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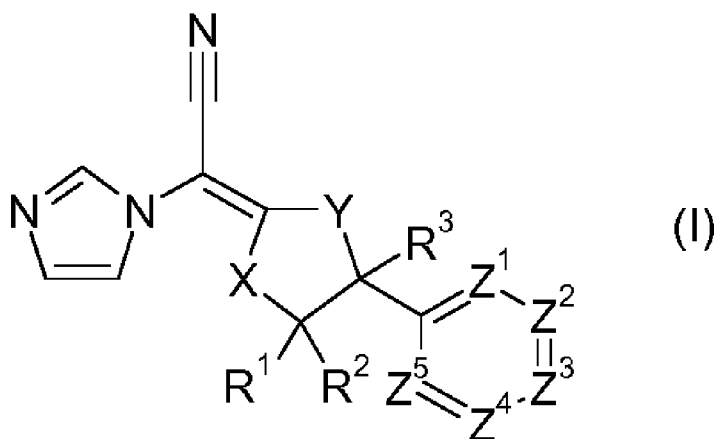
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[Continued on next page]

(54) Title: MICROBIOCIDAL IMIDAZOLE DERIVATIVES



(57) **Abstract:** Compounds of the formula (I) wherein X and Y are independently O, S or NR⁴; Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N; optionally Z¹ and Z², Z² and Z³, Z³ and Z⁴ or Z⁴ and Z⁵ may also belong to a second, annulated saturated, partially unsaturated or aromatic isocyclic or heterocyclic ring to form a bicyclic ring system, with the proviso, that at least two out of Z¹, Z², Z³, Z⁴ and Z⁵ are N; R¹, R² and R³ are independently hydrogen, halogen, C₁-C₆ alkyl, C₁-C₆ haloalkyl, C₁-C₆ alkoxy or C₁-C₆ haloalkoxy; R⁵ is hydrogen, halogen, cyano, hydroxyl, amino, nitro, C₁-C₆ alkyl, C₁-C₆ haloalkyl, C₃-C₆ cycloalkyl, C₃-C₆ halocycloalkyl, C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, C₁-C₆ alkylthio, C₁-C₆ alkylsulfinyl, C₁-C₆ alkylsulfonyl, C₁-C₆ alkylcarbonyl, C₂-C₆ alkenyl, C₂-C₆ haloalkenyl, C₂-C₆ alkenyloxy, C₂-C₆ haloalkenyloxy, C₂-C₆ alkynyl, C₂-C₆ alkynyloxy, C₃-C₆ cycloalkyl-C₂-C₆ alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or substituted heteroaryl, unsubstituted aryloxy or substituted aryloxy, unsubstituted phenylthio or substituted, unsubstituted heteroaryloxy or substituted heteroaryloxy, unsubstituted aryl-C₁-C₆ alkyl or substituted heteroaryl-C₁-C₆alkyl, are useful as pesticides especially fungicide.





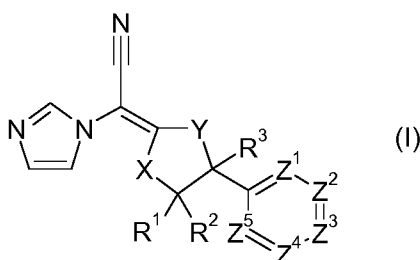
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Microbiocidal Imidazole Derivatives

The present invention relates to microbiocidal imidazole derivatives, e.g. as active ingredients, which have microbiocidal activity, in particular fungicidal activity. The invention also relates to preparation of these imidazole derivatives, to imidazole derivatives used as intermediates in the preparation of these imidazole derivatives, to preparation of these intermediates, to agrochemical compositions which comprise at least one of the imidazole derivatives, to preparation of these compositions and to use of the imidazole derivatives or compositions in agriculture or horticulture for controlling or preventing infestation of plants, harvested food crops, seeds or non-living materials by phytopathogenic microorganisms, preferably fungi.

The present invention provides compounds of formula (I):



wherein,

X and Y are independently O, S or NR⁴;

Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N and at least two out of Z¹, Z², Z³, Z⁴ and Z⁵ are N;

optionally Z¹ and Z², Z² and Z³, Z³ and Z⁴ or Z⁴ and Z⁵ may also belong to a second, annulated saturated, partially unsaturated or aromatic isocyclic or heterocyclic ring to form a bicyclic ring system; R¹, R² and R³ are independently hydrogen, halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy;

R⁵ is independently hydrogen, halogen, cyano, hydroxyl, amino, nitro, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₃-C₆halocycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₁-C₆alkylthio, C₁-C₆alkylsulfinyl, C₁-C₆alkylsulfonyl, C₁-C₆alkylcarbonyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkenyloxy, C₂-C₆haloalkenyloxy, C₂-C₆alkynyl, C₂-C₆alkynyloxy, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or heteroaryl substituted by 1 to 3 substituents R⁶, unsubstituted aryloxy or aryloxy substituted by 1 to 3 substituents R⁶, unsubstituted phenylthio or phenylthio substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryloxy or heteroaryloxy substituted by 1 to 3 substituents R⁶, unsubstituted arylC₁-C₆alkyl or heteroarylC₁-C₆alkyl substituted by 1 to 3 substituents R⁶;

R⁶ is independently hydroxyl, amino, nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆haloalkyl, C₁-C₆haloalkoxy, aryl, aryloxy, or arylC₁-C₆alkyl, in which the alkyl, alkoxy and aryl groups can be substituted by halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy or a salt or a N-oxide thereof.

Where substituents are indicated as being optionally substituted, this means that they may or may not carry one or more identical or different substituents, e.g. one to three substituents. Normally not more than three such optional substituents are present at the same time. Where a group is indicated as being substituted, e.g. alkyl, this includes those groups that are part of other groups, e.g. the alkyl in alkylthio. The same applies for the cycloalkyl, aryl, alkenyl, alkynyl and heteroaryl groups.

The number of substituents does not exceed the number of available C-H and N-H bonds, for example in the group C₁-C₈alkyl substituted by one or more substituents has only one to three substituents if C₁alkyl thus methyl is meant.

The term "halogen" refers to fluorine, chlorine, bromine or iodine, preferably fluorine, chlorine or
5 bromine.

Alkyl substituents may be straight-chained or branched. Alkyl on its own or as part of another substituent is, depending upon the number of carbon atoms mentioned, for example, methyl, ethyl, n-propyl, n-butyl, n-pentyl, n-hexyl and the isomers thereof, for example, iso-propyl, iso-butyl, sec-butyl, tert-butyl or iso-amyl.

10 Cycloalkyl includes cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

Alkenyl substituents can be in the form of straight or branched chains, and the alkenyl moieties, where appropriate, can be of either the (E)- or (Z)-configuration. Examples are vinyl and allyl. The alkenyl groups are preferably C₂-C₆, more preferably C₂-C₄ and most preferably C₂-C₃ alkenyl groups.

Alkynyl substituents can be in the form of straight or branched chains. Examples are ethynyl
15 and propargyl. The alkynyl groups are preferably C₂-C₆, more preferably C₂-C₄ and most preferably C₂-C₃ alkynyl groups.

Haloalkyl groups may contain one or more identical or different halogen atoms and, for example, may stand for CH₂Cl, CHCl₂, CCl₃, CH₂F, CHF₂, CF₃, CF₃CH₂, CH₃CF₂, CF₃CF₂ or CCl₃CCl₂.

Haloalkenyl groups are alkenyl groups, respectively, which are substituted with one or more of
20 the same or different halogen atoms and are, for example, 2,2-difluorovinyl or 1,2-dichloro-2-fluoro-vinyl.

Alkoxy means a radical -OR, where R is alkyl, e.g. as defined above. Alkoxy groups include, but are not limited to, methoxy, ethoxy, 1-methylethoxy, propoxy, butoxy, 1-methylpropoxy and 2-methylpropoxy.

25 Haloalkoxy means a radical -OR, where R is haloalkyl, e.g. is described above. Haloalkoxy groups include, but are not limited to, CH₂ClO, CHCl₂O, CCl₃O, CH₂FO, CHF₂O-, CF₃O-, CF₃CH₂O-, CH₃CF₂O or CCl₃CCl₂O-.

Cyano means a -CN group.

Amino means an -NH₂ group.

30 Hydroxyl or hydroxy stands for a -OH group.

Aryl means a ring system which may be mono-, bi- or tricyclic. Examples of such rings include phenyl, naphthalenyl, anthracenyl, indenyl or phenanthrenyl. A preferred aryl group is phenyl.

Preferred optional substituents for aryl are halogen, cyano, hydroxyl, amino, nitro, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₃-C₆halocycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₁-C₆alkylthio, C₁-
35 C₆alkylsulfinyl, C₁-C₆alkylsulfonyl, C₁-C₆alkylcarbonyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkenyloxy, C₂-C₆haloalkenyloxy, C₂-C₆alkynyl and C₂-C₆alkynyloxy.

Heteroaryl stands for aromatic heterocyclic ring systems, which can be mono-, bi- or tricyclic and wherein at least one oxygen, nitrogen or sulfur atom is present as a ring member, which can be accompanied by other oxygen, nitrogen, sulphur atoms as ring members. Monocyclic and bicyclic
40 aromatic ring systems are preferred. For example, monocyclic heteroaryl may be a 5- or 6-membered ring containing one to three heteroatoms selected from oxygen, nitrogen and sulfur, more preferably

selected from nitrogen and sulfur. Bicyclic heteroaryl may be a 9- or 10-membered bicyclic ring containing one to five heteroatoms, preferably one to three heteroatoms, selected from oxygen, nitrogen and sulfur. Examples of heteroaryl are furyl, thienyl, pyrrolyl, imidazolyl, pyrazolyl, thiazolyl, isothiazolyl, oxazolyl, isoxazolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, pyridyl, pyridazinyl, 5 pyrimidinyl, pyrazinyl, triazinyl, tetrazinyl, indolyl, benzothiophenyl, benzofuranyl, benzimidazolyl, indazolyl, benzotriazolyl, benzothiazolyl, benzoxazolyl, quinolinyl, isoquinolinyl, phthalazinyl, quinoxalinyl, quinazolinyl, cinnolinyl and naphthyridinyl. Heteroaryl rings do not contain adjacent oxygen ring atoms, adjacent sulfur ring atoms or adjacent oxygen and sulfur ring atoms. Preferred optional substituents for heteroaryl are halogen, cyano, hydroxyl, amino, nitro, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₃-C₆halocycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₁-C₆alkylthio, C₁-C₆alkylsulfinyl, C₁-C₆alkylsulfonyl, C₁-C₆alkylcarbonyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkenyloxy, C₂-C₆haloalkenyloxy, C₂-C₆alkynyl and C₂-C₆alkynyloxy.

The presence of one or more possible asymmetric carbon atoms in a compound of formula (I) means that the compounds may occur in optically isomeric forms, i.e. enantiomeric or diastereomeric 15 forms. Also atropisomers may occur as a result of restricted rotation about a single bond. Formula (I) is intended to include all those possible isomeric forms and mixtures thereof. The present invention includes all those possible isomeric forms and mixtures thereof for a compound of formula (I). Likewise, formula (I) is intended to include all possible tautomers. The present invention includes all possible tautomeric forms for a compound of formula (I).

20 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. Albin and S. Pietra, CRC Press, Boca Raton 1991.

25 The following list provides definitions, including preferred definitions, for substituents R¹, R², R³, R⁴, R⁵, R⁶, X, Y, Z¹, Z², Z³, Z⁴ and Z⁵ with reference to compounds of formula (I). For any one of these substituents, any of the definitions given below may be combined with any definition of any other substituent given below or elsewhere in this document.

X and Y are independently O, S or NR⁵.

30 Preferably X and Y are independently O or S, more preferably S.

Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N and that at least two out of Z¹, Z², Z³, Z⁴ and Z⁵ are N.

Preferably two or three out of Z¹, Z², Z³, Z⁴ and Z⁵ are N, more preferably two out of Z¹, Z², Z³, Z⁴ and Z⁵ are N.

35 More preferably the ring formed by Z¹, Z², Z³, Z⁴ and Z⁵ are selected from pyridazinyl, cinnolinyl, pyrimidinyl, pyrazinyl, triazinyl each of the pyridazinyl, cinnolinyl, pyrimidinyl, pyrazinyl, triazinyl can be substituted by F, Cl, Br, C₁-C₆haloalkyl, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted phenox or substituted phenoxy substituted by F, Cl, unsubstituted phenylthio or substituted phenylthio substituted by F. Most preferably the ring formed by Z¹, Z², Z³, Z⁴ and Z⁵ are selected from pyrimidinyl, pyrazinyl 40 each of the from pyrimidinyl, pyrazinyl can be substituted by F, Cl, Br, C₁-C₆haloalkyl, C₃-C₆cycloalkyl-

C₂-C₆alkynyl, unsubstituted phenoxy or phenoxy substituted by F, Cl, unsubstituted phenylthio or substituted phenylthio substituted by F.

R¹, R² and R³ are independently hydrogen, halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

- 5 Preferably R¹, R² and R³ are independently hydrogen, halogen, C₁-C₆alkyl or C₁-C₆haloalkyl; more preferably hydrogen or C₁-C₆alkyl; even more preferably hydrogen or methyl. Most preferably R¹, R² and R³ are all hydrogen.

