pyrametostrobin + TX, pyraoxystrobin +

25 TX, ferbam + TX, mancozeb + TX, maneb + TX, metiram + TX, propineb + TX, zineb + TX, captafol + TX, captan + TX, fluoroimide + TX, folpet + TX, tolylfluanid + TX, bordeaux mixture + TX, copper oxide + TX, mancozeb + TX, oxine-copper + TX, nitrothal-isopropyl + TX, edifenphos + TX, iprobenphos + TX, phosdiphen + TX, tolclofos-methyl + TX, anilazine + TX, benthiavalicarb + TX, blasticidin-S + TX, chloroneb -+ TX, chloro-tha-lonil + TX, cyflufenamid + TX, cymoxanil + TX, cyclobutrifluram + TX,

30 diclocymet + TX, diclomezine -+ TX, dicloran + TX, diethofencarb + TX, dimethomorph -+ TX, flumorph + TX, dithianon + TX, ethaboxam + TX, etridiazole + TX, famoxadone + TX, fenamidone + TX, fenoxanil + TX, ferimzone + TX, fluazinam + TX, fluopicolide + TX, flusulfamide + TX, fluxapyroxad + TX, -fenhexamid + TX, fosetyl-aluminium -+ TX, hymexazol + TX, iprovalicarb + TX, cyazofamid + TX, methasulfocarb + TX, metrafenone + TX, pencycuron + TX, phthalide + TX,

35 polyoxins + TX, propamocarb + TX, pyribencarb + TX, proquinazid + TX, pyroquilon + TX, pyriofenone + TX, quinoxifen + TX, quintozone + TX, tiadinil + TX, triazoxide + TX, tricyclazole + TX, triforine + TX, validamycin + TX, valifenalate + TX, zoxamide + TX, mandipropamid + TX, flubeneteram + TX, isopyrazam + TX, sedaxane + TX, benzovindiflupyr + TX, pydiflumetofen + TX, 3-difluoromethyl-1-methyl-1H-pyrazole-4-carboxylic acid (3',4',5'-trifluoro-biphenyl-2-yl)-amide + TX, isoflucypram + TX,

isotianil + TX, dipymetitrone + TX, 6-ethyl-5,7-dioxo-pyrrolo[4,5][1,4]dithiino[1,2-c]isothiazole-3-carbonitrile + TX, 2-(difluoromethyl)-N-[3-ethyl-1,1-dimethyl-indan-4-yl]pyridine-3-carboxamide + TX, 4-(2,6-difluorophenyl)-6-methyl-5-phenyl-pyridazine-3-carbonitrile + TX, (R)-3-(difluoromethyl)-1-methyl-N-[1,1,3-trimethylindan-4-yl]pyrazole-4-carboxamide + TX, 4-(2-bromo-4-fluoro-phenyl)-N-(2-chloro-6-fluoro-phenyl)-2,5-dimethyl-pyrazol-3-amine + TX, 4-(2-bromo-4-fluorophenyl)-N-(2-chloro-6-fluorophenyl)-1,3-dimethyl-1H-pyrazol-5-amine + TX, fluindapyr + TX, coumethoxystrobin (jiaxiangjunzhi) + TX, lvbenmixianan + TX, dichlobentiazox + TX, mandestrobin + TX, 3-(4,4-difluoro-3,4-dihydro-3,3-dimethylisoquinolin-1-yl)quinolone + TX, 2-[2-fluoro-6-[(8-fluoro-2-methyl-3-quinolyl)oxy]phenyl]propan-2-ol + TX, oxathiapirolin + TX, tert-butyl N-[6-[[[(1-methyltetrazol-5-yl)-phenyl-methylene]amino]oxymethyl]-2-pyridyl]carbamate + TX, pyraziflumid + TX, inpyrfluxam + TX, troprocarb + TX, mefentrifluconazole + TX, ipfentrifluconazole + TX, 2-(difluoromethyl)-N-[(3R)-3-ethyl-1,1-dimethyl-indan-4-yl]pyridine-3-carboxamide + TX, N'-(2,5-dimethyl-4-phenoxy-phenyl)-N-ethyl-N-methyl-formamidine + TX, N'-[4-(4,5-dichlorothiazol-2-yl)oxy-2,5-dimethyl-phenyl]-N-ethyl-N-methyl-formamidine + TX, [2-[3-[2-[1-[2-[3,5-bis(difluoromethyl)pyrazol-1-yl]acetyl]-4-piperidyl]thiazol-4-yl]-4,5-dihydroisoxazol-5-yl]-3-chloro-phenyl]methanesulfonate + TX, but-3-ynyl N-[6-[[[(Z)-[(1-methyltetrazol-5-yl)-phenyl-methylene]amino]oxymethyl]-2-pyridyl]carbamate + TX, methyl N-[5-[4-(2,4-dimethylphenyl)triazol-2-yl]-2-methyl-phenyl]methyl]carbamate + TX, 3-chloro-6-methyl-5-phenyl-4-(2,4,6-trifluorophenyl)pyridazine + TX, pyridachlometyl + TX, 3-(difluoromethyl)-1-methyl-N-[1,1,3-trimethylindan-4-yl]pyrazole-4-carboxamide + TX, 1-[2-[1-(4-chlorophenyl)pyrazol-3-yl]oxymethyl]-3-methyl-phenyl]-4-methyl-tetrazol-5-one + TX, 1-methyl-4-[3-methyl-2-[2-methyl-4-(3,4,5-trimethylpyrazol-1-yl)phenoxy]methyl]phenyl]tetrazol-5-one + TX, aminopyrifin + TX, ametocradin + TX, amisulbrom + TX, penflufen + TX, (Z,2E)-5-[1-(4-chlorophenyl)pyrazol-3-yl]oxy-2-methoxyimino-N,3-dimethyl-pent-3-enamide + TX, florylpicoxamid + TX, fempicoxamid + TX, tebufloquin + TX, ipflufenquin + TX, quinfumelin + TX, isofetamid + TX, N-[2-[2,4-dichlorophenoxy]phenyl]-3-(difluoromethyl)-1-methyl-pyrazole-4-carboxamide + TX, N-[2-[2-chloro-4-(trifluoromethyl)phenoxy]phenyl]-3-(difluoromethyl)-1-methyl-pyrazole-4-carboxamide + TX, benzothiostrubin + TX, phenamacril + TX, 5-amino-1,3,4-thiadiazole-2-thiol zinc salt (2:1) + TX, fluopyram + TX, flutianil + TX, fluopimomide + TX, pyrapropoyne + TX, picarbutrazox + TX, 2-(difluoromethyl)-N-(3-ethyl-1,1-dimethyl-indan-4-yl)pyridine-3-carboxamide + TX, 2-(difluoromethyl)-N-((3R)-1,1,3-trimethylindan-4-yl)pyridine-3-carboxamide + TX, 4-[[6-[2-(2,4-difluorophenyl)-1,1-difluoro-2-hydroxy-3-(1,2,4-triazol-1-yl)propyl]-3-pyridyl]oxy]benzonitrile + TX, metyltetraprole + TX, 2-(difluoromethyl)-N-((3R)-1,1,3-trimethylindan-4-yl)pyridine-3-carboxamide + TX, α -(1,1-dimethylethyl)- α -[4'-(trifluoromethoxy)[1,1'-biphenyl]-4-yl]-5-pyrimidinemethanol + TX, fluoxapirolin + TX, enoxastrobin + TX, 4-[[6-[2-(2,4-difluorophenyl)-1,1-difluoro-2-hydroxy-3-(1,2,4-triazol-1-yl)propyl]-3-pyridyl]oxy]benzonitrile + TX, 4-[[6-[2-(2,4-difluorophenyl)-1,1-difluoro-2-hydroxy-3-(5-sulfanyl-1,2,4-triazol-1-yl)propyl]-3-pyridyl]oxy]benzonitrile + TX, 4-[[6-[2-(2,4-difluorophenyl)-1,1-difluoro-2-hydroxy-3-(5-thioxo-4H-1,2,4-triazol-1-yl)propyl]-3-pyridyl]oxy]benzonitrile + TX, trinexapac + TX, coumethoxystrobin + TX, zhongshengmycin + TX, thiodiazole copper + TX, zinc thiazole + TX, amectotractin + TX, iprodione + TX; N'-[5-bromo-2-methyl-6-[(1S)-1-methyl-2-propoxy-ethoxy]-3-