- R⁵ is independently hydrogen, halogen, cyano, hydroxyl, amino, nitro, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₃-C₆halocycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₁-C₆alkylthio, C₁-C₆alkylsulfinyl, C₁-C₆alkylsulfonyl, C₁-C₆alkylcarbonyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkenyloxy, C₂-C₆haloalkenyloxy, C₂-C₆alkynyl, C₂-C₆alkynyloxy, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted aryloxy, optionally substituted heteroaryloxy, optionally substituted arylC₁-C₆alkyl or optionally substituted heteroarylC₁-C₆alkyl.
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- Preferably R⁵ is independently hydrogen, halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₂-C₆alkenyl, C₂-C₆alkynyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted aryloxy or optionally substituted heteroaryloxy; more preferably hydrogen, halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, optionally substituted aryl or optionally substituted aryloxy; even more preferably hydrogen, halogen, C₁-C₆alkyl, C₁-C₆haloalkyl or optionally substituted aryloxy.
- 15

- Preferably R⁶ is independently hydroxyl, amino, nitro, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy; more preferably R⁶ is independently halogen, C₁-C₆alkyl, C₁-C₆haloalkyl or unsubstituted aryl or substituted aryl which can be substituted by Cl or F.
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- 25 In a preferred embodiment the invention the compounds are of formula (I) wherein

X and Y are independently O or S;

Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N and at least two out of Z¹, Z², Z³, Z⁴ and Z⁵ are N;

R¹, R² and R³ are independently hydrogen, halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy;

- 30 R⁵ is independently hydrogen, halogen, cyano, hydroxyl, amino, nitro, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₃-C₆halocycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₁-C₆alkylthio, C₁-C₆alkylsulfinyl, C₁-C₆alkylsulfonyl, C₁-C₆alkylcarbonyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkenyloxy, C₂-C₆haloalkenyloxy, C₂-C₆alkynyl, C₂-C₆alkynyloxy C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or heteroaryl substituted by 1 to 3 substituents R⁶, unsubstituted aryloxy or aryloxy substituted by 1 to 3 substituents R⁶, unsubstituted phenylthio or phenylthio substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryloxy or heteroaryloxy substituted by 1 to 3 substituents R⁶, unsubstituted arylC₁-C₆alkyl or heteroarylC₁-C₆alkyl substituted by 1 to 3 substituents R⁶;
- 35

- R⁶ is independently hydroxyl, amino, nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy C₁-C₆haloalkyl, C₁-C₆haloalkoxy, aryl, aryloxy, or arylC₁-C₆alkyl, in which the alkyl, alkoxy and aryl groups can be
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independently substituted by one to three substituents selected from halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

In a further preferred embodiment the invention the compounds are of formula (I) wherein

5 X and Y are both S;

Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N with the proviso, that at least two out of Z¹, Z², Z³, Z⁴ and Z⁵ are N;

R¹, R² and R³ are independently hydrogen, C₁-C₆alkyl, C₁-C₆alkoxy or C₃-C₆cycloalkyl;

R⁵ is hydrogen, halogen, cyano, hydroxyl, amino, nitro, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₃-C₆halocycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₁-C₆alkylthio, C₁-C₆alkylsulfinyl, C₁-C₆alkylsulfonyl, C₁-C₆alkylcarbonyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkenyloxy, C₂-C₆haloalkenyloxy, C₂-C₆alkynyl, C₂-C₆alkynyloxy, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or heteroaryl substituted by 1 to 3 substituents R⁶, unsubstituted aryloxy or aryloxy substituted by 1 to 3 substituents R⁶, unsubstituted phenylthio or phenylthio substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryloxy or heteroaryloxy substituted by 1 to 3 substituents R⁶, unsubstituted arylC₁-C₆alkyl or heteroarylC₁-C₆alkyl substituted by 1 to 3 substituents R⁶;

R⁶ is is hydroxyl, amino, nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy C₁-C₆haloalkyl, C₁-C₆haloalkoxy, aryl, aryloxy, or arylC₁-C₆alkyl, in which the alkyl, alkoxy and aryl groups can be independently substituted by one to three substituents selected from halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

In a further preferred embodiment the invention the compounds are of formula (I) wherein

X and Y are both S;

25 Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N with the proviso, that at least two out of Z¹, Z², Z³, Z⁴ and Z⁵ are N;

R¹, R² and R³ are independently hydrogen or C₁-C₆alkyl;

R⁵ is hydrogen, halogen, cyano, hydroxyl, amino, nitro, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₃-C₆halocycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₁-C₆alkylthio, C₁-C₆alkylsulfinyl, C₁-C₆alkylsulfonyl, C₁-C₆alkylcarbonyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkenyloxy, C₂-C₆haloalkenyloxy, C₂-C₆alkynyl, C₂-C₆alkynyloxy, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or heteroaryl substituted by 1 to 3 substituents R⁶, unsubstituted aryloxy or aryloxy substituted by 1 to 3 substituents R⁶, unsubstituted phenylthio or phenylthio substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryloxy or heteroaryloxy substituted by 1 to 3 substituents R⁶, unsubstituted arylC₁-C₆alkyl or heteroarylC₁-C₆alkyl substituted by 1 to 3 substituents R⁶;

R⁶ is is hydroxyl, amino, nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy C₁-C₆haloalkyl, C₁-C₆haloalkoxy, aryl, aryloxy, or arylC₁-C₆alkyl, in which the alkyl, alkoxy and aryl groups can be independently substituted by one to three substituents selected from halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

Preferably in the compounds of formula (I)

X and Y are independently O or S;

Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N with the proviso, that two or three out of Z¹, Z², Z³, Z⁴ and Z⁵ are N;

5 R¹, R² and R³ each hydrogen;

R⁵ is hydrogen, halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or heteroaryl substituted by 1 to 3 substituents R⁶, unsubstituted aryloxy or aryloxy substituted by 1 to 3 substituents R⁶, unsubstituted phenylthio or phenylthio substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryloxy or heteroaryloxy substituted by 1 to 3 substituents R⁶, unsubstituted arylC₁-C₆alkyl or heteroarylC₁-C₆alkyl substituted by 1 to 3 substituents R⁶;

R⁶ is independently hydroxyl, amino, nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆haloalkyl, C₁-C₆haloalkoxy, aryl, aryloxy, or arylC₁-C₆alkyl, in which the alkyl, alkoxy and aryl groups can be

15 independently substituted by one to three substituents selected from halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

In a yet further preferred embodiment the invention the compounds are of formula (I) wherein

X and Y are both S;

20 Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N with the proviso, that two or three out of Z¹, Z², Z³, Z⁴ and Z⁵ are N;

R¹, R² and R³ are all hydrogen;

R⁵ is F, Cl, Br, C₁-C₆haloalkyl, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted phenoxy or substituted phenoxy substituted by F, Cl, unsubstituted phenylthio or substituted phenylthio substituted by F, Cl.

25

In a yet further preferred embodiment X and Y are both S;

the ring formed by Z¹, Z², Z³, Z⁴ and Z⁵ is selected from pyridazinyl, cinnolinyl, pyrimidinyl, pyrazinyl, triazinyl each of them can be substituted by hydrogen, halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or heteroaryl substituted by 1 to 3 substituents R⁶, unsubstituted aryloxy or aryloxy substituted by 1 to 3 substituents R⁶, unsubstituted phenylthio or phenylthio substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryloxy or heteroaryloxy substituted by 1 to 3 substituents R⁶, unsubstituted arylC₁-C₆alkyl or heteroarylC₁-C₆alkyl substituted by 1 to 3 substituents R⁶;

35 R⁶ is is hydroxyl, amino, nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆haloalkyl, C₁-C₆haloalkoxy, aryl, aryloxy, or arylC₁-C₆alkyl, in which the alkyl, alkoxy and aryl groups can be independently substituted by one to three substituents selected from halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

40 More preferably X and Y are both S;

the ring formed by Z^1 , Z^2 , Z^3 , Z^4 and Z^5 is selected from pyridazinyl, cinnoliny, pyrimidinyl, pyrazinyl, triazinyl each of them can be substituted by F, Cl, Br, C_1 - C_6 haloalkyl, C_3 - C_6 cycloalkyl- C_2 - C_6 alkynyl, unsubstituted phenoxy or substituted phenoxy substituted by F, Cl, unsubstituted phenylthio or substituted phenylthio substituted by F, Cl;

5 R^1 , R^2 and R^3 are each is hydrogen.

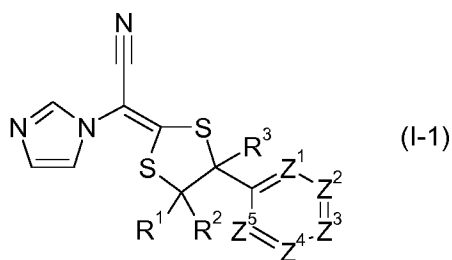
Even more preferably

X and Y are both S;.

the ring formed by Z^1 , Z^2 , Z^3 , Z^4 and Z^5 is selected from pyrimidinyl, pyrazinyl, each of can be
 10 substituted by F, Cl, Br, C_1 - C_6 haloalkyl, C_3 - C_6 cycloalkyl- C_2 - C_6 alkynyl, unsubstituted phenoxy or substituted phenoxy substituted by F, Cl, unsubstituted phenylthio or substituted phenylthio substituted by F, Cl;

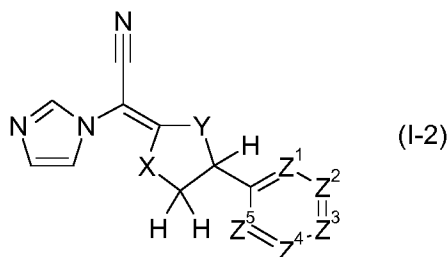
R^1 , R^2 and R^3 are all hydrogen.

15 The invention also relates to compounds of formula (I-1):



in which Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 have the definitions as described for formula (I). Preferred definitions of Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 are as defined for formula (I).

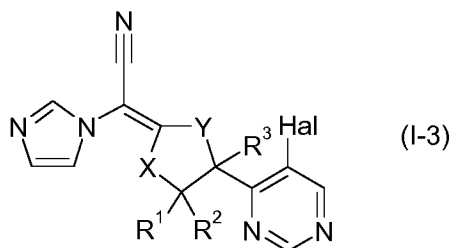
The invention also relates to compounds of formula (I-2):



20

wherein X, Y, Z^1 , Z^2 , Z^3 , Z^4 and Z^5 have the definition as described for formula (I). Preferred definitions of X, Y, Z^1 , Z^2 , Z^3 , Z^4 and Z^5 are as defined for formula (I).

The invention also relates to compounds of formula (I-3):



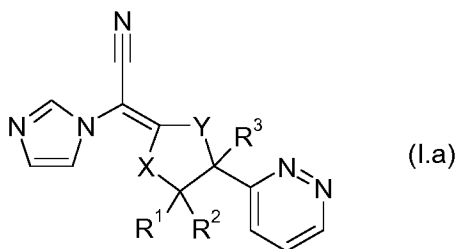
25 wherein X, Y, R^1 , R^2 and R^3 have the definition as described for formula (I) and Hal is halogen. Preferred definitions of X, Y, R^1 , R^2 and R^3 are as defined for formula (I).

In a preferred embodiment the compounds of the present invention are selected from the formula (I.b), (I.c), (I.d), (I.e), (I.f), (I.g), (I.h), (I.i), (I.j), (I.k), (I.l), (I.m), (I.n), (I.o), (I.p), (I.q), (I.r), (I.s), (I.t), (I.u), (I.v), (I.w), (I.x), (I.y), (I.z), (I.aa), (I.ab), (I.ac), (I.ad), (I.ae), (I.af), (I.ag), (I.ah), (I.ai), (I.aj), (I.ak), (I.am), (I.an),
 5 (I.ao), (I.ap), (I.aq), (I.ar), (I.as), (I.at), (I.au), (I.av), (I.aw), (I.ax), (I.ay), (I.az), (I.ba), and (I.bb) and X wherein Y are both S and R¹, R² and R⁴ are independently hydrogen or methyl more preferably R¹, R² and R⁴ are each hydrogen.

In a more preferred embodiment the compounds of the present invention are selected from the
 10 formula (I.k), (I.r), (I.v), (I.aa), (I.ae), (I.ao), and (I.bb) wherein X and Y are both S and R¹, R² and R⁴ are independently hydrogen or methyl more preferably R¹, R² and R⁴ are each hydrogen.

Further preferred embodiments of the present invention are the embodiments E-I.a to E-I.ba, which are defined as compounds of formula (I) which are represented by one formula selected from the
 15 group consisting of the formula (I.a) to (I.ba) as described below, wherein in formulae (I.a) to (I.ba) the meanings of the substituents X, Y, R¹, R² and R³ have the preferred meanings as mentioned above or one of the meanings 1 to 28 given in the corresponding Table 1.

For example, embodiment E-I.a is represented by the compounds of formula (I.a)

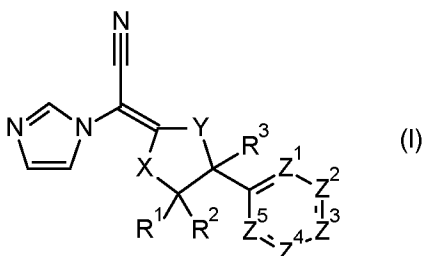


20

and the substituents X, Y, R¹, R² and R³ have the meanings as defined above or one of the meanings 1 to 28 given in the Table 1.

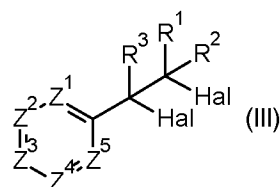
Embodiments E-I.b to E-I.ba are defined accordingly and the substituents X, Y, R¹, R² and R³ have the
 25 meanings as defined above or one of the meanings 1 to 28 given in the corresponding Table 1.

The invention further relates to a process for the preparation of a compound of formula (I) and to compounds obtainable by this process



30

comprising reacting imidazol-1-yl-acetonitrile with compound of formula (III)



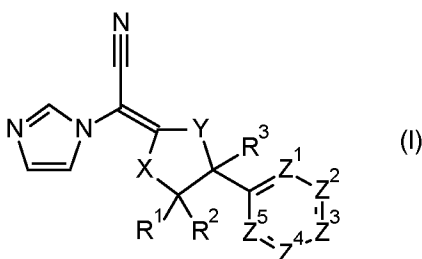
and with a compound of formula (IV)



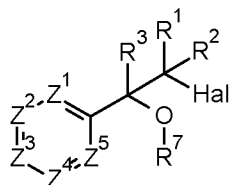
In the presence of a base, wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 are as defined as in any of the
 5 claims 1 to 7 and Hal is halogen. Preferably the base is sodium hydroxide or potassium hydroxide. Preferably the reaction is carried out at room temperature (= ambient temperature) which denotes a temperature between 10°C and 30 °C, more preferable at 20°C to 25°C. Preferably the reaction is carried out in a polar aprotic solvent, preferably in dimethylformamide (DMF), acetonitrile (MeCN), or dimethyl sulfoxide (DMSO), most preferably in dimethyl sulfoxide (DMSO).