pyridyl]-N-ethyl-N-methyl-formamidine + TX, N'-[5-bromo-2-methyl-6-[(1R)-1-methyl-2-propoxy-ethoxy]-3-pyridyl]-N-ethyl-N-methyl-formamidine + TX, N'-[5-bromo-2-methyl-6-(1-methyl-2-propoxy-ethoxy)-3-pyridyl]-N-ethyl-N-methyl-formamidine + TX, N'-[5-chloro-2-methyl-6-(1-methyl-2-propoxy-ethoxy)-3-pyridyl]-N-ethyl-N-methyl-formamidine + TX, N'-[5-bromo-2-methyl-6-(1-methyl-2-propoxy-ethoxy)-3-pyridyl]-N-isopropyl-N-methyl-formamidine + TX (these compounds may be prepared from the methods described in WO2015/155075); N'-[5-bromo-2-methyl-6-(2-propoxypropoxy)-3-pyridyl]-N-ethyl-N-methyl-formamidine + TX (this compound may be prepared from the methods described in IPCOM000249876D); N-isopropyl-N'-[5-methoxy-2-methyl-4-(2,2,2-trifluoro-1-hydroxy-1-phenylethyl)phenyl]-N-methyl-formamidine + TX, N'-[4-(1-cyclopropyl-2,2,2-trifluoro-1-hydroxy-ethyl)-5-methoxy-2-methyl-phenyl]-N-isopropyl-N-methyl-formamidine + TX (these compounds may be prepared from the methods described in WO2018/228896); N-ethyl-N'-[5-methoxy-2-methyl-4-[2-trifluoromethyl]oxetan-2-yl]phenyl]-N-methyl-formamidine + TX, N-ethyl-N'-[5-methoxy-2-methyl-4-[2-trifluoromethyl]tetrahydrofuran-2-yl]phenyl]-N-methyl-formamidine + TX (these compounds may be prepared from the methods described in WO2019/110427); N-[(1R)-1-benzyl-3-chloro-1-methyl-but-3-enyl]-8-fluoro-quinoline-3-carboxamide + TX, N-[(1S)-1-benzyl-3,3,3-trifluoro-1-methyl-propyl]-8-fluoro-quinoline-3-carboxamide + TX, N-[(1S)-1-benzyl-1,3-dimethyl-butyl]-7,8-difluoro-quinoline-3-carboxamide + TX, 8-fluoro-N-[1-[(3-fluorophenyl)methyl]-1,3-dimethyl-butyl]quinoline-3-carboxamide + TX, N-(1-benzyl-1,3-dimethyl-butyl)-8-fluoro-quinoline-3-carboxamide + TX, N-[(1R)-1-benzyl-1,3-dimethyl-butyl]-8-fluoro-quinoline-3-carboxamide + TX, N-[(1S)-1-benzyl-1,3-dimethyl-butyl]-8-fluoro-quinoline-3-carboxamide + TX, N-(1-benzyl-3-chloro-1-methyl-but-3-enyl)-8-fluoro-quinoline-3-carboxamide + TX (these compounds may be prepared from the methods described in WO2017/153380); 1-(6,7-dimethylpyrazolo[1,5-a]pyridin-3-yl)-4,4,5-trifluoro-3,3-dimethyl-isoquinoline + TX, 1-(6,7-dimethylpyrazolo[1,5-a]pyridin-3-yl)-4,4,6-trifluoro-3,3-dimethyl-isoquinoline + TX, 4,4-difluoro-3,3-dimethyl-1-(6-methylpyrazolo[1,5-a]pyridin-3-yl)isoquinoline + TX, 4,4-difluoro-3,3-dimethyl-1-(7-methylpyrazolo[1,5-a]pyridin-3-yl)isoquinoline + TX, 1-(6-chloro-7-methyl-pyrazolo[1,5-a]pyridin-3-yl)-4,4-difluoro-3,3-dimethyl-isoquinoline + TX (these compounds may be prepared from the methods described in WO2017/025510); 1-(4,5-dimethylbenzimidazol-1-yl)-4,4,5-trifluoro-3,3-dimethyl-isoquinoline + TX, 1-(4,5-dimethylbenzimidazol-1-yl)-4,4-difluoro-3,3-dimethyl-isoquinoline + TX, 6-chloro-4,4-difluoro-3,3-dimethyl-1-(4-methylbenzimidazol-1-yl)isoquinoline + TX, 4,4-difluoro-1-(5-fluoro-4-methyl-benzimidazol-1-yl)-3,3-dimethyl-isoquinoline + TX, 3-(4,4-difluoro-3,3-dimethyl-1-isoquinolyl)-7,8-dihydro-6H-cyclopenta[e]benzimidazole + TX (these compounds may be prepared from the methods described in WO2016/156085); N-methoxy-N-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]cyclopropanecarboxamide + TX, N,2-dimethoxy-N-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]propanamide + TX, N-ethyl-2-methyl-N-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]propanamide + TX, 1-methoxy-3-methyl-1-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]urea + TX, 1,3-dimethoxy-1-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]urea + TX, 3-ethyl-1-methoxy-1-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]urea + TX, N-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]propanamide + TX, 4,4-dimethyl-2-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]isoxazolidin-3-one +

TX, 5,5-dimethyl-2-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]isoxazolidin-3-one + TX, ethyl 1-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]pyrazole-4-carboxylate + TX, N,N-dimethyl-1-[[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methyl]-1,2,4-triazol-3-amine + TX. The compounds in this paragraph may be prepared from the methods described in WO 2017/055473, WO 5 2017/055469, WO 2017/093348 and WO 2017/118689; 2-[6-(4-chlorophenoxy)-2-(trifluoromethyl)-3-pyridyl]-1-(1,2,4-triazol-1-yl)propan-2-ol + TX (this compound may be prepared from the methods described in WO 2017/029179); 2-[6-(4-bromophenoxy)-2-(trifluoromethyl)-3-pyridyl]-1-(1,2,4-triazol-1-yl)propan-2-ol + TX (this compound may be prepared from the methods described in WO 2017/029179); 3-[2-(1-chlorocyclopropyl)-3-(2-fluorophenyl)-2-hydroxy-propyl]imidazole-4-carbonitrile 10 + TX (this compound may be prepared from the methods described in WO 2016/156290); 3-[2-(1-chlorocyclopropyl)-3-(3-chloro-2-fluoro-phenyl)-2-hydroxy-propyl]imidazole-4-carbonitrile + TX (this compound may be prepared from the methods described in WO 2016/156290); (4-phenoxyphenyl)methyl 2-amino-6-methyl-pyridine-3-carboxylate + TX (this compound may be prepared from the methods described in WO 2014/006945); 2,6-Dimethyl-1H,5H-[1,4]dithiino[2,3-c:5,6-c']dipyrrole-1,3,5,7(2H,6H)-tetrone + TX (this compound may be prepared from the methods 15 described in WO 2011/138281); N-methyl-4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]benzenecarbothioamide + TX; N-methyl-4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]benzamide + TX; (Z,E)-5-[1-(2,4-dichlorophenyl)pyrazol-3-yl]oxy-2-methoxyimino-N,3-dimethyl-pent-3-enamide + TX (this compound may be prepared from the methods described in WO 2018/153707); N'-(2-chloro-5-methyl-4-phenoxy-phenyl)-N-ethyl-N-methyl-formamidine + TX; N'-[2-chloro-4-(2-fluorophenoxy)-5-methyl-phenyl]-N-ethyl-N-methyl-formamidine + TX (this compound may be prepared from the 20 methods described in WO 2016/202742); 2-(difluoromethyl)-N-[(3S)-3-ethyl-1,1-dimethyl-indan-4-yl]pyridine-3-carboxamide + TX (this compound may be prepared from the methods described in WO 2014/095675); (5-methyl-2-pyridyl)-[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methanone + TX, (3-methylisoxazol-5-yl)-[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]methanone + TX (these 25 compounds may be prepared from the methods described in WO 2017/220485); 2-oxo-N-propyl-2-[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]acetamide + TX (this compound may be prepared from the methods described in WO 2018/065414); ethyl 1-[[5-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]-2-thienyl]methyl]pyrazole-4-carboxylate + TX (this compound may be prepared from the methods 30 described in WO 2018/158365); 2,2-difluoro-N-methyl-2-[4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]phenyl]acetamide + TX, N-(E)-methoxyiminomethyl-4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]benzamide + TX, N-(Z)-methoxyiminomethyl-4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]benzamide + TX, N-[N-methoxy-C-methyl-carbonimidoyl]-4-[5-(trifluoromethyl)-1,2,4-oxadiazol-3-yl]benzamide + TX (these compounds may be prepared from the methods described in WO 2018/202428); 35
microbials including: *Acinetobacter lwoffii* + TX, *Acremonium alternatum* + TX + TX, *Acremonium cephalosporium* + TX + TX, *Acremonium diospyri* + TX, *Acremonium obclavatum* + TX, *Adoxophyes orana granulovirus* (AdoxGV) (Capex®) + TX, *Agrobacterium radiobacter* strain K84 (Galltrol-A®) + TX, *Alternaria alternate* + TX, *Alternaria cassia* + TX, *Alternaria destruens* (Smolder®) + TX, *Ampelomyces quisqualis* (AQ10®) + TX, *Aspergillus flavus* AF36 (AF36®) + TX, *Aspergillus flavus*

- NRRL 21882 (Aflaguard®) + TX, *Aspergillus* spp. + TX, *Aureobasidium pullulans* + TX, *Azospirillum* + TX, (MicroAZ® + TX, TAZO B®) + TX, *Azotobacter* + TX, *Azotobacter chroococcum* (Azotomeal®) + TX, *Azotobacter* cysts (Bionatural Blooming Blossoms®) + TX, *Bacillus amyloliquefaciens* + TX, *Bacillus cereus* + TX, *Bacillus chitinosporus* strain CM-1 + TX, *Bacillus chitinosporus* strain AQ746 + TX, *Bacillus licheniformis* strain HB-2 (Biostart™ Rhizoboost®) + TX, *Bacillus licheniformis* strain 3086 (EcoGuard® + TX, Green Releaf®) + TX, *Bacillus circulans* + TX, *Bacillus firmus* (BioSafe® + TX, BioNem-WP® + TX, VOTiVO®) + TX, *Bacillus firmus* strain I-1582 + TX, *Bacillus macerans* + TX, *Bacillus marismortui* + TX, *Bacillus megaterium* + TX, *Bacillus mycoides* strain AQ726 + TX, *Bacillus papillae* (Milky Spore Powder®) + TX, *Bacillus pumilus* spp. + TX, *Bacillus pumilus* strain GB34 (Yield Shield®) + TX, *Bacillus pumilus* strain AQ717 + TX, *Bacillus pumilus* strain QST 2808 (Sonata® + TX, Ballad Plus®) + TX, *Bacillus spahericus* (VectoLex®) + TX, *Bacillus* spp. + TX, *Bacillus* spp. strain AQ175 + TX, *Bacillus* spp. strain AQ177 + TX, *Bacillus* spp. strain AQ178 + TX, *Bacillus subtilis* strain QST 713 (CEASE® + TX, Serenade® + TX, Rhapsody®) + TX, *Bacillus subtilis* strain QST 714 (JAZZ®) + TX, *Bacillus subtilis* strain AQ153 + TX, *Bacillus subtilis* strain AQ743 + TX, *Bacillus subtilis* strain QST3002 + TX, *Bacillus subtilis* strain QST3004 + TX, *Bacillus subtilis* var. *amyloliquefaciens* strain FZB24 (Taegro® + TX, Rhizopro®) + TX, *Bacillus thuringiensis* Cry 2Ae + TX, *Bacillus thuringiensis* Cry1Ab + TX, *Bacillus thuringiensis aizawai* GC 91 (Agree®) + TX, *Bacillus thuringiensis israelensis* (BMP123® + TX, Aquabac® + TX, VectoBac®) + TX, *Bacillus thuringiensis kurstaki* (Javelin® + TX, Deliver® + TX, CryMax® + TX, Bonide® + TX, Scutella WP® + TX, Turilav WP® + TX, Astuto® + TX, Dipel WP® + TX, Biobit® + TX, Foray®) + TX, *Bacillus thuringiensis kurstaki* BMP 123 (Baritone®) + TX, *Bacillus thuringiensis kurstaki* HD-1 (Bioprotec-CAF / 3P®) + TX, *Bacillus thuringiensis* strain BD#32 + TX, *Bacillus thuringiensis* strain AQ52 + TX, *Bacillus thuringiensis* var. *aizawai* (XenTari® + TX, DiPel®) + TX, bacteria spp. (GROWMEND® + TX, GROWSWEET® + TX, Shootup®) + TX, bacteriophage of *Clavipacter michiganensis* (AgriPhage®) + TX, Bakflor® + TX, *Beauveria bassiana* (Beaugenic® + TX, Brocaril WP®) + TX, *Beauveria bassiana* GHA (Mycotrol ES® + TX, Mycotrol O® + TX, BotaniGuard®) + TX, *Beauveria brongniartii* (Engerlingspilz® + TX, Schweizer Beauveria® + TX, Melocont®) + TX, *Beauveria* spp. + TX, *Botrytis cineria* + TX, *Bradyrhizobium japonicum* (TerraMax®) + TX, *Brevibacillus brevis* + TX, *Bacillus thuringiensis tenebrionis* (Novodor®) + TX, BtBooster + TX, *Burkholderia cepacia* (Deny® + TX, Intercept® + TX, Blue Circle®) + TX, *Burkholderia gladii* + TX, *Burkholderia gladioli* + TX, *Burkholderia* spp. + TX, Canadian thistle fungus (CBH Canadian Bioherbicide®) + TX, *Candida butyri* + TX, *Candida famata* + TX, *Candida fructus* + TX, *Candida glabrata* + TX, *Candida guilliermondii* + TX, *Candida melibiosica* + TX, *Candida oleophila* strain O + TX, *Candida parapsilosis* + TX, *Candida pelliculosa* + TX, *Candida pulcherrima* + TX, *Candida reukaufii* + TX, *Candida saitoana* (Bio-Coat® + TX, Biocure®) + TX, *Candida sake* + TX, *Candida* spp. + TX, *Candida tenuis* + TX, *Cedecea dravisae* + TX, *Cellulomonas flavigena* + TX, *Chaetomium cochliodes* (Nova-Cide®) + TX, *Chaetomium globosum* (Nova-Cide®) + TX, *Chromobacterium subtsugae* strain PR4A4-1T (Grandevo®) + TX, *Cladosporium cladosporioides* + TX, *Cladosporium oxysporum* + TX, *Cladosporium chlorocephalum* + TX, *Cladosporium* spp. + TX, *Cladosporium tenuissimum* + TX, *Clonostachys rosea* (EndoFine®) + TX, *Colletotrichum acutatum* +

TX, *Coniothyrium minitans* (Cotans WG®) + TX, *Coniothyrium* spp. + TX, *Cryptococcus albidus* (YIELDPLUS®) + TX, *Cryptococcus humicola* + TX, *Cryptococcus infirmo-minutus* + TX, *Cryptococcus laurentii* + TX, *Cryptophlebia leucotreta granulovirus* (Cryptex®) + TX, *Cupriavidus campinensis* + TX, *Cydia pomonella granulovirus* (CYD-X®) + TX, *Cydia pomonella granulovirus* (Madex® + TX, Madex Plus® + TX, Madex Max/ Carpovirusine®) + TX, *Cylindrobasidium laeve* (Stumpout®) + TX, *Cylindrocladium* + TX, *Debaryomyces hansenii* + TX, *Drechslera hawaiiensis* + TX, *Enterobacter cloacae* + TX, *Enterobacteriaceae* + TX, *Entomophthora virulenta* (Vektor®) + TX, *Epicoccum nigrum* + TX, *Epicoccum purpurascens* + TX, *Epicoccum* spp. + TX, *Filobasidium floriforme* + TX, *Fusarium acuminatum* + TX, *Fusarium chlamydosporum* + TX, *Fusarium oxysporum* (Fusaclean® / Biofox C®) + TX, *Fusarium proliferatum* + TX, *Fusarium* spp. + TX, *Galactomyces geotrichum* + TX, *Gliocladium catenulatum* (Primastop® + TX, Prestop®) + TX, *Gliocladium roseum* + TX, *Gliocladium* spp. (SoilGard®) + TX, *Gliocladium virens* (Soilgard®) + TX, *Granulovirus* (Granupom®) + TX, *Halobacillus halophilus* + TX, *Halobacillus litoralis* + TX, *Halobacillus trueperi* + TX, *Halomonas* spp. + TX, *Halomonas subglaciescola* + TX, *Halovibrio variabilis* + TX, *Hanseniaspora uvarum* + TX, *Helicoverpa armigera nucleopolyhedrovirus* (Helicovex®) + TX, *Helicoverpa zea nuclear polyhedrosis virus* (Gemstar®) + TX, Isoflavone – formononetin (Myconate®) + TX, *Kloeckera apiculata* + TX, *Kloeckera* spp. + TX, *Lagenidium giganteum* (Laginex®) + TX, *Lecanicillium longisporum* (Vertiblast®) + TX, *Lecanicillium muscarium* (Vertikil®) + TX, *Lymantria Dispar nucleopolyhedrosis virus* (Disparvirus®) + TX, *Marinococcus halophilus* + TX, *Meira geulakonigii* + TX, *Metarhizium anisopliae* (Met52®) + TX, *Metarhizium anisopliae* (Destruxin WP®) + TX, *Metschnikowia fruticola* (Shemer®) + TX, *Metschnikowia pulcherrima* + TX, *Microdochium dimerum* (Antibot®) + TX, *Micromonospora coerulea* + TX, *Microsphaeropsis ochracea* + TX, *Muscodor albus* 620 (Muscudor®) + TX, *Muscodor roseus* strain A3-5 + TX, *Mycorrhizae* spp. (AMykor® + TX, Root Maximizer®) + TX, *Myrothecium verrucaria* strain AARC-0255 (DiTera®) + TX, BROS PLUS® + TX, *Ophiostoma piliferum* strain D97 (Sylvanex®) + TX, *Paecilomyces farinosus* + TX, *Paecilomyces fumosoroseus* (PFR-97® + TX, PreFeRa®) + TX, *Paecilomyces linacinus* (Biostat WP®) + TX, *Paecilomyces lilacinus* strain 251 (MeloCon WG®) + TX, *Paenibacillus polymyxa* + TX, *Pantoea agglomerans* (BlightBan C9-1®) + TX, *Pantoea* spp. + TX, *Pasteuria* spp. (Econem®) + TX, *Pasteuria nishizawae* + TX, *Penicillium aurantiogriseum* + TX, *Penicillium billai* (Jumpstart® + TX, TagTeam®) + TX, *Penicillium brevicompactum* + TX, *Penicillium frequentans* + TX, *Penicillium griseofulvum* + TX, *Penicillium purpurogenum* + TX, *Penicillium* spp. + TX, *Penicillium viridicatum* + TX, *Phlebiopsis gigantea* (Rotstop®) + TX, phosphate solubilizing bacteria (Phosphomeal®) + TX, *Phytophthora cryptogea* + TX, *Phytophthora palmivora* (Devine®) + TX, *Pichia anomala* + TX, *Pichia guilhermondii* + TX, *Pichia membranaefaciens* + TX, *Pichia onychis* + TX, *Pichia stipites* + TX, *Pseudomonas aeruginosa* + TX, *Pseudomonas aureofasciens* (Spot-Less Biofungicide®) + TX, *Pseudomonas cepacia* + TX, *Pseudomonas chlororaphis* (AtEze®) + TX, *Pseudomonas corrugate* + TX, *Pseudomonas fluorescens* strain A506 (BlightBan A506®) + TX, *Pseudomonas putida* + TX, *Pseudomonas reactans* + TX, *Pseudomonas* spp. + TX, *Pseudomonas syringae* (Bio-Save®) + TX, *Pseudomonas viridiflava* + TX, *Pseudomonas fluorescens* (Zequanox®) + TX, *Pseudozyma flocculosa* strain PF-A22 UL (Sporodex