10

The invention further relates to a process for the preparation of a compound of formula (I) and to compounds obtainable by this process



15 comprising reacting imidazol-1-yl-acetonitrile with compound of formula (XI)



and with a compound of formula (IV)

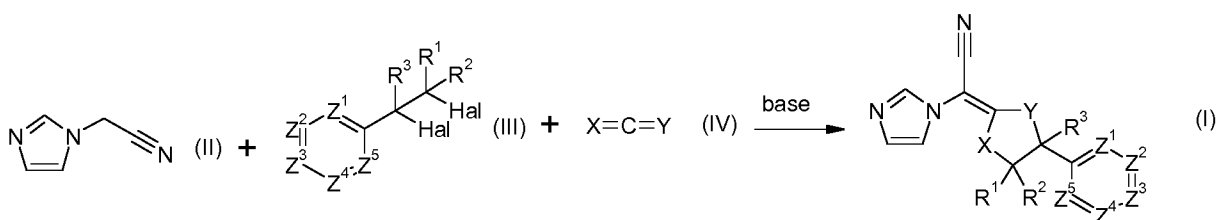


In the presence of a base, wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 are as defined as in any of the
 20 claims 1 to 7, R^7 is a sulfonyl group, and Hal is halogen. Preferably the base is sodium hydroxide or potassium hydroxide. Preferably R^7 is sodium hydroxide or potassium hydroxide. Preferably the reaction is carried out at room temperature (= ambient temperature) which denotes a temperature between 10°C and 30 °C, more preferable at 20°C to 25°C. Preferably the reaction is carried out in a polar aprotic solvent, preferably in dimethylformamide (DMF), acetonitrile (MeCN), or dimethyl
 25 sulfoxide (DMSO), most preferably in dimethyl sulfoxide (DMSO).

Compounds of the present invention can be made as shown in the following schemes, in which, unless otherwise stated, the definition of each variable is as defined above for a compound of formula (I).

The compounds of formula (I), wherein X, Y, Z¹, Z², Z³, Z⁴, Z⁵, R¹, R² and R³ are as defined for formula (I), can be obtained by transformation of imidazol-1-yl-acetonitrile (II), a compound of formula (III), wherein Z¹, Z², Z³, Z⁴, Z⁵, R¹, R² and R³ are as defined for formula (I) and Hal is halogen, and a compound of formula (IV), wherein X and Y are as defined for formula (I), with a base, such as sodium hydroxide or potassium hydroxide. This is shown in Scheme 1.

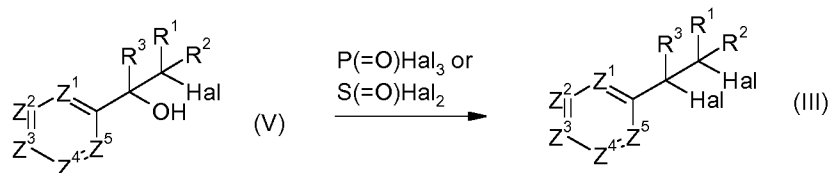
Scheme 1



10

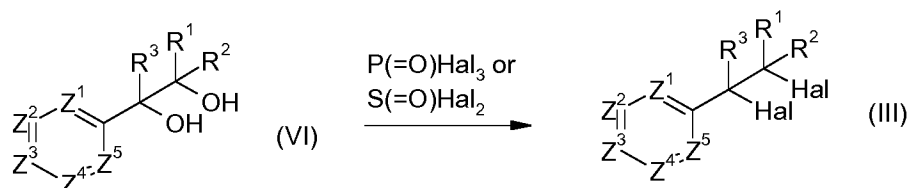
The compounds of formula (III), wherein Z¹, Z², Z³, Z⁴, Z⁵, R¹, R² and R³ are as defined for formula (I) and Hal is halogen, can be obtained by transformation of a compound of formula (V), wherein Z¹, Z², Z³, Z⁴, Z⁵, R¹, R² and R³ are as defined for formula (I) and Hal is halogen, with a halogenation reagent, such as phosphorus oxychloride, phosphorus oxybromide, thionyl chloride or thionyl bromide. This is shown in Scheme 2.

Scheme 2



Alternatively, the compounds of formula (III), wherein Z¹, Z², Z³, Z⁴, Z⁵, R¹ and R² are as defined for formula (I) and Hal is halogen, can be obtained by transformation of a compound of formula (VI), wherein Z¹, Z², Z³, Z⁴, Z⁵, R¹ and R² are as defined for formula (I), with a halogenation reagent, such as phosphorus oxychloride, phosphorus oxybromide, thionyl chloride or thionyl bromide. This is shown in Scheme 3.

Scheme 3

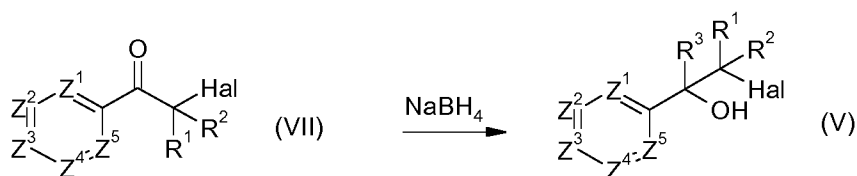


25

The compounds of formula (V) wherein Z¹, Z², Z³, Z⁴, Z⁵, R¹ and R² are as defined for formula (I), R³ is hydrogen and Hal is halogen, can be obtained by transformation of a compound of formula (VII), wherein Z¹, Z², Z³, Z⁴, Z⁵, R¹ and R² are as defined for formula (I) and Hal is halogen, with a reducing agent, such as sodium borohydride. This is shown in Scheme 4.

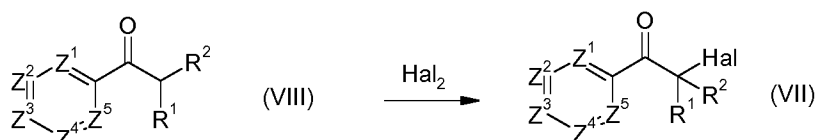
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Scheme 4



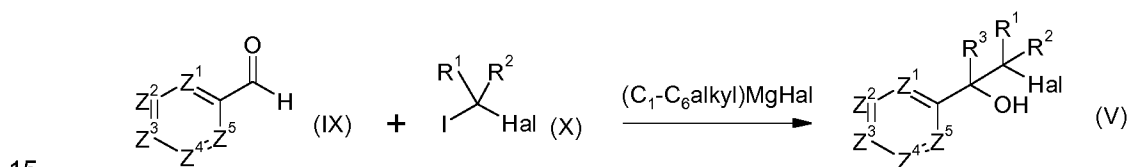
The compounds of formula (VII), wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 and R^2 are as defined for formula (I) and Hal is halogen, can be obtained by transformation of a compound of formula (VIII), wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 and R^2 are as defined for formula (I), with a halogenation reagent, such as chlorine or bromine. This is shown in Scheme 5.

Scheme 5



Alternatively, the compounds of formula (V), wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 and R^2 are as defined for formula (I), R^3 is hydrogen and Hal is halogen, can be obtained by transformation of a compound of formula (IX), wherein Z^1 , Z^2 , Z^3 , Z^4 and Z^5 are as defined for formula (I), with a compound of formula (X), wherein R^1 and R^2 are as defined for formula (I) and Hal is halogen, preferably chloro or bromo, and a base. This is shown in Scheme 6.

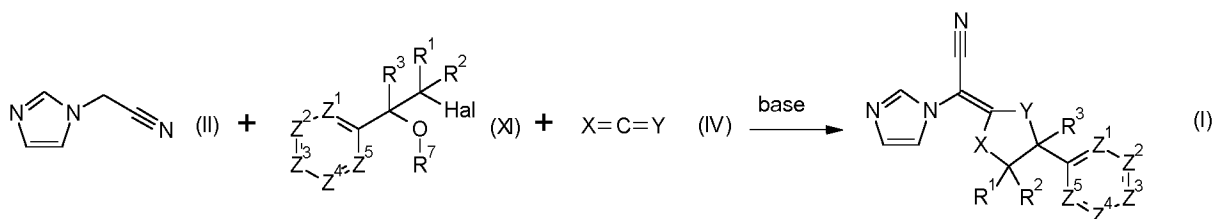
Scheme 6



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Alternatively, the compounds of formula (I), wherein X, Y, Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 are as defined for formula (I), can be obtained by transformation of imidazol-1-yl-acetonitrile (II), a compound of formula (XI), wherein X, Y, Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 are as defined for formula (I), R^6 is a sulfonyl group, such as methylsulfonyl or p-tolylsulfonyl, and Hal is halogen, and a compound of formula (IV), wherein X and Y are as defined for formula (I), with a base, such as sodium hydroxide or potassium hydroxide. This is shown in Scheme 7.

Scheme 7

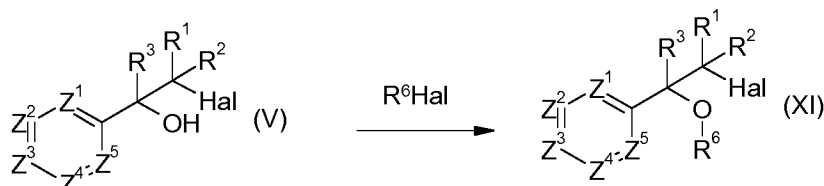


The compounds of formula (XI), wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 are as defined for formula (I), R^7 is a sulfonyl group, such as methylsulfonyl or p-tolylsulfonyl, and Hal is halogen, can be obtained by transformation of a compound of formula (V), wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3

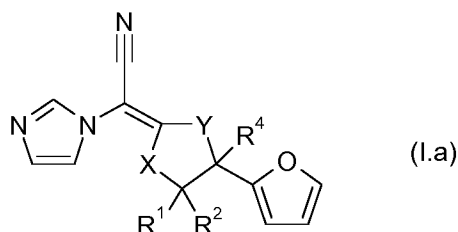
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are as defined for formula (I) and Hal is halogen, with a sulfonyl halide, such as mesyl chloride or tosyl chloride. This is shown in Scheme 8.

Scheme 8



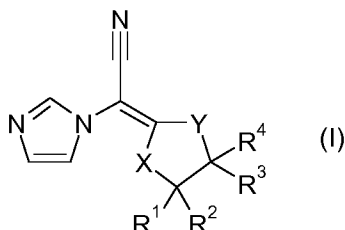
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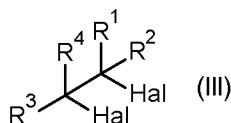
and the substituents X, Y, R¹, R² and R⁴ have the meanings as defined above or one of the meanings
10 1 to 28 given in the Table 1.

Embodiments E-I.b to E-I.bb are defined accordingly and the substituents X, Y, R¹, R² and R⁴ have the meanings as defined above or one of the meanings 1 to 28 given in the corresponding Table 1.

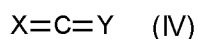
The invention further relates to a process for the preparation of a compound of formula (I) and to
15 compounds obtainable by this process



comprising reacting imidazol-1-yl-acetonitrile with compound of formula (III)



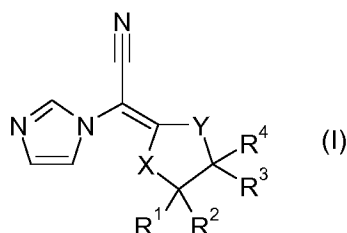
20 and with a compound of formula (IV)



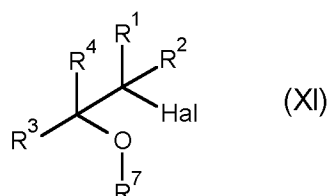
In the presence of a base, wherein X, Y, R¹, R², R³ and R⁴ are as defined as in any of the claims 1 to 7 and Hal is halogen. Preferably the base is sodium hydroxide or potassium hydroxide. Preferably the reaction is carried out at room temperature (= ambient temperature) which denotes a temperature
25 between 10°C and 30 °C, more preferable at 20°C to 25°C. Preferably the reaction is carried out in a

polar aprotic solvent, preferably in dimethylformamide (DMF), acetonitrile (MeCN), or dimethyl sulfoxide (DMSO), most preferably in dimethyl sulfoxide (DMSO).

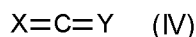
The invention further relates to a process for the preparation of a compound of formula (I) and to
5 compounds obtainable by this process



comprising reacting imidazol-1-yl-acetonitrile with compound of formula (XI)



10 and with a compound of formula (IV)

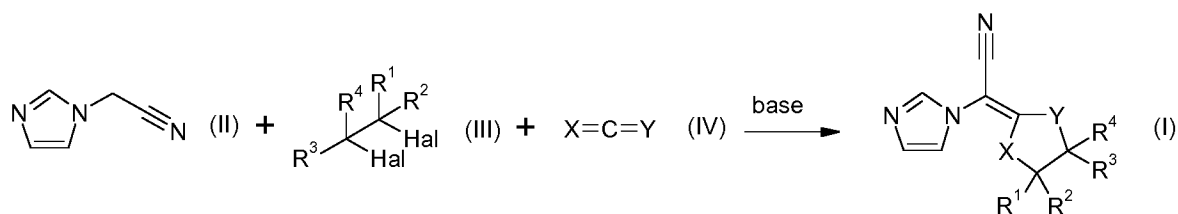


In the presence of a base, wherein X, Y, R¹, R², R³ and R⁴ are as defined as in any of the claims 1 to 7, R⁷ is a sulfonyl group, and Hal is halogen. Preferably the base is sodium hydroxide or potassium hydroxide. Preferably R⁷ is sodium hydroxide or potassium hydroxide. Preferably the reaction is
15 carried out at room temperature (= ambient temperature) which denotes a temperature between 10°C and 30 °C, more preferable at 20°C to 25°C. Preferably the reaction is carried out in a polar aprotic solvent, preferably in dimethylformamide (DMF), acetonitrile (MeCN), or dimethyl sulfoxide (DMSO), most preferably in dimethyl sulfoxide (DMSO).