-132-

- L®) + TX, *Puccinia canaliculata* + TX, *Puccinia thlaspeos* (Wood Warrior®) + TX, *Pythium paroecandrum* + TX, *Pythium oligandrum* (Polygandron® + TX, Polyversum®) + TX, *Pythium periplocum* + TX, *Rhanella aquatilis* + TX, *Rhanella* spp. + TX, *Rhizobia* (Dormal® + TX, Vault®) + TX, *Rhizoctonia* + TX, *Rhodococcus globerulus* strain AQ719 + TX, *Rhodosporidium diobovatum* + TX,
- 5 *Rhodosporidium toruloides* + TX, *Rhodotorula* spp. + TX, *Rhodotorula glutinis* + TX, *Rhodotorula graminis* + TX, *Rhodotorula mucilagnosa* + TX, *Rhodotorula rubra* + TX, *Saccharomyces cerevisiae* + TX, *Salinococcus roseus* + TX, *Sclerotinia minor* + TX, *Sclerotinia minor* (SARRITOR®) + TX, *Scytalidium* spp. + TX, *Scytalidium uredinicola* + TX, *Spodoptera exigua nuclear polyhedrosis virus* (Spod-X® + TX, Spexit®) + TX, *Serratia marcescens* + TX, *Serratia plymuthica* + TX, *Serratia* spp. +
- 10 TX, *Sordaria fimicola* + TX, *Spodoptera littoralis nucleopolyhedrovirus* (Littovir®) + TX, *Sporobolomyces roseus* + TX, *Stenotrophomonas maltophilia* + TX, *Streptomyces ahygroscopicus* + TX, *Streptomyces albaduncus* + TX, *Streptomyces exfoliates* + TX, *Streptomyces galbus* + TX, *Streptomyces griseoplanus* + TX, *Streptomyces griseoviridis* (Mycostop®) + TX, *Streptomyces lydicus* (Actinovate®) + TX, *Streptomyces lydicus* WYEC-108 (ActinoGrow®) + TX, *Streptomyces violaceus* +
- 15 TX, *Tilletiopsis minor* + TX, *Tilletiopsis* spp. + TX, *Trichoderma asperellum* (T34 Biocontrol®) + TX, *Trichoderma gamsii* (Tenet®) + TX, *Trichoderma atroviride* (Plantmate®) + TX, *Trichoderma hamatum* TH 382 + TX, *Trichoderma harzianum rifai* (Mycostar®) + TX, *Trichoderma harzianum* T-22 (Trianum-P® + TX, PlantShield HC® + TX, RootShield® + TX, Trianum-G®) + TX, *Trichoderma harzianum* T-39 (Trichodex®) + TX, *Trichoderma inhamatum* + TX, *Trichoderma koningii* + TX, *Trichoderma* spp. LC
- 20 52 (Sentinel®) + TX, *Trichoderma lignorum* + TX, *Trichoderma longibrachiatum* + TX, *Trichoderma polysporum* (Binab T®) + TX, *Trichoderma taxi* + TX, *Trichoderma virens* + TX, *Trichoderma virens* (formerly *Gliocladium virens* GL-21) (SoilGuard®) + TX, *Trichoderma viride* + TX, *Trichoderma viride* strain ICC 080 (Remedier®) + TX, *Trichosporon pullulans* + TX, *Trichosporon* spp. + TX, *Trichothecium* spp. + TX, *Trichothecium roseum* + TX, *Typhula phacorrhiza* strain 94670 + TX,
- 25 *Typhula phacorrhiza* strain 94671 + TX, *Ulocladium atrum* + TX, *Ulocladium oudemansii* (Botry-Zen®) + TX, *Ustilago maydis* + TX, various bacteria and supplementary micronutrients (Natural II®) + TX, various fungi (Millennium Microbes®) + TX, *Verticillium chlamydosporium* + TX, *Verticillium lecanii* (Mycotal® + TX, Vertalec®) + TX, Vip3Aa20 (VIPtera®) + TX, *Virgibacillus marismortui* + TX, *Xanthomonas campestris* pv. *Poae* (Camperico®) + TX, *Xenorhabdus bovienii* + TX, *Xenorhabdus*
- 30 *nematophilus*;
- Plant extracts including: pine oil (Retenol®) + TX, azadirachtin (Plasma Neem Oil® + TX, AzaGuard® + TX, MeemAzal® + TX, Molt-X® + TX, Botanical IGR (Neemazad® + TX, Neemix®) + TX, canola oil (Lilly Miller Vegol®) + TX, *Chenopodium ambrosioides near ambrosioides* (Requiem®) + TX, *Chrysanthemum* extract (Crisant®) + TX, extract of neem oil (Trilogy®) + TX, essentials oils of
- 35 *Labiatae* (Botania®) + TX, extracts of clove rosemary peppermint and thyme oil (Garden insect killer®) + TX, Glycinebetaine (Greenstim®) + TX, garlic + TX, lemongrass oil (GreenMatch®) + TX, neem oil + TX, *Nepeta cataria* (Catnip oil) + TX, *Nepeta catarina* + TX, nicotine + TX, oregano oil (MossBuster®) + TX, *Pedaliaceae* oil (Nematon®) + TX, pyrethrum + TX, *Quillaja saponaria* (NemaQ®) + TX, *Reynoutria sachalinensis* (Regalia® + TX, Sakalia®) + TX, rotenone (Eco Roten®) + TX, *Rutaceae*

- plant extract (Soleo®) + TX, soybean oil (Ortho ecosense®) + TX, tea tree oil (Timorex Gold®) + TX, thymus oil + TX, AGNIQUE® MMF + TX, BugOil® + TX, mixture of rosemary sesame peppermint thyme and cinnamon extracts (EF 300®) + TX, mixture of clove rosemary and peppermint extract (EF 400®) + TX, mixture of clove peppermint garlic oil and mint (Soil Shot®) + TX, kaolin (Screen®) + TX,
- 5 storage glucan of brown algae (Laminarin®);
- pheromones including: blackheaded fireworm pheromone (3M Sprayable Blackheaded Fireworm Pheromone®) + TX, Codling Moth Pheromone (Paramount dispenser-(CM)/ Isomate C-Plus®) + TX, Grape Berry Moth Pheromone (3M MEC-GBM Sprayable Pheromone®) + TX, Leafroller pheromone (3M MEC – LR Sprayable Pheromone®) + TX, Muscamone (Snip7 Fly Bait® + TX, Starbar Premium
- 10 Fly Bait®) + TX, Oriental Fruit Moth Pheromone (3M oriental fruit moth sprayable pheromone®) + TX, Peachtree Borer Pheromone (Isomate-P®) + TX, Tomato Pinworm Pheromone (3M Sprayable pheromone®) + TX, Entostat powder (extract from palm tree) (Exosex CM®) + TX, (E + TX,Z + TX,Z)-3 + TX,8 + TX,11 Tetradecatrienyl acetate + TX, (Z + TX,Z + TX,E)-7 + TX,11 + TX,13-Hexadecatrienal + TX, (E + TX,Z)-7 + TX,9-Dodecadien-1-yl acetate + TX, 2-Methyl-1-butanol + TX,
- 15 Calcium acetate + TX, Scenturion® + TX, Biolure® + TX, Check-Mate® + TX, Lavandulyl senecioate;
- Macrobiotics including: *Aphelinus abdominalis* + TX, *Aphidius ervi* (Aphelinus-System®) + TX, *Acerophagus papaya* + TX, *Adalia bipunctata* (Adalia-System®) + TX, *Adalia bipunctata* (Adaline®) + TX, *Adalia bipunctata* (Aphidalia®) + TX, *Ageniaspis citricola* + TX, *Ageniaspis fuscicollis* + TX, *Amblyseius andersoni* (Anderline® + TX, Andersoni-System®) + TX, *Amblyseius californicus*
- 20 (Amblyline® + TX, Spical®) + TX, *Amblyseius cucumeris* (Thripex® + TX, Bugline cucumeris®) + TX, *Amblyseius fallacis* (Fallacis®) + TX, *Amblyseius swirskii* (Bugline swirskii® + TX, Swirskii-Mite®) + TX, *Amblyseius womersleyi* (WomerMite®) + TX, *Amitus hesperidum* + TX, *Anagrus atomus* + TX, *Anagrus fusciventris* + TX, *Anagrus kamali* + TX, *Anagrus loecki* + TX, *Anagrus pseudococci* (Citripar®) + TX, *Anicetus benefices* + TX, *Anisopteromalus calandrae* + TX, *Anthocoris nemoralis*
- 25 (Anthocoris-System®) + TX, *Aphelinus abdominalis* (Apheline® + TX, Aphiline®) + TX, *Aphelinus asychis* + TX, *Aphidius colemani* (Ahipar®) + TX, *Aphidius ervi* (Ervipar®) + TX, *Aphidius gifuensis* + TX, *Aphidius matricariae* (Ahipar-M®) + TX, *Aphidoletes aphidimyza* (Aphidend®) + TX, *Aphidoletes aphidimyza* (Aphidoline®) + TX, *Aphytis lingnanensis* + TX, *Aphytis melinus* + TX, *Aprostocetus hagenowii* + TX, *Atheta coriaria* (Staphyline®) + TX, *Bombus* spp. + TX, *Bombus terrestris* (Natupol
- 30 Beehive®) + TX, *Bombus terrestris* (Beeline® + TX, Tripol®) + TX, *Cephalonomia stephanoderis* + TX, *Chilocorus nigritus* + TX, *Chrysoperla carnea* (Chrysoline®) + TX, *Chrysoperla carnea* (Chrysopa®) + TX, *Chrysoperla rufilabris* + TX, *Cirrospilus ingenuus* + TX, *Cirrospilus quadristriatus* + TX, *Citrostichus phyllocnistoides* + TX, *Closterocerus chamaeleon* + TX, *Closterocerus* spp. + TX, *Coccidoxenoides perminutus* (Planopar®) + TX, *Coccophagus cowperi* + TX, *Coccophagus lycimnia* +
- 35 TX, *Cotesia flavipes* + TX, *Cotesia plutellae* + TX, *Cryptolaemus montrouzieri* (Cryptobug® + TX, Cryptoline®) + TX, *Cybocephalus nipponicus* + TX, *Dacnusa sibirica* + TX, *Dacnusa sibirica* (Minusa®) + TX, *Diglyphus isaea* (Diminex®) + TX, *Delphastus catalinae* (Delphastus®) + TX, *Delphastus pusillus* + TX, *Diachasmimorpha krausii* + TX, *Diachasmimorpha longicaudata* + TX, *Diaparsis jucunda* + TX, *Diaphorencyrtus aligarhensis* + TX, *Diglyphus isaea* + TX, *Diglyphus isaea*