20 Compounds of the present invention can be made as shown in the following schemes, in which, unless otherwise stated, the definition of each variable is as defined above for a compound of formula (I).

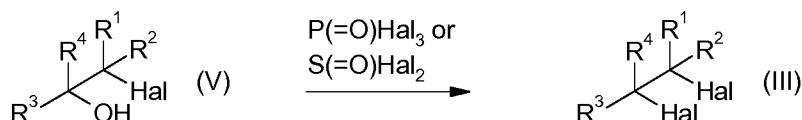
The compounds of formula (I), wherein X, Y, R¹, R², R³ and R⁴ are as defined for formula (I), can be obtained by transformation of imidazol-1-yl-acetonitrile (II), a compound of formula (III), wherein R¹,
25 R², R³ and R⁴ are as defined for formula (I) and Hal is halogen, and a compound of formula (IV), wherein X and Y are as defined for formula (I), with a base, such as sodium hydroxide or potassium hydroxide. This is shown in Scheme 1.

Scheme 1



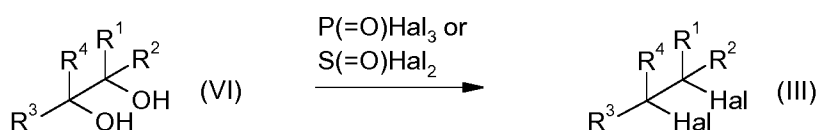
The compounds of formula (III), wherein R^1 , R^2 , R^3 and R^4 are as defined for formula (I) and Hal is halogen, can be obtained by transformation of a compound of formula (V), wherein R^1 , R^2 , R^3 and R^4 are as defined for formula (I) and Hal is halogen, with a halogenation reagent, such as phosphorus
 5 oxychloride, phosphorus oxybromide, thionyl chloride or thionyl bromide. This is shown in Scheme 2.

Scheme 2



Alternatively, the compounds of formula (III), wherein R^1 , R^2 , R^3 and R^4 are as defined for formula (I) and Hal is halogen, can be obtained by transformation of a compound of formula (VI), wherein R^1 , R^2 , R^3 and R^4 are as defined for formula (I), with a halogenation reagent, such as
 10 phosphorus oxychloride, phosphorus oxybromide, thionyl chloride or thionyl bromide. This is shown in Scheme 3.

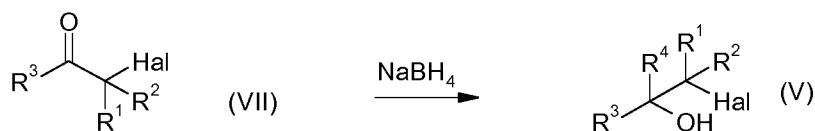
Scheme 3



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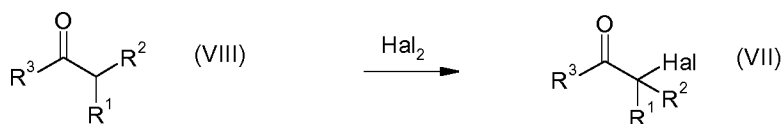
The compounds of formula (V) wherein R^1 , R^2 and R^3 are as defined for formula (I), R^4 is hydrogen and Hal is halogen, can be obtained by transformation of a compound of formula (VII), wherein R^1 , R^2 and R^3 are as defined for formula (I) and Hal is halogen, with a reducing agent, such as sodium borohydride. This is shown in Scheme 4.

20 Scheme 4



The compounds of formula (VII), wherein R^1 , R^2 and R^3 are as defined for formula (I) and Hal is halogen, can be obtained by transformation of a compound of formula (VIII), wherein R^1 , R^2 and R^3 are as defined for formula (I), with a halogenation reagent, such as chlorine or bromine. This is shown
 25 in Scheme 5.

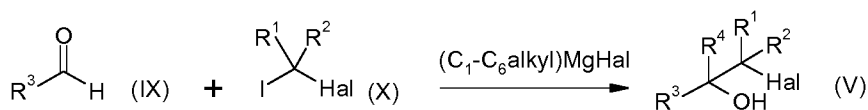
Scheme 5



Alternatively, the compounds of formula (V), wherein R^1 , R^2 and R^3 are as defined for formula (I), R^4 is hydrogen and Hal is halogen, can be obtained by transformation of a compound of formula (IX), wherein R^3 is as defined for formula (I), with a compound of formula (X), wherein R^1 and R^2 are as defined for formula (I) and Hal is halogen, preferably chloro or bromo, and a base. This is shown in

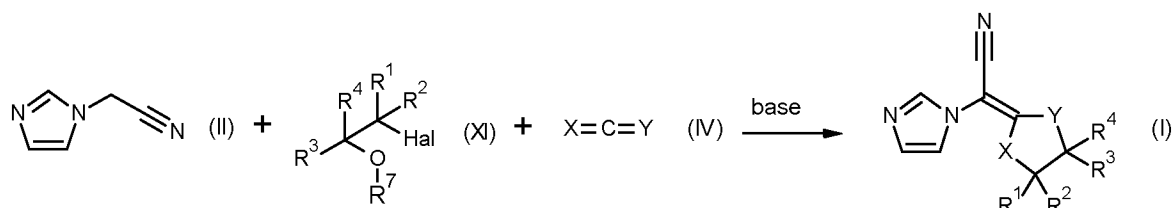
5 Scheme 6.

Scheme 6



Alternatively, the compounds of formula (I), wherein X, Y, R^1 , R^2 , R^3 and R^4 are as defined for formula (I), can be obtained by transformation of imidazol-1-yl-acetonitrile (II), a compound of formula (XI), wherein R^1 , R^2 , R^3 and R^4 are as defined for formula (I), R^6 is a sulfonyl group, such as methylsulfonyl or p-tolylsulfonyl, and Hal is halogen, and a compound of formula (IV), wherein X and Y are as defined for formula (I), with a base, such as sodium hydroxide or potassium hydroxide. This is shown in Scheme 7.

Scheme 7

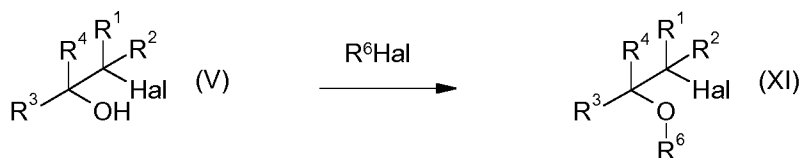


15

The compounds of formula (XI), wherein R^1 , R^2 , R^3 and R^4 are as defined for formula (I), R^7 is a sulfonyl group, such as methylsulfonyl or p-tolylsulfonyl, and Hal is halogen, can be obtained by transformation of a compound of formula (V), wherein R^1 , R^2 , R^3 and R^4 are as defined for formula (I) and Hal is halogen, with a sulfonyl halide, such as mesyl chloride or tosyl chloride. This is shown in

20 Scheme 8.

Scheme 8



The compounds of formula (I) of this invention are useful as plant disease control agents. The present invention therefore further comprises a method for controlling plant diseases caused by fungal plant pathogens comprising applying to the plant or portion thereof to be protected, or to the plant seed to be protected, an effective amount of a compound of the invention or a fungicidal composition containing said compound.

Compounds of formula (I) and fungicidal compositions containing them may be used to control plant diseases caused by a broad spectrum of fungal plant pathogens in the Basidiomycete, Ascomycete, Oomycete and/or Deuteromycete, Blasocladiomycete, Chytridiomycete, Glomeromycete and/or Mucoromycete classes.

30

They are effective in controlling a broad spectrum of plant diseases, such as foliar pathogens of ornamental, turf, vegetable, field, cereal, and fruit crops.

These pathogens may include:

- Oomycetes, including Phytophthora diseases such as those caused by *Phytophthora capsici*,
 5 *Phytophthora infestans*, *Phytophthora sojae*, *Phytophthora fragariae*, *Phytophthora nicotianae*,
Phytophthora cinnamomi, *Phytophthora citricola*, *Phytophthora citrophthora* and *Phytophthora erythroseptica*; Pythium diseases such as those caused by *Pythium aphanidermatum*, *Pythium arrhenomanes*, *Pythium graminicola*, *Pythium irregulare* and *Pythium ultimum*; diseases caused by Peronosporales such as *Peronospora destructor*, *Peronospora parasitica*, *Plasmopara viticola*,
 10 *Plasmopara halstedii*, *Pseudoperonospora cubensis*, *Albugo candida*, *Sclerophthora macrospora* and *Bremia lactucae*; and others such as *Aphanomyces cochlioides*, *Labyrinthula zosterae*,
Peronosclerospora sorghi and *Sclerospora graminicola*.
- Ascomycetes, including blotch, spot, blast or blight diseases and/or rots for example those caused by Pleosporales such as *Stemphylium solani*, *Stagonospora tainanensis*, *Spilocaea oleaginea*,
 15 *Setosphaeria turcica*, *Pyrenochaeta lycopersici*, *Pleospora herbarum*, *Phoma destructiva*,
Phaeosphaeria herpotrichoides, *Phaeocryptocus gaeumannii*, *Ophiosphaerella graminicola*,
Ophiobolus graminis, *Leptosphaeria maculans*, *Hendersonia creberrima*, *Helminthosporium tritici-repentis*, *Setosphaeria turcica*, *Drechslera glycines*, *Didymella bryoniae*, *Cycloconium oleagineum*, *Corynespora cassiicola*, *Cochliobolus sativus*, *Bipolaris cactivora*, *Venturia inaequalis*,
 20 *Pyrenophora teres*, *Pyrenophora tritici-repentis*, *Alternaria alternata*, *Alternaria brassicicola*, *Alternaria solani* and *Alternaria tomatophila*, Capnodiales such as *Septoria tritici*, *Septoria nodorum*, *Septoria glycines*, *Cercospora arachidicola*, *Cercospora sojae*, *Cercospora zeae-maydis*, *Cercospora capsellae* and *Cercospora herpotrichoides*, *Cladosporium carpophilum*, *Cladosporium effusum*,
Passalora fulva, *Cladosporium oxysporum*, *Dothistroma septosporum*, *Isariopsis clavispora*,
 25 *Mycosphaerella fijiensis*, *Mycosphaerella graminicola*, *Mycovellosiella koepkei*, *Phaeoisariopsis bataticola*, *Pseudocercospora vitis*, *Pseudocercospora herpotrichoides*, *Ramularia beticola*,
Ramularia collo-cygni, Magnaporthales such as *Gaeumannomyces graminis*, *Magnaporthe grisea*,
Pyricularia oryzae, Diaporthales such as *Anisogramma anomala*, *Apiognomonina errabunda*, *Cytospora platani*, *Diaporthe phaseolorum*, *Discula destructiva*, *Gnomonia fructicola*, *Greeneria uvicola*,
 30 *Melanconium juglandinum*, *Phomopsis viticola*, *Sirococcus clavigignenti-juglandacearum*, *Tubakia dryina*, *Dicarpella* spp., *Valsa ceratosperma*, and others such as *Actinothyrium graminis*, *Ascochyta pisi*, *Aspergillus flavus*, *Aspergillus fumigatus*, *Aspergillus nidulans*, *Asperisporium caricae*, *Blumeriella jaapii*, *Candida* spp., *Capnodium ramosum*, *Cephaloascus* spp., *Cephalosporium gramineum*,
Ceratocystis paradoxa, *Chaetomium* spp., *Hymenoscyphus pseudoalbidus*, *Coccidioides* spp.,
 35 *Cylindrosporium padi*, *Diplocarpon malae*, *Drepanopeziza campestris*, *Elsinoe ampelina*, *Epicoccum nigrum*, *Epidermophyton* spp., *Eutypa lata*, *Geotrichum candidum*, *Gibellina cerealis*, *Gloeocercospora sorghi*, *Gloeodes pomigena*, *Gloeosporium perennans*; *Gloeotinia temulenta*, *Griphosphaeria corticola*,
Kabatiella lini, *Leptographium microsporum*, *Leptosphaerulina crassiasca*, *Lophodermium seeditiosum*,
Marssonina graminicola, *Microdochium nivale*, *Monilinia fructicola*, *Monographella albescens*,
 40 *Monosporascus cannonballus*, *Naemacyclus* spp., *Ophiostoma novo-ulmi*, *Paracoccidioides brasiliensis*, *Penicillium expansum*, *Pestalotia rhododendri*, *Petriellidium* spp., *Pezicula* spp.,

- Phialophora gregata*, *Phyllachora pomigena*, *Phymatotrichum omnivora*, *Physalospora abdita*, *Plectosporium tabacinum*, *Polyscytalum pustulans*, *Pseudopeziza medicaginis*, *Pyrenopeziza brassicae*, *Ramulispora sorghi*, *Rhabdocline pseudotsugae*, *Rhynchosporium secalis*, *Sacrocladium oryzae*, *Scedosporium* spp., *Schizothyrium pomi*, *Sclerotinia sclerotiorum*, *Sclerotinia minor*,
5 *Sclerotium* spp., *Typhula ishikariensis*, *Seimatosporium mariae*, *Lepteutypa cupressi*, *Septocytia ruborum*, *Sphaceloma perseae*, *Sporonema phacidiodioides*, *Stigmina palmivora*, *Tapesia yallundae*, *Taphrina bullata*, *Thielviopsis basicola*, *Trichoseptoria fructigena*, *Zygophiala jamaicensis*; powdery mildew diseases for example those caused by Erysiphales such as *Blumeria graminis*, *Erysiphe polygoni*, *Uncinula necator*, *Sphaerotheca fuligena*, *Podosphaera leucotricha*, *Podosphaera macularis*
10 *Golovinomyces cichoracearum*, *Leveillula taurica*, *Microsphaera diffusa*, *Oidiopsis gossypii*, *Phyllactinia guttata* and *Oidium arachidis*; molds for example those caused by Botryosphaerales such as *Dothiorella aromatica*, *Diplodia seriata*, *Guignardia bidwellii*, *Botrytis cinerea*, *Botryotinia allii*, *Botryotinia fabae*, *Fusicoccum amygdali*, *Lasiodiplodia theobromae*, *Macrophoma theicola*, *Macrophomina phaseolina*, *Phyllosticta cucurbitacearum*; anthracnoses for example those caused by
15 Glomerelales such as *Colletotrichum gloeosporioides*, *Colletotrichum lagenarium*, *Colletotrichum gossypii*, *Glomerella cingulata*, and *Colletotrichum graminicola*; and wilts or blights for example those caused by Hypocreales such as *Acremonium strictum*, *Claviceps purpurea*, *Fusarium culmorum*, *Fusarium graminearum*, *Fusarium virguliforme*, *Fusarium oxysporum*, *Fusarium subglutinans*, *Fusarium oxysporum* f.sp. *cubense*, *Gerlachia nivale*, *Gibberella fujikuroi*, *Gibberella zeae*,
20 *Gliocladium* spp., *Myrothecium verrucaria*, *Nectria ramulariae*, *Trichoderma viride*, *Trichothecium roseum*, and *Verticillium theobromae*.

- Basidiomycetes, including smuts for example those caused by Ustilaginales such as *Ustilago inoidea* *virens*, *Ustilago nuda*, *Ustilago tritici*, *Ustilago zeae*, rusts for example those caused by Pucciniales such as *Cerotelium fici*, *Chrysomyxa arctostaphyli*, *Coleosporium ipomoeae*, *Hemileia*
25 *vastatrix*, *Puccinia arachidis*, *Puccinia cacabata*, *Puccinia graminis*, *Puccinia recondita*, *Puccinia sorghi*, *Puccinia hordei*, *Puccinia striiformis* f.sp. *Hordei*, *Puccinia striiformis* f.sp. *Secalis*, *Pucciniastrum coryli*, or Uredinales such as *Cronartium ribicola*, *Gymnosporangium juniperi-viginianae*, *Melampsora medusae*, *Phakopsora pachyrhizi*, *Phragmidium mucronatum*, *Physopella ampelosisidis*, *Tranzschelia discolor* and *Uromyces viciae-fabae*; and other rots and diseases such as
30 those caused by *Cryptococcus* spp., *Exobasidium vexans*, *Marasmiellus inoderma*, *Mycena* spp., *Sphacelotheca reiliana*, *Typhula ishikariensis*, *Urocystis agropyri*, *Itersonilia perplexans*, *Corticium invisum*, *Laetisaria fuciformis*, *Waitea circinata*, *Rhizoctonia solani*, *Thanetophorus cucurmeris*, *Entyloma dahliae*, *Entylomella microspora*, *Neovossia molinae* and *Tilletia caries*.

Blastocladiomycetes, such as *Physoderma maydis*.

- 35 Mucoromycetes, such as *Choanephora cucurbitarum*.; *Mucor* spp.; *Rhizopus arrhizus*,
As well as diseases caused by other species and genera closely related to those listed above.

- In addition to their fungicidal activity, the compounds and compositions comprising them may also have activity against bacteria such as *Erwinia amylovora*, *Erwinia caratovora*, *Xanthomonas campestris*, *Pseudomonas syringae*, *Strptomyces scabies* and other related species as well as certain
40 protozoa.

Compounds of formula (I) may be mixed with one or more of compounds selected from those in the following chemical or functional classes:- 1,2,4-thiadiazoles, 2,6-dinitroanilines, acylalanines, aliphatic nitrogenous compounds, amidines, aminopyrimidinols, anilides, anilino-pyrimidines, anthraquinones, antibiotics, aryl-phenylketones, benzamides, benzene-sulfonamides, benzimidazoles, 5 benzothiazoles, benzothiadiazoles, benzothiophenes, benzoylpyridines, benzthiadiazoles, benzylcarbamates, butylamines, carbamates, carboxamides, carpropamids, chloronitriles, cinnamic acid amides, copper containing compounds, cyanoacetamideoximes, cyanoacrylates, cyanoimidazoles, cyanomethylene-thiazolidines, dicyanonitriles, dicarboxamides, dicarboximides, dimethylsulphamates, dinitrophenol carbonates, dinitrophenyl, dinitrophenyl crotonates, diphenyl 10 phosphates, dithiino compounds, dithiocarbamates, dithioethers, dithiolanes, ethyl-amino-thiazole carboxamides, ethyl-phosphonates, furan carboxamides, glucopyranosyls, glucopyranoxyls, glutaronitriles, guanidines, herbicides/plant growth regulators, hexopyranosyl antibiotics, hydroxy(2-amino)pyrimidines, hydroxyanilides, hydroxyisoxazoles, imidazoles, imidazolinones, insecticides/plant growth regulators, isobenzofuranones, isoxazolidinyl-pyridines, isoxazolines, maleimides, mandelic 15 acid amides, mectin derivatives, morpholines, norpholines, n-phenyl carbamates, organotin compounds, oxathiin carboxamides, oxazoles, oxazolidine-diones, phenols, phenoxy quinolines, phenyl-acetamides, phenylamides, phenylbenzamides, phenyl-oxo-ethyl-thiophenes amides, phenylpyrroles, phenylureas, phosphorothiolates, phosphorus acids, phthalamic acids, phthalimides, picolinamides, piperazines, piperidines, plant extracts, polyoxins, propionamides, pthalimides, 20 pyrazole-4-carboxamides, pyrazolinones, pyridazinones, pyridines, pyridine carboxamides, pyridinyl-ethyl benzamides, pyrimdinamines, pyrimidines, pyrimidine-amines, pyrimidione-hydrazone, pyrrolidines, pyrrolquinoliones, quinazolinones, quinolines, quinoline derivatives, quinoline-7-carboxylic acids, quinoxalines, spiroketalamines, strobilurins, sulfamoyl triazoles, sulphamides, tetrazolyloximes, thiadiazines, thiadiazole carboxamides, thiazole carboxanides, thiocyanates, 25 thiophene carboxamides, toluamides, triazines, triazobenthiazoles, triazoles, triazole-thiones, triazolo-pyrimidylamine, valinamide carbamates, ammonium methyl phosphonates, arsenic-containing compounds, benylimidazolylcarbamates, carbonitriles, carboxanilides, carboximidamides, carboxylic phenylamides, diphenyl pyridines, furanilides, hydrazine carboxamides, imidazoline acetates, isophthalates, isoxazolones, mercury salts, organomercury compounds, organophosphates, 30 oxazolidinediones, pentylsulfonyl benzenes, phenyl benzamides, phosphonothionates, phosphorothioates, pyridyl carboxamides, pyridyl furfuryl ethers, pyridyl methyl ethers, SDHIs, thiadiazinanethiones, thiazolidines.

Particularly preferred fungicidal combinations include the following where "I" designates compounds of formula (I): I + (.+/-)-cis-1-(4-chlorophenyl)-2-(1H-1,2,4-triazol-1-yl)-cycloheptanol 35 (huanjunzuo), I + (2RS)-2-bromo-2-(bromomethyl)glutaronitrile (bromothalonil), I + (E)-N-methyl-2-[2-(2,5-dimethylphenoxy)methyl]phenyl]-2-methoxy-iminoacetamide, (mandestrobin), I + 1-(5-bromo-2-pyridyl)-2-(2,4-difluorophenyl)-1,1-difluoro-3-(1,2,4-triazol-1-yl)propan-2-ol, I + 1-methylcyclopropene, I + 2-methyl-, [[4-methoxy-2-[[[(3S,7R,8R,9S)-9-methyl-8-(2-methyl-1-oxopropoxy)-2,6-dioxo-7-(phenylmethyl)-1,5-dioxonan-3-yl]amino]carbonyl]-3-pyridinyl]oxy]propanoic 40 acid methyl ester, I + 2-(1-tert-butyl)-1-(2-chlorophenyl)-3-(1,2,4-triazol-1-yl)-propan-2-ol (TCDP), I + 2,4-D, I + 2,4-DB, I + 2,6-dichloro-N-(4-trifluoromethylbenzyl)-benzamide, I + 2,6-dimethyl-

[1,4]dithiino[2,3-c:5,6-c']dipyrrole-1,3,5,7(2H,6H)-tetraone, I + 2-[[[(1R,5S)-5-[(4-fluorophenyl)methyl]-1-hydroxy-2,2-dimethyl-cyclopentyl]methyl]-4H-1,2,4-triazole-3-thione I + 2-[[3-(2-chlorophenyl)-2-(2,4-difluorophenyl)oxiran-2-yl]methyl]-4H-1,2,4-triazole-3-thione I + ametocradin (imidium), I + 2-[2-[(7,8-difluoro-2-methyl-3-quinolyl)oxy]-6-fluoro-phenyl]propan-2-ol I + 2-[2-fluoro-6-[(8-fluoro-2-methyl-3-quinolyl)oxy]phenyl]propan-2-ol I + cyflufenamid, I + 2-benzyl-4-chlorophenol (Chlorophene), I + 3-(difluoromethyl)-N-(7-fluoro-1,1,3,3-tetramethyl-indan-4-yl)-1-methyl-pyrazole-4-carboxamide I + diclocymet, I + 3-(difluoromethyl)-N-methoxy-1-methyl-N-[1-methyl-2-(2,4,6-trichlorophenyl)ethyl]pyrazole-4-carboxamide, I + 3'-chloro-2-methoxy-N-[(3RS)-tetrahydro-2-oxofuran-3-yl]acet-2',6'-xylidide (clozylacon), I + 3-iodo-2-propinyl n-butylcarbamate (IPBC), I + 4,4,5-trifluoro-3,3-dimethyl-1-(3-quinolyl)isoquinoline I + 4,4-difluoro-3,3-dimethyl-1-(3-quinolyl)isoquinoline I + 5-fluoro-3,3,4,4-tetramethyl-1-(3-quinolyl)isoquinoline I + 9-fluoro-2,2-dimethyl-5-(3-quinolyl)-3H-1,4-benzoxazepine I + tebufloquin, I + 4-CPA, I + 5-fluoro-2-(p-tolylmethoxy)pyrimidin-4-amine I + ferimzone, I + acibenzolar, I + acibenzolar-S-methyl, I + allyl alcohol, I + ametocradin, I + amisulbrom, I + anilazine, I + aureofungin, I + azaconazole, I + azafenidin, I + azithiram, I + azoxystrobin, I + benalaxyl, I + benalaxyl-M, I + benalaxyl-M (kiralaxyl), I + benomyl, I + benthiavalicarb, I + benthiazole (TCMTB), I + benzalkonium chloride, I + benzamorf, I + benzovindiflupyr (solatenol), I + bethoxazin, I + biphenyl, I + bitertanol (biloxazol), I + bixafen, I + BLAD, I + blasticidin-S, I + Bordeaux mixture, I + boscalid, I + bromuconazole, I + bupirimate, I + but-3-ynyl N-[6-[[[(Z)-[(1-methyltetrazol-5-yl)-phenyl-methylene]amino]oxymethyl]-2-pyridyl]carbamate I + dazomet, I + butylamine, I + calcium polysulfide, I + captafol, I + captan, I + carbaryl, I + carbendazim, I + carbendazim chlorhydrate, I + carboxin, I + CAS 517875-34-2 (DAS777), I + chinomethionate, I + chinomethionate (oxythioquinox, quinoxymethionate), I + chitosan, I + chlobenthiazole, I + chlorfenazole, I + chlormequat, I + chloroneb, I + chloropicrin, I + chlorothalonil, I + chlozolate, I + climbazole, I + clofencet, I + copper acetate, I + copper carbonate, I + copper hydroxide, I + copper naphthenate, I + copper oleate, I + copper oxychloride, I + copper oxyquinolate, I + copper silicate, I + copper sulphate, I + copper tallate, I + coumoxystrobin, I + cresol, I + cuprous oxide, I + cyazofamid, I + cyclafuramid, I + cymoxanil, I + cyproconazole, I + cyprodinil, I + daracide, I + dichlofluanid, I + dichlorophen (dichlorophene), I + dichlorprop, I + diclomezine, I + dicloran, I + diethofencarb, I + difenoconazole, I + difenzoquat, I + diflumetorim, I + dimetachlone (dimethaclone), I + dimetconazole, I + dimethipin, I + dimethirimol, I + dimethomorph, I + dimoxystrobin, I + dingjunezuo (Jun Si Qi), I + diniconazole, I + diniconazole-M, I + , I + dinobuton, I + dinocap, I + dinoceton, I + dinopenton, I + diphenylamine, I + dipyrithione, I + ditalimfos, I + dithianon, I + dithioether, I + dodemorph, I + dodicin, I + dodine, I + doguadine, I + drazoxolon, I + edifenphos, I + endothal, I + enestroburin, enoxastrobin I + fenamistobin, I + epoxiconazole, I + etaconazole, I + etem, I + ethaboxam, I + ethephon, I + ethoxyquin, I + famoxadone, I + fenamidone, I + fenarimol, I + fenbuconazole, I + fenfuram, I + fenhexamid, I + fenoxanil, I + fenpiclonil, I + fenpropidin, I + fenpropimorph, I + fenpyrazamine, I + fentin acetate, I + fentin hydroxide, I + ferbam, I + fluazinam, I + fludioxonil, I + flufenoxystrobin, I + flumetralin, I + flumorph, I + fluopicolide, I + fluopicolide (flupicolide), I + fluopyram, I + fluoroimide, I + fluoxastrobin, I + fluquinconazole, I + flusilazole, I + flusulfamide, I + flutianil, I + flutolanil, I + flutriafol, I + fluxapyroxad, I + folpet, I + forchlorfenuron, I + fosetyl, I + fuberidazole, I + furalaxyl, I