(Miglyphus® + TX, Digline®) + TX, *Dacnusa sibirica* (DacDigline® + TX, Minex®) + TX, *Diversinervus* spp. + TX, *Encarsia citrina* + TX, *Encarsia formosa* (Encarsia max® + TX, Encarline® + TX, En-Strip®) + TX, *Eretmoceris eremicus* (Enermix®) + TX, *Encarsia guadeloupae* + TX, *Encarsia haitiensis* + TX, *Episyrphus balteatus* (Syrphidend®) + TX, *Eretmoceris siphonini* + TX, *Eretmoceris californicus* + TX, *Eretmoceris eremicus* (Ercal® + TX, Eretline e®) + TX, *Eretmoceris eremicus* (Bemimix®) + TX, *Eretmoceris hayati* + TX, *Eretmoceris mundus* (Bemipar® + TX, Eretline m®) + TX, *Eretmoceris siphonini* + TX, *Exochomus quadripustulatus* + TX, *Feltiella acarisuga* (Spidend®) + TX, *Feltiella acarisuga* (Feltiline®) + TX, *Fopius arisanus* + TX, *Fopius ceratitivorus* + TX, Formononetin (Wirless Beehome®) + TX, *Franklinothrips vespiformis* (Vespop®) + TX, *Galendromus* 10 *occidentalis* + TX, *Goniozus legneri* + TX, *Habrobracon hebetor* + TX, *Harmonia axyridis* (HarmoBeetle®) + TX, *Heterorhabditis* spp. (Lawn Patrol®) + TX, *Heterorhabditis bacteriophora* (NemaShield HB® + TX, Nemaseek® + TX, Terranem-Nam® + TX, Terranem® + TX, Larvanem® + TX, B-Green® + TX, NemAttack® + TX, Nematop®) + TX, *Heterorhabditis megidis* (Nemasys H® + TX, BioNem H® + TX, Exhibitline hm® + TX, Larvanem-M®) + TX, *Hippodamia convergens* + TX, 15 *Hypoaspis aculeifer* (Aculeifer-System® + TX, Entomite-A®) + TX, *Hypoaspis miles* (Hypoline m® + TX, Entomite-M®) + TX, *Lbalia leucospoides* + TX, *Lecanoideus floccissimus* + TX, *Lemophagus errabundus* + TX, *Leptomastidea abnormis* + TX, *Leptomastix dactylopii* (Leptopar®) + TX, *Leptomastix epona* + TX, *Lindorus lophanthae* + TX, *Lipolexis oregmae* + TX, *Lucilia caesar* (NatuFly®) + TX, *Lysiphlebus testaceipes* + TX, *Macrolophus caliginosus* (Mirical-N® + TX, Macroline c® + TX, Mirical®) + TX, *Mesoseiulus longipes* + TX, *Metaphycus flavus* + TX, *Metaphycus lounsburyi* 20 + TX, *Micromus angulatus* (Milacewing®) + TX, *Microterys flavus* + TX, *Muscidifurax raptorellus* and *Spalangia cameroni* (Biopar®) + TX, *Neodryinus typhlocybae* + TX, *Neoseiulus californicus* + TX, *Neoseiulus cucumeris* (THRYPEX®) + TX, *Neoseiulus fallaxis* + TX, *Nesideocoris tenuis* (NesidioBug® + TX, Nesibug®) + TX, *Ophyra aenescens* (Biofly®) + TX, *Orius insidiosus* (Thripor-l® 25 + TX, Oriline i®) + TX, *Orius laevigatus* (Thripor-L® + TX, Oriline l®) + TX, *Orius majusculus* (Oriline m®) + TX, *Orius strigicollis* (Thripor-S®) + TX, *Pauesia juniperorum* + TX, *Pediobius foveolatus* + TX, *Phasmarhabditis hermaphrodita* (Nemaslug®) + TX, *Phymastichus coffea* + TX, *Phytoseiulus macropilus* + TX, *Phytoseiulus persimilis* (Spidex® + TX, Phytoline p®) + TX, *Podisus maculiventris* (Podisus®) + TX, *Pseudacteon curvatus* + TX, *Pseudacteon obtusus* + TX, *Pseudacteon tricusps* + 30 TX, *Pseudaphycus maculipennis* + TX, *Pseudleptomastix mexicana* + TX, *Psyllaephagus pilosus* + TX, *Psytalia concolor* (complex) + TX, *Quadrastichus* spp. + TX, *Rhyzobius lophanthae* + TX, *Rodolia cardinalis* + TX, *Rumina decollate* + TX, *Semiolacher petiolatus* + TX, *Sitobion avenae* (Ervibank®) + TX, *Steinernema carpocapsae* (Nematac C® + TX, Millenium® + TX, BioNem C® + TX, NemAttack® + TX, Nemastar® + TX, Capsanem®) + TX, *Steinernema feltiae* (NemaShield® + TX, Nemasys F® + 35 TX, BioNem F® + TX, Steinernema-System® + TX, NemAttack® + TX, Nemaplus® + TX, Exhibitline sf® + TX, Scia-rid® + TX, Entonem®) + TX, *Steinernema kraussei* (Nemasys L® + TX, BioNem L® + TX, Exhibitline srb®) + TX, *Steinernema riobrave* (BioVector® + TX, BioVektor®) + TX, *Steinernema scapterisci* (Nematac S®) + TX, *Steinernema* spp. + TX, *Steinemematid* spp. (Guardian Nematodes®) + TX, *Stethorus punctillum* (Stethorus®) + TX, *Tamarixia radiata* + TX, *Tetrastichus setifer* + TX,

-135-

Thripobius semiluteus + TX, *Torymus sinensis* + TX, *Trichogramma brassicae* (Tricholine b®) + TX, *Trichogramma brassicae* (Tricho-Strip®) + TX, *Trichogramma evanescens* + TX, *Trichogramma minutum* + TX, *Trichogramma ostrinae* + TX, *Trichogramma platneri* + TX, *Trichogramma pretiosum* + TX, *Xanthopimpla stemmator*; and

- 5 other biologicals including: abscisic acid + TX, bioSea® + TX, *Chondrostereum purpureum* (Chontrol Paste®) + TX, *Colletotrichum gloeosporioides* (Collego®) + TX, Copper Octanoate (Cueva®) + TX, Delta traps (Trapline d®) + TX, *Erwinia amylovora* (Harpin) (ProAct® + TX, Ni-HIBIT Gold CST®) + TX, Ferri-phosphate (Ferramol®) + TX, Funnel traps (Trapline y®) + TX, Gallex® + TX, Grower's Secret® + TX, Homo-brassonolide + TX, Iron Phosphate (Lilly Miller Worry Free Ferramol Slug & Snail Bait®) + TX, MCP hail trap (Trapline f®) + TX, *Microctonus hyperodae* + TX, *Mycoleptodiscus terrestris* (Des-X®) + TX, BioGain® + TX, Aminomite® + TX, Zenox® + TX, Pheromone trap (Thripline ams®) + TX, potassium bicarbonate (MilStop®) + TX, potassium salts of fatty acids (Sanova®) + TX, potassium silicate solution (Sil-Matrix®) + TX, potassium iodide + potassiumthiocyanate (Enzicur®) + TX, SuffOil-X® + TX, Spider venom + TX, *Nosema locustae* (Semaspore Organic Grasshopper Control®) + TX, Sticky traps (Trapline YF® + TX, Rebell Amarillo®) + TX and Traps (Takitrapiine y + b®) + TX.
- 10
- 15

The references in brackets behind the active ingredients, e.g. [3878-19-1] refer to the Chemical Abstracts Registry number. The above described mixing partners are known. Where the active ingredients are included in "The Pesticide Manual" [The Pesticide Manual - A World Compendium; Thirteenth Edition; Editor: C. D. S. Tomlin; The British Crop Protection Council], they are described therein under the entry number given in round brackets hereinabove for the particular compound; for example, the compound "abamectin" is described under entry number (1). Where "[CCN]" is added hereinabove to the particular compound, the compound in question is included in the "Compendium of Pesticide Common Names", which is accessible on the internet [A. Wood; Compendium of Pesticide Common Names, Copyright © 1995-2004]; for example, the compound "acetoprole" is described under the internet address <http://www.alanwood.net/pesticides/acetoprole.html>.

20

25

- Most of the active ingredients described above are referred to hereinabove by a so-called "common name", the relevant "ISO common name" or another "common name" being used in individual cases. If the designation is not a "common name", the nature of the designation used instead is given in round brackets for the particular compound; in that case, the IUPAC name, the IUPAC/Chemical Abstracts name, a "chemical name", a "traditional name", a "compound name" or a "development code" is used or, if neither one of those designations nor a "common name" is used, an "alternative name" is employed. "CAS Reg. No" means the Chemical Abstracts Registry Number.
- 30
- 35

The active ingredient mixture of the compounds of formula I selected selected from the compounds defined in the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27 and with active ingredients described above comprises a compound selected from one compound defined in

-136-

the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27 and an active ingredient as described above preferably in a mixing ratio of from 100:1 to 1:6000, especially from 50:1 to 1:50, more especially in a ratio of from 20:1 to 1:20, even more especially from 10:1 to 1:10, very especially from 5:1 and 1:5, special preference being given to a ratio of from 2:1 to 1:2, and a ratio of from 4:1 to 2:1 being likewise preferred, above all in a ratio of 1:1, or 5:1, or 5:2, or 5:3, or 5:4, or 4:1, or 4:2, or 4:3, or 3:1, or 3:2, or 2:1, or 1:5, or 2:5, or 3:5, or 4:5, or 1:4, or 2:4, or 3:4, or 1:3, or 2:3, or 1:2, or 1:600, or 1:300, or 1:150, or 1:35, or 2:35, or 4:35, or 1:75, or 2:75, or 4:75, or 1:6000, or 1:3000, or 1:1500, or 1:350, or 2:350, or 4:350, or 1:750, or 2:750, or 4:750. Those mixing ratios are by weight.