- + furametpyr, I + gibberellic acid, I + gibberellins, I + guazatine, I + hexachlorobenzene, I + hexaconazole, I + hymexazol, I + hymexazole, hydroxyisoxazole I + imazalil, I + I + etridiazole, I + imazalil, I + imazalil sulphate, I + imibenconazole, I + iminoctadine, I + iminoctadine triacetate, I + iodocarb (isopropanyl butylcarbamate), I + ipconazole, I + iprobenfos, I + iprodione, I + iprovalicarb,
- 5 I + isofetamid, I + isopropanyl butylcarbamate (iodocarb), I + isoprothiolane, I + isopyrazam, I + isotianil, I + kasugamycin, I + kresoxim-methyl, I + KSF-1002, I + maleic hydrazide, I + mancozeb, I + mandestrobin, I + mandipropamid, I + maneb, I + mepanipyrim, I + mepiquat, I + mepronil, I + meptyldinocap, I + metalaxyl, I + metalaxyl-M (mefenoxam), I + metam, I + metaminostrobin, I + metconazole, I + methyl bromide, I + methyl iodide, I + methyl isothiocyanate, I + metiram
- 10 (polyram), I + metiram-zinc, I + metominostrobin, I + metrafenone, I + m-phenylphenol, I + myclobutanil, I + N'-(2,5-Dimethyl-4-phenoxy-phenyl)-N-ethyl-N-methyl-formamidine, I + N'-[4-(4,5-Dichloro-thiazol-2-yloxy)-2,5-dimethyl-phenyl]-N-ethyl-N-methyl-formamidine, I + N'-[4-[[3-(4-chlorophenyl)methyl]-1,2,4-thiadiazol-5-yl]oxy]-2,5-dimethyl-phenyl]-N-ethyl-N-methyl-formamidine, I + ethirimol, I + N-(2-p-chlorobenzoyl-ethyl)-hexaminium chloride, I + N-[(5-chloro-2-isopropyl-
- 15 phenyl)methyl]-N-cyclopropyl-3-(difluoromethyl)-5-fluoro-1-methyl-pyrazole-4-carboxamide I + N-cyclopropyl-3-(difluoromethyl)-5-fluoro-N-[(2-isopropylphenyl)methyl]-1-methyl-pyrazole-4-carboxamide I + carpropamid, I + nabam, I + naphthalene acetamide, I + NNF-0721, I + octhilinone, I + ofurace, I + orysastrobin, I + osthonol, I + oxadixyl, I + oxasulfuron, I + oxathiapiprolin, I + oxine-copper, I + oxolinic acid, I + oxpoconazole, I + oxycarboxin, I + paclobutrazol, I + pefurazoate, I +
- 20 penconazole, I + pencycuron, I + penflufen, I + penthiopyrad, I + phenamacril, I + phosdiphen, I + phosetyl-Al, I + phosetyl-Al (fosetyl-al), I + phosphorus acids, I + phthalide (fthalide), I + picarbutrazox, I + picoxystrobin, I + piperalin, I + polycarbamate, I + polyoxin D (polyoxrim), I + p-phenylphenol, I + probenazole, I + prochloraz, I + procymidone, I + prohexadione, I +
- prohexadione-calcium, I + propamidine, I + propamocarb, I + propiconazole, I + propineb, I +
- 25 propionic acid, I + proquinazid, I + prothioconazole, I + pyraclostrobin, I + pyrametostrobin, I + pyraoxystrobin, I + pyrazophos, I + pyribencarb (KIF-7767), I + pyrifenoxy, I + pyrimethanil, I + pyriofenone (IKF-309), I + pyroquilon, I + quinoxifen, I + quintozone, I + sedaxane, I + silthiofam, I + simeconazole, I + spiroxamine, I + streptomycin, I + sulphur, I + tebuconazole, I + tebufloquin, I + tecloftalam, I + tecnazene, (TCNB), I + tetraconazole, I + thiabendazole, I + thicyofen, I +
- 30 thidiazuron, I + thifluzamide, I + thiophanate-methyl, I + thiram, I + tiadinil, I + tioxyimid, I + tolclofos-methyl, I + tolprocarb, I + tolylfluanid, I + triadimefon, I + triadimenol, I + triazoxide, I + tribromophenol (TBP), I + tribufos (tributyl phosphorotrithioate), I + triclopyricarb, I + tricyclazole, I + tridemorph, I + trifloxystrobin, I + triflumizole, I + triforine, I + trinexapac, I + triticonazole, I +
- uniconazole, I + validamycin, I + valifenalate, I + vapam, I + vapam (metam sodium), I + vinclozolin,
- 35 I + zineb, I + ziram, I + zoxamide, I + α -naphthalene acetic acid.

Compounds of this invention can also be mixed with one or more further pesticides including insecticides, nematocides, bactericides, acaricides, growth regulators, chemosterilants, semiochemicals, repellents, attractants, pheromones, feeding stimulants or other biologically active compounds to form a multi-component pesticide giving an even broader spectrum of agricultural

40 protection.

Examples of such agricultural protectants with which compounds of this invention can be formulated are:

Insecticides such as abamectin, acephate, acetamiprid, amidoflumet (S-1955), avermectin, azadirachtin, azinphos-methyl, bifenthrin, bifentazate, buprofezin, carbofuran, cartap, 5 chlorantraniliprole (DPX-E2Y45), chlorfenapyr, chlorfluazuron, chlorpyrifos, chlorpyrifos-methyl, chromafenozide, clothianidin, cyflumetofen, cyfluthrin, beta-cyfluthrin, cyhalothrin, lambda-cyhalothrin, cypermethrin, cyromazine, deltamethrin, diafenthiuron, diazinon, dieldrin, diflubenzuron, dimefluthrin, dimethoate, dinotefuran, diofenolan, emamectin, endosulfan, esfenvalerate, ethiprole, fenothiocarb, fenoxycarb, fenpropathrin, fenvalerate, fipronil, flonicamid, flubendiamide, flucythrinate, tau-fluvalinate, 10 flufenimer (UR-50701), flufenoxuron, fonophos, halofenozide, hexaflumuron, hydramethylnon, imidacloprid, indoxacarb, isofenphos, lufenuron, malathion, metaflumizone, metaldehyde, methamidophos, methidathion, methomyl, methoprene, methoxychlor, metofluthrin, monocrotophos, methoxyfenozide, nitenpyram, nithiazine, novaluron, noviflumuron (XDE-007), oxamyl, parathion, parathion-methyl, permethrin, phorate, phosalone, phosmet, phosphamidon, pirimicarb, profenofos, 15 profluthrin, pymetrozine, pyrafluprole, pyrethrin, pyridalyl, pyrifluquinazon, pyriprole, pyriproxifen, rotenone, ryanodine, spinetoram, spinosad, spirotetramat, spiromesifen (BSN 2060), spirotetramat, sulprofos, tebufenozide, teflubenzuron, tefluthrin, terbufos, tetrachlorvinphos, thiacloprid, thiamethoxam, thiodicarb, thiosultap-sodium, tralomethrin, triazamate, trichlorfon and triflumuron;

Bactericides such as streptomycin;

20 Acaricides such as amitraz, chinomethionat, chlorobenzilate, cyenopyrafen, cyhexatin, dicofol, dienochlor, etoxazole, fenazaquin, fenbutatin oxide, fenpropathrin, fenpyroximate, hexythiazox, propargite, pyridaben and tebufenpyrad; and

Biological agents such as *Bacillus thuringiensis*, *Bacillus thuringiensis* delta endotoxin, baculovirus, and entomopathogenic bacteria, virus and fungi.

25 Plant disease control is ordinarily accomplished by applying an effective amount of a compound of this invention either pre- or post-infection, to the portion of the plant to be protected such as the roots, stems, foliage, fruit, seeds, tubers or bulbs, or to the media (soil or sand) in which the plants to be protected are growing. The compounds may also be applied to seeds to protect the seeds and seedlings developing from the seeds. The compounds may also be applied through irrigation water to 30 treat plants.

The present invention envisages application of the compounds of the invention to plant propagation material prior to, during, or after planting, or any combination of these.

Although active ingredients can be applied to plant propagation material in any physiological state, a common approach is to use seeds in a sufficiently durable state to incur no damage during 35 the treatment process. Typically, seed would have been harvested from the field; removed from the plant; and separated from any cob, stalk, outer husk, and surrounding pulp or other non-seed plant material. Seed would preferably also be biologically stable to the extent that treatment would not cause biological damage to the seed. It is believed that treatment can be applied to seed at any time between seed harvest and sowing of seed including during the sowing process.

40 Methods for applying or treating active ingredients on to plant propagation material or to the locus of planting are known in the art and include dressing, coating, pelleting and soaking as well

as nursery tray application, in furrow application, soil drenching, soil injection, drip irrigation, application through sprinklers or central pivot, or incorporation into soil (broad cast or in band). Alternatively or in addition active ingredients may be applied on a suitable substrate sown together with the plant propagation material.

5 Rates of application for these compounds can be influenced by many factors of the environment and should be determined under actual use conditions. Foliage can normally be protected when treated at a rate of from less than about 1 g/ha to about 5,000 g/ha of active ingredient. Seed and seedlings can normally be protected when seed is treated at a rate of from about 0.1 to about 10g per kilogram of seed.

10 Crops of useful plants in which the composition according to the invention can be used include perennial and annual crops, such as berry plants for example blackberries, blueberries, cranberries, raspberries and strawberries; cereals for example barley, maize (corn), millet, oats, rice, rye, sorghum, triticale and wheat; fibre plants for example cotton, flax, hemp, jute and sisal; field crops for example sugar and fodder beet, coffee, hops, mustard, oilseed rape (canola), poppy, sugar cane, sunflower, 15 tea and tobacco; fruit trees for example apple, apricot, avocado, banana, cherry, citrus, nectarine, peach, pear and plum; grasses for example Bermuda grass, bluegrass, bentgrass, centipede grass, fescue, ryegrass, St. Augustine grass and Zoysia grass; herbs such as basil, borage, chives, coriander, lavender, lovage, mint, oregano, parsley, rosemary, sage and thyme; legumes for example beans, lentils, peas and soya beans; nuts for example almond, cashew, ground nut, hazelnut, peanut, 20 pecan, pistachio and walnut; palms for example oil palm; ornamentals for example flowers, shrubs and trees; other trees, for example cacao, coconut, olive and rubber; vegetables for example asparagus, aubergine, broccoli, cabbage, carrot, cucumber, garlic, lettuce, marrow, melon, okra, onion, pepper, potato, pumpkin, rhubarb, spinach and tomato; and vines for example grapes.

Crops are to be understood as being those which are naturally occurring, obtained by 25 conventional methods of breeding, or obtained by genetic engineering. They include crops which contain so-called output traits (e.g. improved storage stability, higher nutritional value and improved flavour).

Crops are to be understood as also including those crops which have been rendered tolerant to herbicides like bromoxynil or classes of herbicides such as ALS-, EPSPS-, GS-, HPPD- and PPO- 30 inhibitors. An example of a crop that has been rendered tolerant to imidazolinones, e.g. imazamox, by conventional methods of breeding is Clearfield® summer canola. Examples of crops that have been rendered tolerant to herbicides by genetic engineering methods include e.g. glyphosate- and glufosinate-resistant maize varieties commercially available under the trade names RoundupReady®, Herculex I® and LibertyLink®.

35 Crops are also to be understood as being those which naturally are or have been rendered resistant to harmful insects. This includes plants transformed by the use of recombinant DNA techniques, for example, to be capable of synthesising one or more selectively acting toxins, such as are known, for example, from toxin-producing bacteria. Examples of toxins which can be expressed include δ -endotoxins, vegetative insecticidal proteins (Vip), insecticidal proteins of bacteria colonising 40 nematodes, and toxins produced by scorpions, arachnids, wasps and fungi.

An example of a crop that has been modified to express the *Bacillus thuringiensis* toxin is the Bt maize KnockOut® (Syngenta Seeds). An example of a crop comprising more than one gene that codes for insecticidal resistance and thus expresses more than one toxin is VipCot® (Syngenta Seeds). Crops or seed material thereof can also be resistant to multiple types of pests (so-called
5 stacked transgenic events when created by genetic modification). For example, a plant can have the ability to express an insecticidal protein while at the same time being herbicide tolerant, for example Herculex I® (Dow AgroSciences, Pioneer Hi-Bred International).

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
10 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-
15 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.

20 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.

25 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
30 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
35 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,
40 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.

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).

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.

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). Many oil derivatives are known from the Compendium of Herbicide Adjuvants, 10th Edition, Southern Illinois University, 2010.

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)
active ingredients	25 %	50 %	75 %
sodium lignosulfonate	5 %	5 %	-
sodium lauryl sulfate	3 %	-	5 %
sodium diisobutyl naphthalenesulfonate	-	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.

5

<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

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	3 %
(4-5 mol of ethylene oxide)	
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

10 this concentrate by dilution with water.

<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 5 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 %
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
 10 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
 15 an apparatus suitable for that purpose.

The Examples which follow serve to illustrate the invention. The 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
 20 Examples, using lower application rates if necessary, for example 50 ppm, 12.5 ppm, 6 ppm, 3 ppm, 1.5 ppm or 0.8 ppm.