- 5 2:1 being likewise preferred, above all in a ratio of 1:1, or 5:1, or 5:2, or 5:3, or 5:4, or 4:1, or 4:2, or 4:3, or 3:1, or 3:2, or 2:1, or 1:5, or 2:5, or 3:5, or 4:5, or 1:4, or 2:4, or 3:4, or 1:3, or 2:3, or 1:2, or 1:600, or 1:300, or 1:150, or 1:35, or 2:35, or 4:35, or 1:75, or 2:75, or 4:75, or 1:6000, or 1:3000, or 1:1500, or 1:350, or 2:350, or 4:350, or 1:750, or 2:750, or 4:750. Those mixing ratios are by weight.
- 10 The mixtures as described above can be used in a method for controlling pests, which comprises applying a composition comprising a mixture as described above to the pests or their environment, with the exception of a method for treatment of the human or animal body by surgery or therapy and diagnostic methods practised on the human or animal body.
- 15 The mixtures comprising a compound of formula I selected from the compounds defined in the Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27 and one or more active ingredients as described above can be applied, for example, in a single "ready-mix" form, in a combined spray mixture composed from separate formulations of the single active ingredient components, such as a "tank-mix", and in a combined use of the single active ingredients when applied in a sequential
- 20 manner, i.e. one after the other with a reasonably short period, such as a few hours or days. The order of applying the compounds of formula I and the active ingredients as described above is not essential for working the present invention.

- 25 The compositions according to the invention can also comprise further solid or liquid auxiliaries, such as stabilizers, for example unepoxidized or epoxidized vegetable oils (for example epoxidized coconut oil, rapeseed oil or soya oil), antifoams, for example silicone oil, preservatives, viscosity regulators, binders and/or tackifiers, fertilizers or other active ingredients for achieving specific effects, for example bactericides, fungicides, nematocides, plant activators, molluscicides or herbicides.

- 30 The compositions according to the invention are prepared in a manner known per se, in the absence of auxiliaries for example by grinding, screening and/or compressing a solid active ingredient and in the presence of at least one auxiliary for example by intimately mixing and/or grinding the active ingredient with the auxiliary (auxiliaries). These processes for the preparation of the compositions and the use of the compounds I for the preparation of these compositions are also a subject of the
- 35 invention.

The application methods for the compositions, that is the methods of controlling pests of the abovementioned type, such as spraying, atomizing, dusting, brushing on, dressing, scattering or pouring - which are to be selected to suit the intended aims of the prevailing circumstances - and the

-137-

use of the compositions for controlling pests of the abovementioned type are other subjects of the invention. Typical rates of concentration are between 0.1 and 1000 ppm, preferably between 0.1 and 500 ppm, of active ingredient. The rate of application per hectare is generally 1 to 2000 g of active ingredient per hectare, in particular 10 to 1000 g/ha, preferably 10 to 600 g/ha.

5

A preferred method of application in the field of crop protection is application to the foliage of the plants (foliar application), it being possible to select frequency and rate of application to match the danger of infestation with the pest in question. Alternatively, the active ingredient can reach the plants via the root system (systemic action), by drenching the locus of the plants with a liquid composition or by incorporating the active ingredient in solid form into the locus of the plants, for example into the soil, for example in the form of granules (soil application). In the case of paddy rice crops, such granules can be metered into the flooded paddy-field.

10

The compounds of formula I of the invention and compositions thereof are also be suitable for the protection of plant propagation material, for example seeds, such as fruit, tubers or kernels, or nursery plants, against pests of the abovementioned type. The propagation material can be treated with the compound prior to planting, for example seed can be treated prior to sowing. Alternatively, the compound can be applied to seed kernels (coating), either by soaking the kernels in a liquid composition or by applying a layer of a solid composition. It is also possible to apply the compositions when the propagation material is planted to the site of application, for example into the seed furrow during drilling. These treatment methods for plant propagation material and the plant propagation material thus treated are further subjects of the invention. Typical treatment rates would depend on the plant and pest/fungi to be controlled and are generally between 1 to 200 grams per 100 kg of seeds, preferably between 5 to 150 grams per 100 kg of seeds, such as between 10 to 100 grams per 100 kg of seeds.

15

20

25

The term seed embraces seeds and plant propagules of all kinds including but not limited to true seeds, seed pieces, suckers, corns, bulbs, fruit, tubers, grains, rhizomes, cuttings, cut shoots and the like and means in a preferred embodiment true seeds.

30

The present invention also comprises seeds coated or treated with or containing a compound of formula I. The term "coated or treated with and/or containing" generally signifies that the active ingredient is for the most part on the surface of the seed at the time of application, although a greater or lesser part of the ingredient may penetrate into the seed material, depending on the method of application. When the said seed product is (re)planted, it may absorb the active ingredient. In an embodiment, the present invention makes available a plant propagation material adhered thereto with a compound of formula I. Further, it is hereby made available, a composition comprising a plant propagation material treated with a compound of formula I.

35

-138-

Seed treatment comprises all suitable seed treatment techniques known in the art, such as seed dressing, seed coating, seed dusting, seed soaking and seed pelleting. The seed treatment application of the compound formula I can be carried out by any known methods, such as spraying or by dusting the seeds before sowing or during the sowing/planting of the seeds.

5

In each aspect and embodiment of the invention, "consisting essentially" and inflections thereof are a preferred embodiment of "comprising" and its inflections, and "consisting of" and inflections thereof are a preferred embodiment of "consisting essentially of" and its inflections.

10 The disclosure in the present application makes available each and every combination of embodiments disclosed herein.

It should be noted that the disclosure herein in respect of a compound of formula I applies equally in respect of a compound of each of formulae I*, I'a, I-A, I'-A, and Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27. Further the preferred enantiomer of formula I'a or I'-A applies also to compounds of Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27 and Table P. Also, made available herein is an agrochemically acceptable salt, stereoisomer, enantiomer, tautomer and/or N-oxide of the compound of formula formulae I*, I'a, I-A, I'-A, and Tables A-1 to A-27, B-1 to B-27, C-1 to C-27, D-1 to D-27 and E-1 to E-27 and Table P.

20

The compounds of the invention can be distinguished from other similar compounds by virtue of greater efficacy at low application rates and/or different pest control, which can be verified by the person skilled in the art using the experimental procedures, using lower concentrations if necessary, for example 10 ppm, 5 ppm, 2 ppm, 1 ppm or 0.2 ppm; or lower application rates, such as 300, 200 or 100, mg of AI per m². The greater efficacy can be observed by an increased safety profile (against non-target organisms above and below ground (such as fish, birds and bees), improved physico-chemical properties, or increased biodegradability).

25

Biological Examples:

30 The Examples which follow serve to illustrate the invention. Certain compounds of the invention can be distinguished from known compounds by virtue of greater efficacy at low application rates, which can be verified by the person skilled in the art using the experimental procedures outlined in the Examples, using lower application rates if necessary, for example 50 ppm, 24 ppm, 12.5 ppm, 6 ppm, 3 ppm, 1.5 ppm, 0.8 ppm or 0.2 ppm.

35

Example B1: *Diabrotica balteata* (Corn root worm)

Maize sprouts placed onto an agar layer in 24-well microtiter plates were treated with aqueous test solutions prepared from 10'000 ppm DMSO stock solutions by spraying. After drying, the plates were

-139-

infested with L2 larvae (6 to 10 per well). The samples were assessed for mortality and growth inhibition in comparison to untreated samples 4 days after infestation.

The following compounds gave an effect of at least 80% control in at least one of the two categories (mortality or growth inhibition) at an application rate of 200 ppm:

5 P3, P4, P6, P7, P8, P9, P10, P11, P13, P15, P19, P21, P22, P23, P24, P25, P26, P28, P29, P30, P33, P34, P35, P37

Example B2: *Euschistus heros* (Neotropical Brown Stink Bug)

10 Soybean leaves on agar in 24-well microtiter plates were sprayed with aqueous test solutions prepared from 10'000 ppm DMSO stock solutions. After drying the leaves were infested with N2 nymphs. The samples were assessed for mortality and growth inhibition in comparison to untreated samples 5 days after infestation.

The following compounds gave an effect of at least 80% control in at least one of the two categories (mortality or growth inhibition) at an application rate of 200 ppm:

15 P1, P3, P5, P24, P28, P33, P35, P37.

Example B3: *Bemisia tabaci* (Cotton white fly): Feeding/contact activity

20 Cotton leaf discs were placed on agar in 24-well microtiter plates and sprayed with aqueous test solutions prepared from 10'000 ppm DMSO stock solutions. After drying the leaf discs were infested with adult white flies. The samples were checked for mortality 6 days after incubation.

The following compounds resulted in at least 80% mortality at an application rate of 200 ppm:
P35

Example B4: *Chilo suppressalis* (Striped rice stemborer)

25 24-well microtiter plates with artificial diet were treated with aqueous test solutions prepared from 10'000 ppm DMSO stock solutions by pipetting. After drying, the plates were infested with L2 larvae (6-8 per well). The samples were assessed for mortality, anti-feeding effect, and growth inhibition in comparison to untreated samples 6 days after infestation. Control of *Chilo suppressalis* by a test sample is given when at least one of the categories mortality, anti-feedant effect, and growth inhibition
30 is higher than the untreated sample.

The following compounds resulted in at least 80% control in at least one of the three categories (mortality, anti-feeding or growth inhibition) at an application rate of 200 ppm:

P1, P2, P3, P4, P5, P6, P7, P8, P9, P10, P11, P12, P13, P14, P15, P16, P18, P19, P20, P21, P22, P23, P24, P25, P26, P27, P28, P29, P30, P31, P32, P33, P34, P35, P37, P38.

35

Example B5: *Plutella xylostella* (Diamond back moth)

24-well microtiter plates with artificial diet were treated with aqueous test solutions prepared from 10'000 ppm DMSO stock solutions by pipetting. After drying, *Plutella* eggs were pipetted through a

-140-

plastic stencil onto a gel blotting paper and the plate was closed with it. The samples were assessed for mortality and growth inhibition in comparison to untreated samples 8 days after infestation.

The following compounds gave an effect of at least 80% control in at least one of the two categories (mortality or growth inhibition) at an application rate of 200 ppm:

- 5 P1, P2, P3, P4, P5, P6, P7, P8, P9, P10, P11, P12, P13, P14, P15, P16, P17, P18, P19, P20, P21, P22, P23, P24, P25, P26, P27, P28, P29, P30, P31, P32, P33, P34, P35, P36, P37, P38.