Example 1: This example illustrates the preparation of (2E)-2-[4-(3-bromopyrazin-2-yl)-1,3-dithiolan-2-ylidene]-2-imidazol-1-yl-acetonitrile (Compound I.a.o.25)

25

a) Preparation of 2-bromo-1-(3-bromopyrazin-2-yl)ethanone

A 50 mL flask is charged with 1-(3-chloropyrazin-2-yl)ethanone (611 mg, 3.9 mmol) and acetic acid (20 mL). A solution of HBr (33% in acetic acid, 8.2 mL) is added slowly, followed by bromine (0.2 mL, 3.9 mmol) to give an orange suspension. After one hour stirring at room temperature, a second
 30 equivalent of bromine (0.2 mL, 3.9 mmol) is added. The reaction mixture is further stirred for one hour at room temperature and is diluted with methyl-tert-butylether (10 mL) and is cooled at 0°C. The suspension is filtered, and the precipitated [product, HBr] salt is washed with methyl-tert-butylether (2 × 5 mL). The collected solid is dissolved in a saturated aqueous solution of sodium carbonate (5 mL). The aqueous solution is extracted with dichloromethane, the combined organic phases are washed
 35 with brine, dried with sodium sulphate, filtered and are evaporated to give 2-bromo-1-(3-bromopyrazin-

2-yl)ethanone. ¹H-NMR (400 MHz, CDCl₃): δ = 8.59 (d, *J* = 2.2 Hz, 1H), 8.54 (d, *J* = 2.2 Hz, 1H), 4.66 (s, 2H). GC-MS (CI): *m/z* = 281 (M+1).

b) Preparation of 2-bromo-1-(3-bromopyrazin-2-yl)ethanol

5 Sodium borohydride (24 mg, 0.62 mmol) is added as a solid at 0°C to a solution of 2-bromo-1-(3-bromopyrazin-2-yl)ethanone (645 mg, 2.3 mmol) in methanol (4 mL). The reaction mixture is further stirred at room temperature for one hour and is quenched carefully with a saturated aqueous solution of ammonium chloride (5 mL). The aqueous mixture is extracted with methyl-*tert*-butylether. The combined organic phases are washed with brine, dried with sodium sulphate, filtered and are
10 evaporated. Purification by chromatography on silica gel (heptanes / ethylacetate 20:1) give 2-bromo-1-(3-bromopyrazin-2-yl)ethanol as a colorless oil. ¹H-NMR (400 MHz, CDCl₃): δ = 8.57 (d, *J* = 2.2 Hz, 1H), 8.39 (d, *J* = 2.2 Hz, 1H), 5.33 (td, *J* = 4.4, 8.4 Hz, 1H), 4.13 (d, *J* = 8.4 Hz, 1H), 3.86 (d, *J* = 4.4 Hz, 2H). GC-MS (CI): *m/z* = 283 (M+1).

15 c) Preparation of [2-bromo-1-(3-bromopyrazin-2-yl)ethyl] methanesulfonate

Methanesulfonyl chloride (128 μL, 1.63 mmol) is added at 0°C to a solution of 2-bromo-1-(3-bromopyrazin-2-yl)ethanol (461 mg, 1.63 mmol) and triethylamine (0.25 mL, 1.8 mmol) in ethyl acetate (3 mL). The resulting white suspension is stirred at 0°C for 30 minutes and is filtered. The filtrate is diluted with H₂O (10 mL) and the biphasic mixture is decanted. The aqueous phase is extracted with
20 ethyl acetate and the combined organic phases are washed with brine, dried with sodium sulphate, filtered and evaporated to give [2-bromo-1-(3-bromopyrazin-2-yl)ethyl] methanesulfonate as a pale yellow oil. ¹H-NMR (400 MHz, CDCl₃): δ = 8.63 (d, *J* = 2.2 Hz, 1H), 8.43 (d, *J* = 2.2 Hz, 1H), 6.27 (t, *J* = 6.8 Hz, 1H), 3.96 (t, *J* = 10.5 Hz, 1H), 3.87 (dd, *J* = 6.8, 10.5 Hz, 1H), 3.13 – 3.10 (m, 4H). GC-MS (CI): *m/z* = 361 (M+1).

25

d) Preparation of (2E)-2-[4-(3-bromopyrazin-2-yl)-1,3-dithiolan-2-ylidene]-2-imidazol-1-yl-acetonitrile (Compound I.a.25)

A 10 mL flask is charged with powdered KOH (159 mg, 2.43 mmol), DMSO (3 mL), purged with Argon and is cooled at 10°C with a water bath. A solution of 2-imidazol-1-ylacetonitrile (125 mg, 1.17 mmol) and carbon disulfide (0.147 mL, 2.5 mmol) in DMSO (3 mL) is then added slowly to give
30 an orange mixture. The cooling bath is removed and the reaction is stirred at room temperature for 30 minutes. A solution of [2-bromo-1-(3-bromopyrazin-2-yl)ethyl] methanesulfonate (350 mg, 0.97 mmol) in DMSO (2 mL) is then added dropwise. After two hours, the reaction mixture is poured into H₂O (15 mL). The aqueous phase is extracted with dichloromethane, the combined organic phases are washed
35 with brine, dried with Na₂SO₄, filtered and evaporated to give a crude pale yellow residue. Purification by chromatography on silica gel (heptanes / EtOAc, 5:1→1:1→1:3) afford (2E)-2-[4-(3-bromopyrazin-2-yl)-1,3-dithiolan-2-ylidene]-2-imidazol-1-yl-acetonitrile (Compound I.a.25) as a yellow gum. ¹H-NMR (400 MHz, CDCl₃): δ = 8.55 (d, *J* = 2.2 Hz, 1H), 8.41 (d, *J* = 2.2 Hz, 1H), 7.61 (s, 1H), 7.16 (s, 1H), 7.03 (s, 1H), 5.67 (t, *J* = 5.5 Hz, 1H), 4.14 (dd, *J* = 5.5, 11.8 Hz, 1H), 3.91 (dd, *J* = 5.5, 11.8 Hz, 1H).
40 MS (ESI): *m/z* = 366, 368 (M+1).

Table 1 below illustrates examples of individual compounds of formula (I) according to the invention.

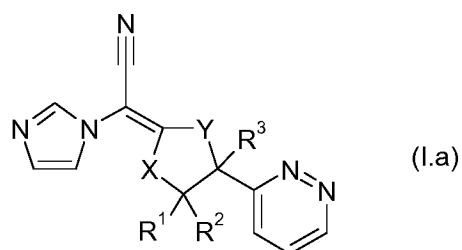
Table 1: individual compounds of formula (I) according to the invention

5

Comp. No.	X	Y	R ¹	R ²	R ³
1	NH	NH	H	H	H
2	NH	NH	CH ₃	H	H
3	NH	NH	CH ₃	CH ₃	H
4	NH	NH	H	H	CH ₃
5	NH	O	H	H	H
6	NH	O	CH ₃	H	H
7	NH	O	CH ₃	CH ₃	H
8	NH	O	H	H	CH ₃
9	NH	S	H	H	H
10	NH	S	CH ₃	H	H
11	NH	S	CH ₃	CH ₃	H
12	NH	S	H	H	CH ₃
13	O	NH	H	H	H
14	O	NH	CH ₃	H	H
15	O	NH	CH ₃	CH ₃	H
16	O	NH	H	H	CH ₃
17	O	O	H	H	H
18	O	O	CH ₃	H	H
19	O	O	CH ₃	CH ₃	H
20	O	O	H	H	CH ₃
21	S	NH	H	H	H
22	S	NH	CH ₃	H	H
23	S	NH	CH ₃	CH ₃	H
24	S	NH	H	H	CH ₃
25	S	S	H	H	H
26	S	S	CH ₃	H	H
27	S	S	CH ₃	CH ₃	H
28	S	S	H	H	CH ₃

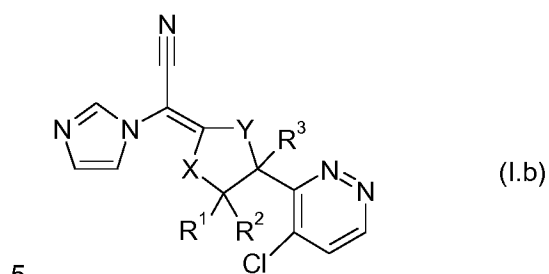
where

a) 28 compounds of formula (I.a):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

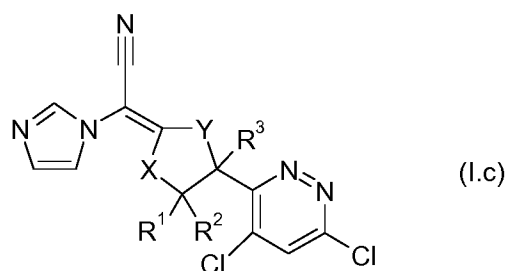
b) 28 compounds of formula (I.b):



5

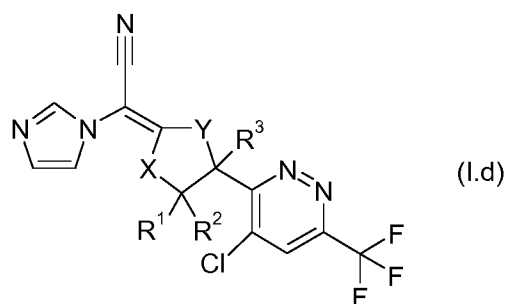
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

c) 28 compounds of formula (I.c):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

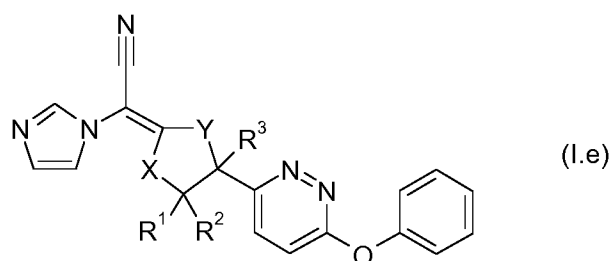
d) 28 compounds of formula (I.d):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

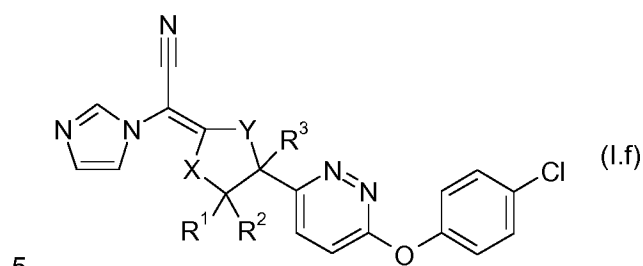
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e) 28 compounds of formula (I.e):



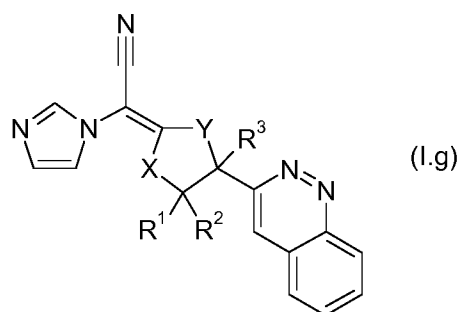
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

f) 28 compounds of formula (I.f):



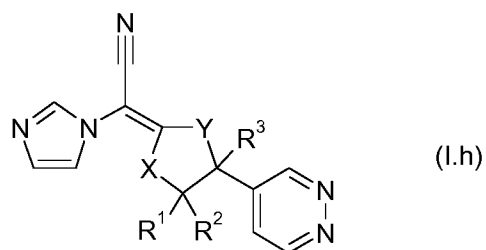
5 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

g) 28 compounds of formula (I.g):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

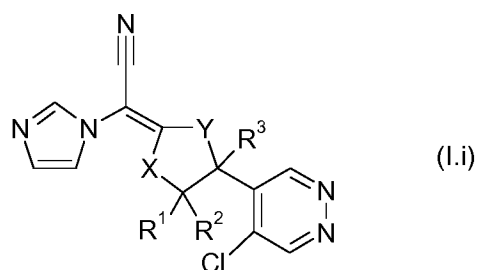
h) 28 compounds of formula (I.h):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

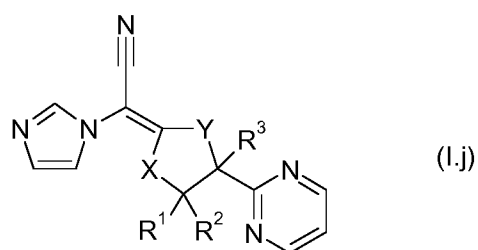
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i) 28 compounds of formula (I.i):



Wherein X, Y, R¹, R² and R³ are as defined in Table 1.