Example B6: *Myzus persicae* (Green peach aphid): Intrinsic activity

10 Test compounds prepared from 10'000 ppm DMSO stock solutions were applied by pipette into 24-well microtiter plates and mixed with sucrose solution. The plates were closed with a stretched Parafilm. A plastic stencil with 24 holes was placed onto the plate and infested pea seedlings were placed directly on the Parafilm. The infested plate was closed with a gel blotting paper and another plastic stencil and then turned upside down. The samples were assessed for mortality 5 days after infestation.

- 15 The following compounds resulted in at least 80% mortality at a test rate of 12 ppm:
P3, P22, P23, P26, P27, P28, P35

Example B7: *Spodoptera littoralis* (Egyptian cotton leaf worm)

20 Cotton leaf discs were placed onto agar in 24-well microtiter plates and sprayed with aqueous test solutions prepared from 10'000 ppm DMSO stock solutions. After drying the leaf discs were infested with five L1 larvae. The samples were assessed for mortality, anti-feeding effect, and growth inhibition in comparison to untreated samples 3 days after infestation. Control of *Spodoptera littoralis* by a test sample is given when at least one of the categories mortality, anti-feedant effect, and growth inhibition is higher than the untreated sample.

- 25 The following compounds resulted in at least 80% control in at least one of the three categories (mortality, anti-feeding or growth inhibition) at an application rate of 200 ppm:

P1, P2, P3, P4, P5, P6, P7, P8, P9, P10, P11, P12, P13, P15, P17, P18, P19, P20, P21, P22, P23, P24, P25, P26, P27, P28, P29, P30, P31, P32, P33, P34, P35, P36, P37, P38.

30 Example B8: *Spodoptera littoralis* (Egyptian cotton leaf worm)

Test compounds were applied by pipette from 10'000 ppm DMSO stock solutions into 24-well plates and mixed with agar. Lettuce seeds were placed onto the agar and the multi well plate was closed by another plate which contained also agar. After 7 days the compound was absorbed by the roots and the lettuce grew into the lid plate. The lettuce leaves were then cut off into the lid plate. *Spodoptera*
35 eggs were pipetted through a plastic stencil onto a humid gel blotting paper and the lid plate was closed with it. The samples were assessed for mortality, anti-feedant effect and growth inhibition in comparison to untreated samples 6 days after infestation.

The following compounds gave an effect of at least 80% control in at least one of the three categories (mortality, anti-feeding, or growth inhibition) at a test rate of 12.5 ppm:

-141-

P22, P23, P24, P34, P35, P37

Example B9: *Myzus persicae* (Green Peach Aphid)

Test compounds prepared from 10'000 ppm DMSO stock solutions were applied by a liquid handling robot into 96-well microtiter plates and mixed with a sucrose solution. Parafilm was stretched over the 96-well microtiter plate and a plastic stencil with 96 holes was placed onto the plate. Aphids were sieved into the wells directly onto the Parafilm. The infested plates were closed with a gel blotting card and a second plastic stencil and then turned upside down. The samples were assessed for mortality 5 days after infestation.

The following compounds resulted in at least 80% mortality at an application rate of 50 ppm: P22, P23, P26, P28

Example B10: *Plutella xylostella* (Diamondback Moth)

96-well microtiter plates containing artificial diet were treated with aqueous test solutions, prepared from 10'000 ppm DMSO stock solutions, by a liquid handling robot. After drying, eggs (~30 per well) were infested onto a netted lid which was suspended above the diet. The eggs hatch and L1 larvae move down to the diet. The samples were assessed for mortality 9 days after infestation.

The following compounds gave an effect of at least 80% mortality at an application rate of 500 ppm: P12, P13, P14, P15, P16, P17, P18, P19, P20, P21, P22, P23, P24, P25, P26, P27, P28, P29, P30, P31, P34

Example B11: *Tetranychus urticae* (Two-spotted spider mite): Feeding/contact activity

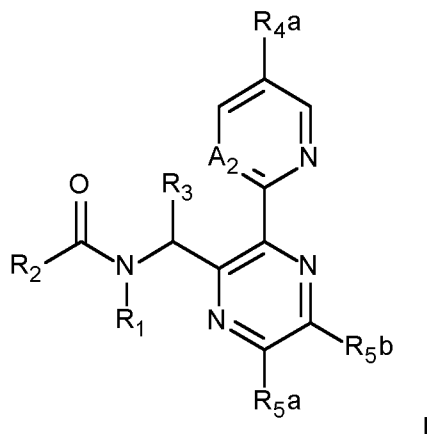
Bean leaf discs on agar in 24-well microtiter plates were sprayed with aqueous test solutions prepared from 10'000 ppm DMSO stock solutions. After drying the leaf discs were infested with a mite population of mixed ages. The samples were assessed for mortality on mixed population (mobile stages) 8 days after infestation.

The following compounds resulted in at least 80% mortality at an application rate of 200 ppm: P16

-142-

CLAIMS

1. A compound of the formula I



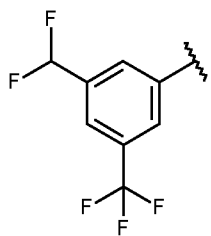
wherein

- 5 R_1 is H, C_1 - C_6 alkyl, C_1 - C_6 cyanoalkyl, aminocarbonyl C_1 - C_6 alkyl, hydroxycarbonyl C_1 - C_6 alkyl, C_1 - C_6 nitroalkyl, trimethylsilane C_1 - C_6 alkyl, C_1 - C_6 haloalkyl, C_2 - C_6 alkenyl, C_2 - C_6 haloalkenyl, C_2 - C_6 alkynyl, C_2 - C_6 haloalkynyl, C_3 - C_4 cycloalkyl C_1 - C_2 alkyl-, C_3 - C_4 cycloalkyl C_1 - C_2 alkyl- wherein the C_3 - C_4 cycloalkyl group is substituted with 1 or 2 halogen atoms, oxetan-3-yl- CH_2 -, benzyl or benzyl substituted with halogen or C_1 - C_6 haloalkyl;
- 10 R_2 is selected from phenyl, pyridine, pyrimidine, pyrazine, pyridazine, and phenyl, pyridine, pyrimidine, pyrazine and pyridazine, each of which is substituted with one to three substituents, provided the substituent(s) are not on either carbon adjacent to the carbon $C=X$ is attached, and each substituent is independently selected from: C_1 - C_3 alkyl, C_1 - C_3 haloalkyl, C_1 - C_3 haloalkylthio, C_1 - C_3 alkoxy, C_1 - C_3 haloalkoxy, halogen, SF_5 , CN, $CONH_2$, and $C(S)NH_2$; R_3 is C_1 - C_3 alkyl or C_1 - C_3 haloalkyl;
- 15 A_2 is CR_{4b} or N;
 R_{4b} is hydrogen, or halogen;
 R_{4a} is cyano, or C_1 - C_3 haloalkoxy;
 R_{5a} and R_{5b} are, independently of each other, selected from hydrogen, halogen, CN, C_1 - C_3 alkyl, C_1 - C_3 haloalkyl, C_3 - C_4 cycloalkyl, C_1 - C_3 alkoxy, and C_1 - C_3 haloalkoxy; or agrochemically acceptable salts,
- 20 stereoisomers, enantiomers, tautomers and N-oxides of the compounds of formula I.
2. The compound according to claim 1 wherein R_3 is methyl.
3. The compound according to either claim 1 or claim 2 wherein R_1 is hydrogen, methyl, ethyl, n-propyl, isobutyl, cyclopropylmethyl or $HCH\equiv CCH_2$ -.
- 25 4. The compound according to any one of claims 1 to 3 wherein R_2 is phenyl, pyridine, pyrimidine, pyrazine, pyridazine, or the phenyl, pyridine, pyrimidine, pyrazine and pyridazine, each of which is substituted with one to three substituents, provided the substituent(s) are not on either carbon

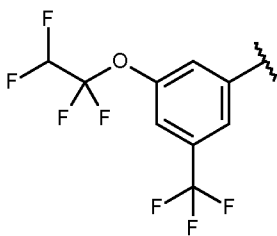
-143-

adjacent to the carbon C=X is attached, and each substituent is independently selected from: C₁-C₃haloalkyl, C₁-C₃haloalkylthio, C₁-C₃alkoxy, C₁-C₃haloalkoxy, and halogen.

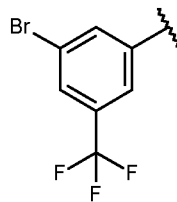
5. The compound according to any one of claims 1 to 4 wherein R₂ is one of M-1 to M-13



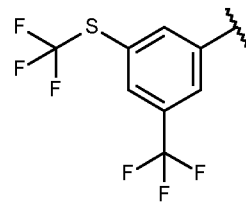
M-1



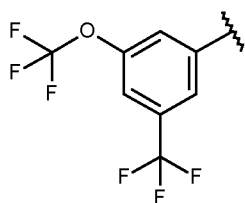
M-2



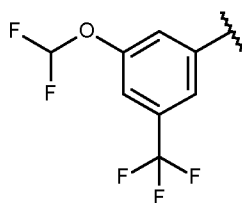
M-3



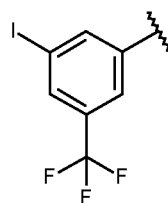
M-4



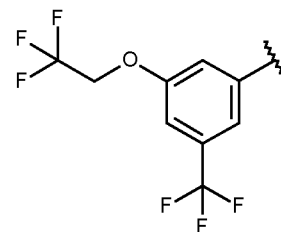
M-5



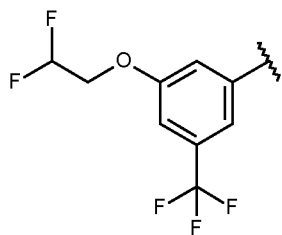
M-6



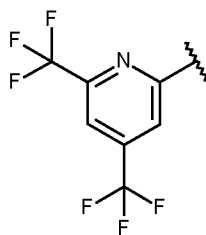
M-7



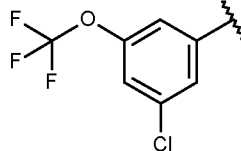
M-8



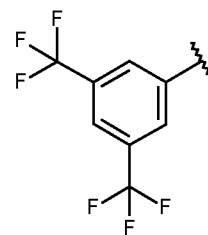
M-9



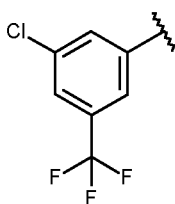
M-10



M-11



M-12



M-13

5

6. The compound according to any one of claims 1 to 5, wherein R_{4a} is cyano, or C₁-C₃fluoroalkoxy.

10 7. The compound according to any one of claims 1 to 6, wherein A₂ is N.

8. The compound according to any one of claims 1 to 7, wherein A₂ is CH.

-144-

9. The compound according to any one of claims 1 to 8, wherein R_{5a} and R_{5b} , independent of each other, are selected from hydrogen, bromo, chloro, methyl, and methoxy.

10 A composition comprising a compound as defined in any one of claims 1 to 9, one or more auxiliaries and diluent, and optionally one more other active ingredient.

11. A method

10 (i) of combating and controlling insects, acarines, nematodes or molluscs which comprises applying to a pest, to a locus of a pest, or to a plant susceptible to attack by a pest an insecticidally, acaricidally, nematocidally or molluscicidally effective amount of a compound as defined as defined in any one of claims 1 to 9 or a composition as defined claim 10; or

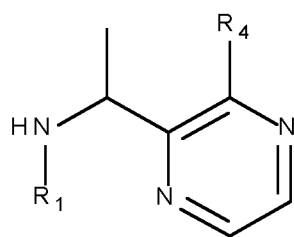
15 (ii) for the protection of plant propagation material from the attack by insects, acarines, nematodes or molluscs, which comprises treating the propagation material or the site, where the propagation material is planted, with an effective amount of a compound as defined in any one of claims 1 to 9 or a composition as defined claim 10; or

20 (iii) of controlling parasites in or on an animal in need thereof comprising administering an effective amount of a compound as defined in any one of claims 1 to 9 or a composition as defined claim 10.

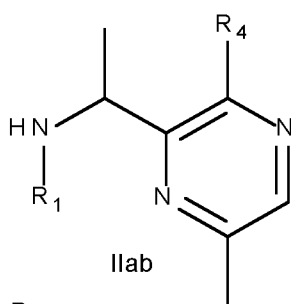
12. A plant propagation material, such as a seed, comprising, or treated with or adhered thereto, a compound as defined in any one of claims 1 to 9 or a composition as defined claim 10.

25 13. A compound of formulae Ilaa to Ilae

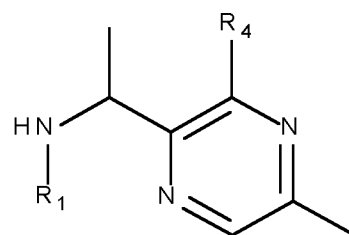
-145-



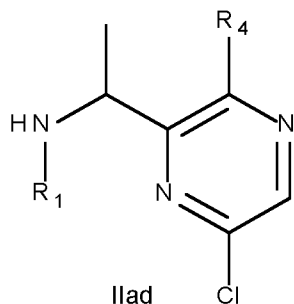
IIaa



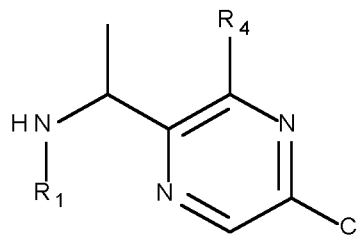
IIab



IIac



IIad



IIae

wherein R_1 is as defined in either claim 1 or 3, and R_4 is the cyclic group containing A_2 and the substituent R_{4a} in formula I, wherein A_2 and R_{4a} are as defined in any one of claims 1, 6, 7 and 8.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2020/059924

A. CLASSIFICATION OF SUBJECT MATTER INV. C07D401/04 C07D403/04 A01N43/26 A61P33/00 ADD.																	
According to International Patent Classification (IPC) or to both national classification and IPC																	
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) C07D A01N A61P Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EPO-Internal, CHEM ABS Data																	
C. DOCUMENTS CONSIDERED TO BE RELEVANT <table border="1"> <thead> <tr> <th>Category*</th> <th>Citation of document, with indication, where appropriate, of the relevant passages</th> <th>Relevant to claim No.</th> </tr> </thead> <tbody> <tr> <td>X</td> <td>WO 2013/092943 A1 (BASF SE [DE]) 27 June 2013 (2013-06-27)</td> <td>1-4,6-13</td> </tr> <tr> <td>A</td> <td>claims 1-61; table C.2; compound 2.47 -----</td> <td>5</td> </tr> <tr> <td>A</td> <td>WO 2017/192385 A1 (ELANCO TIERGESUNDHEIT AG [CH]; LILLY CO ELI [US]) 9 November 2017 (2017-11-09) cited in the application claims 1-16; example 57 -----</td> <td>1-13</td> </tr> <tr> <td>A</td> <td>WO 2016/091776 A1 (EVOTEC AG [DE]) 16 June 2016 (2016-06-16) claims 1-38; compound 324 -----</td> <td>1-13</td> </tr> </tbody> </table>			Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.	X	WO 2013/092943 A1 (BASF SE [DE]) 27 June 2013 (2013-06-27)	1-4,6-13	A	claims 1-61; table C.2; compound 2.47 -----	5	A	WO 2017/192385 A1 (ELANCO TIERGESUNDHEIT AG [CH]; LILLY CO ELI [US]) 9 November 2017 (2017-11-09) cited in the application claims 1-16; example 57 -----	1-13	A	WO 2016/091776 A1 (EVOTEC AG [DE]) 16 June 2016 (2016-06-16) claims 1-38; compound 324 -----	1-13
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.															
X	WO 2013/092943 A1 (BASF SE [DE]) 27 June 2013 (2013-06-27)	1-4,6-13															
A	claims 1-61; table C.2; compound 2.47 -----	5															
A	WO 2017/192385 A1 (ELANCO TIERGESUNDHEIT AG [CH]; LILLY CO ELI [US]) 9 November 2017 (2017-11-09) cited in the application claims 1-16; example 57 -----	1-13															
A	WO 2016/091776 A1 (EVOTEC AG [DE]) 16 June 2016 (2016-06-16) claims 1-38; compound 324 -----	1-13															
☐ Further documents are listed in the continuation of Box C. ☒ See patent family annex.																	
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family																	
Date of the actual completion of the international search		Date of mailing of the international search report															
24 April 2020		06/05/2020															
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer Sáez Díaz, R															

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2020/059924

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2013092943	A1	27-06-2013	AR 090045 A1 15-10-2014
		AU 2012356947 A1 10-07-2014	
		BR 112014015553 A2 13-06-2017	
		CA 2858766 A1 27-06-2013	
		CL 2014001688 A1 07-11-2014	
		CN 104169278 A 26-11-2014	
		CO 6980665 A2 27-06-2014	
		CR 20140276 A 23-07-2014	
		EP 2794601 A1 29-10-2014	
		ES 2727479 T3 16-10-2019	
		JP 6415983 B2 31-10-2018	
		JP 2015502967 A 29-01-2015	
		KR 20140115329 A 30-09-2014	
		US 2014364466 A1 11-12-2014	
		WO 2013092943 A1 27-06-2013	
		ZA 201405336 B 29-11-2017	
WO 2017192385	A1	09-11-2017	AR 108222 A1 01-08-2018
		AU 2017261187 A1 04-10-2018	
		BR 112018069578 A2 22-01-2019	
		CA 3023208 A1 09-11-2017	
		CL 2018003021 A1 15-02-2019	
		CN 109311841 A 05-02-2019	
		CO 2018011825 A2 22-11-2018	
		DK 3452460 T3 16-03-2020	
		EA 201892191 A1 30-04-2019	
		EP 3452460 A1 13-03-2019	
		JP 6370504 B2 08-08-2018	
		JP 2018517716 A 05-07-2018	
		KR 20180132100 A 11-12-2018	
		PT 3452460 T 24-03-2020	
		TW 201803863 A 01-02-2018	
		TW 201938548 A 01-10-2019	
		US 2018186778 A1 05-07-2018	
		US 2019256501 A1 22-08-2019	
		WO 2017192385 A1 09-11-2017	
WO 2016091776	A1	16-06-2016	AR 102948 A1 05-04-2017
		AU 2015359626 A1 29-06-2017	
		BR 112017012327 A2 27-02-2018	
		CA 2969952 A1 16-06-2016	
		CL 2017001488 A1 23-02-2018	
		CN 107207507 A 26-09-2017	
		CN 110256418 A 20-09-2019	
		CR 20170242 A 02-02-2018	
		CU 20170077 A7 07-11-2017	
		DO P2017000137 A 31-07-2017	
		DO P2018000182 A 15-09-2018	
		EA 201791261 A1 29-12-2017	
		EA 201891120 A1 31-10-2018	
		EP 3230281 A1 18-10-2017	
		EP 3587417 A1 01-01-2020	
		IL 252665 A 31-10-2019	
		JP 6544665 B2 17-07-2019	
		JP 6647371 B2 14-02-2020	
		JP 2017537122 A 14-12-2017	
		JP 2019059742 A 18-04-2019	
		KR 20170093203 A 14-08-2017	

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2020/059924

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
		PE 20180227 A1	31-01-2018
		PH 12017501079 A1	18-10-2017
		SG 11201704717V A	30-08-2017
		SV 2017005461 A	30-04-2018
		TN 2017000244 A1	19-10-2018
		TW 201629053 A	16-08-2016
		US 2018093980 A1	05-04-2018
		US 2018118731 A1	03-05-2018
		US 2019185466 A1	20-06-2019
		UY 36422 A	30-06-2016
		WO 2016091776 A1	16-06-2016