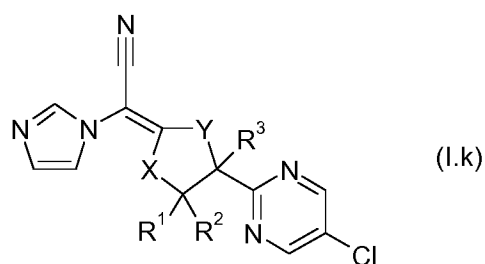
j) 28 compounds of formula (I.j):



5

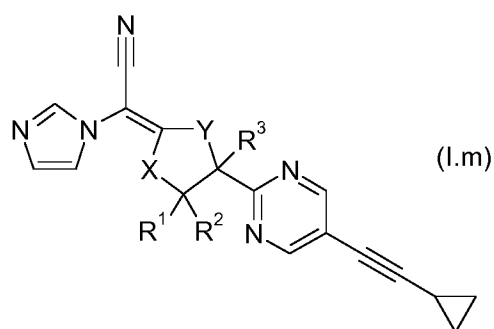
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

k) 28 compounds of formula (I.k):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

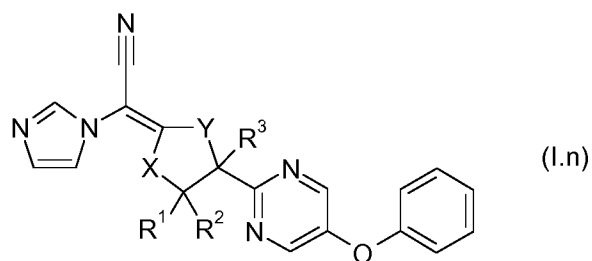
m) 28 compounds of formula (I.m):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

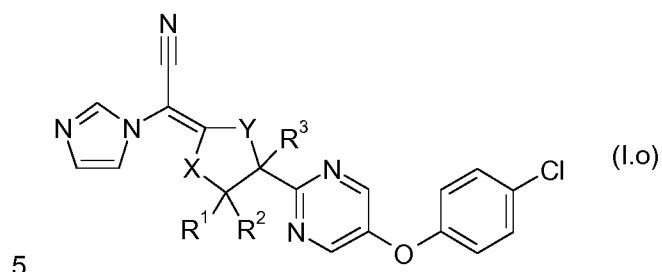
15

n) 28 compounds of formula (I.n):



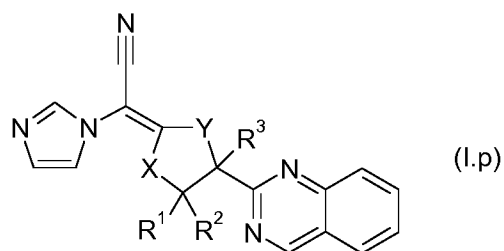
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

o) 28 compounds of formula (I.o):



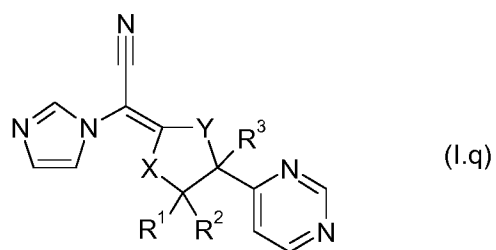
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

p) 28 compounds of formula (I.p):



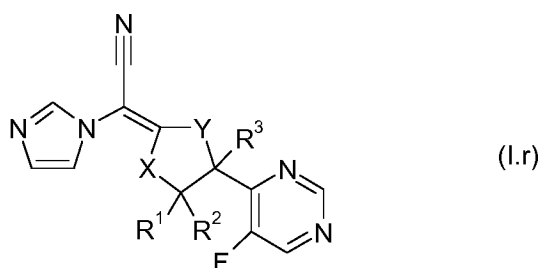
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

q) 28 compounds of formula (I.q):



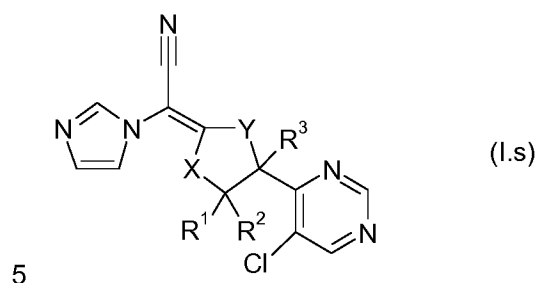
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

r) 28 compounds of formula (I.r):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

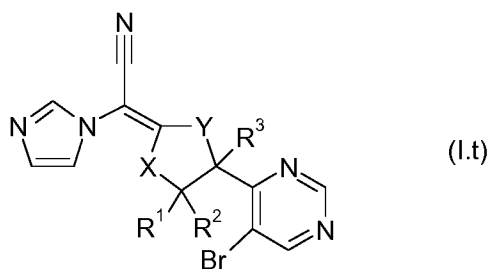
s) 28 compounds of formula (I.s):



5

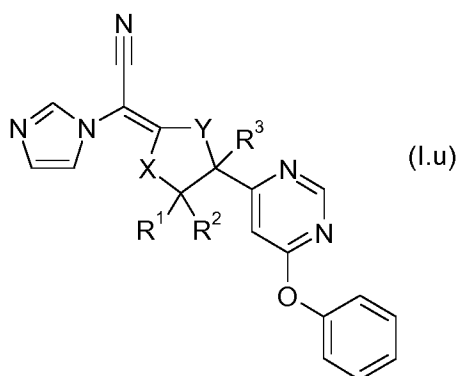
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

t) 28 compounds of formula (I.t):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

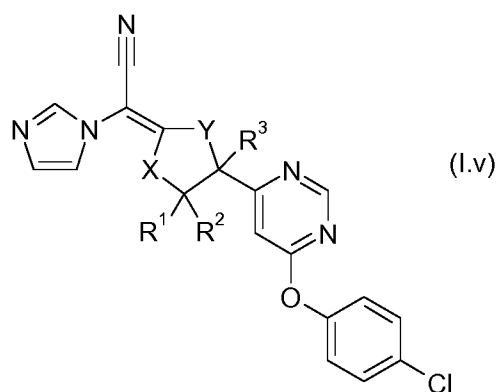
u) 28 compounds of formula (I.u):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

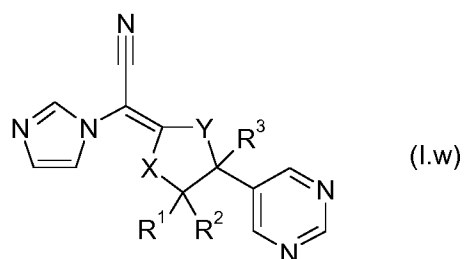
15

v) 28 compounds of formula (I.v):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

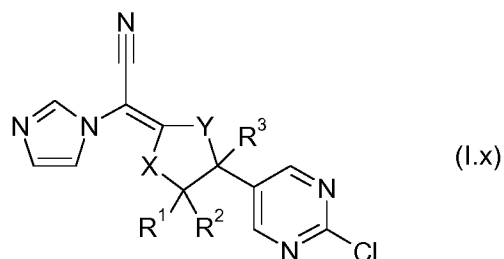
w) 28 compounds of formula (I.w):



5

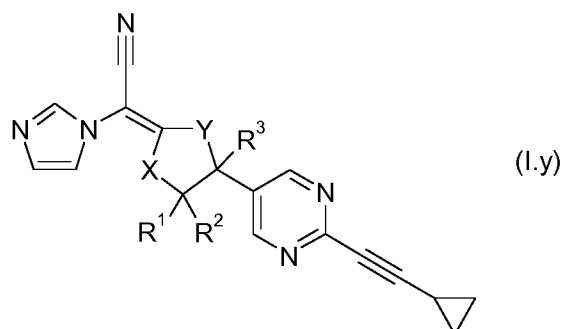
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

x) 28 compounds of formula (I.x):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

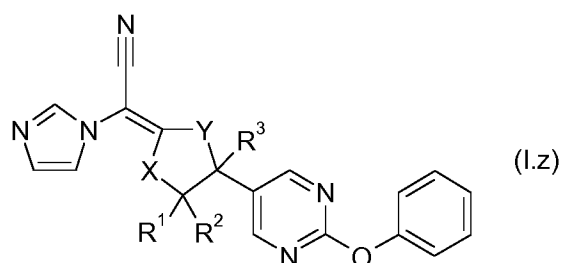
y) 28 compounds of formula (I.y):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

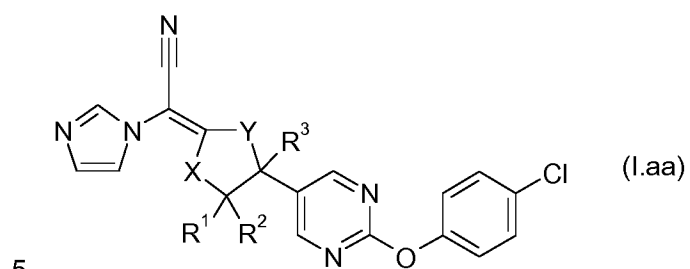
15

z) 28 compounds of formula (I.z):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

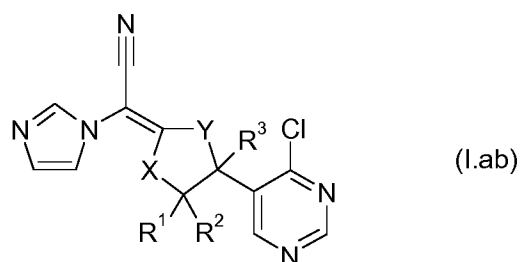
aa) 28 compounds of formula (I.aa):



5

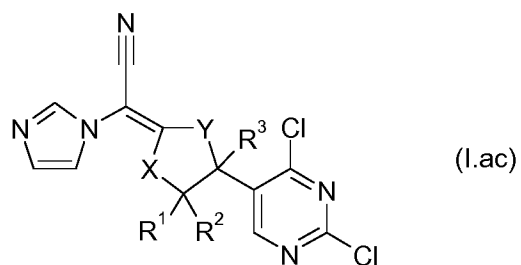
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

ab) 28 compounds of formula (I.ab):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

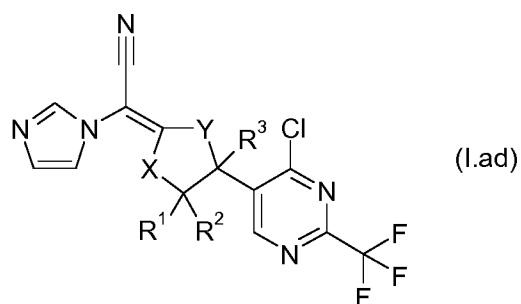
ac) 28 compounds of formula (I.ac):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

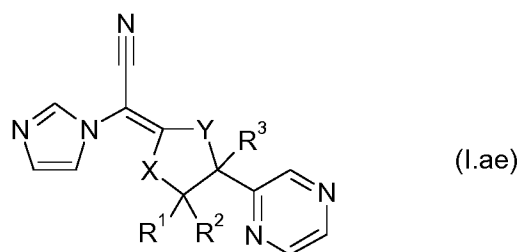
15

ad) 28 compounds of formula (I.ad):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

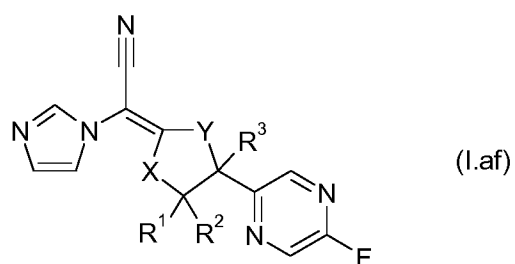
ae) 28 compounds of formula (I.ae):



5

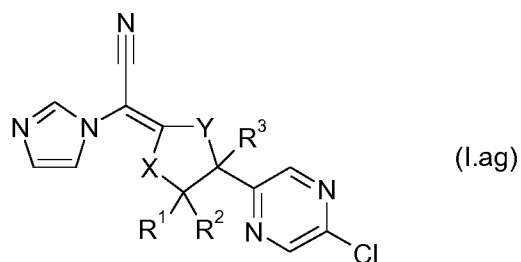
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

af) 28 compounds of formula (I.af):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

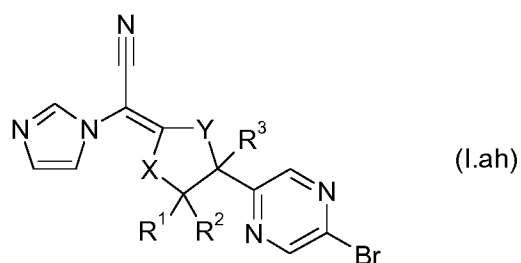
ag) 28 compounds of formula (I.ag):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

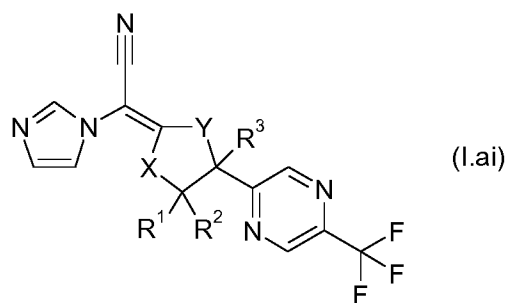
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ah) 28 compounds of formula (I.ah):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

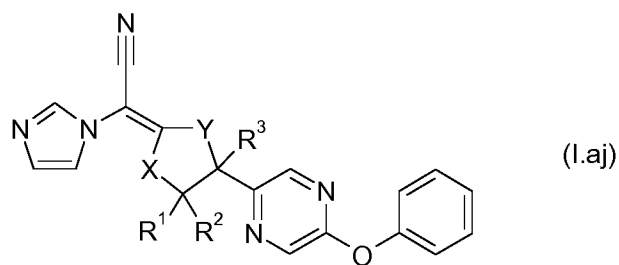
ai) 28 compounds of formula (I.ai):



5

wherein X, Y, R¹, R² and R³ are as defined in Table 1.

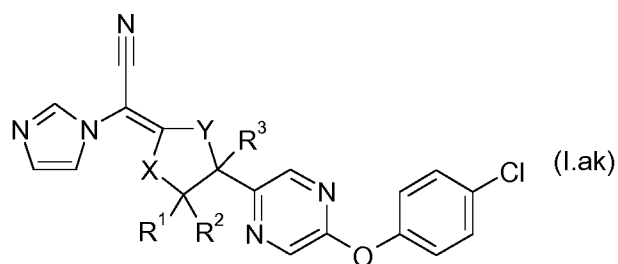
aj) 28 compounds of formula (I.aj):



10

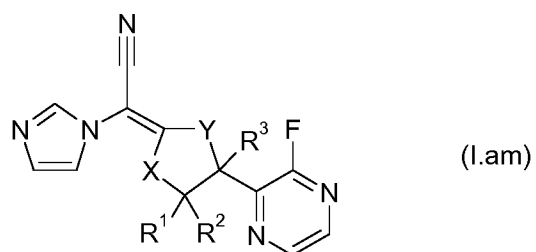
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

ak) 28 compounds of formula (I.ak):



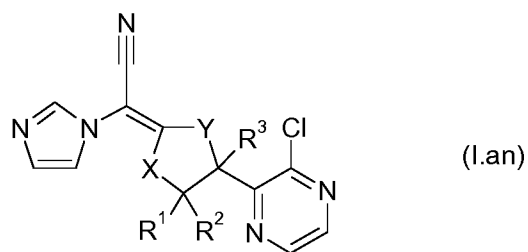
15 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

am) 28 compounds of formula (I.am):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

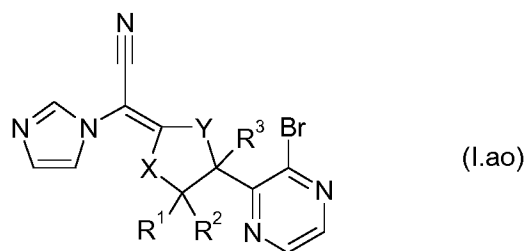
an) 28 compounds of formula (I.an):



5

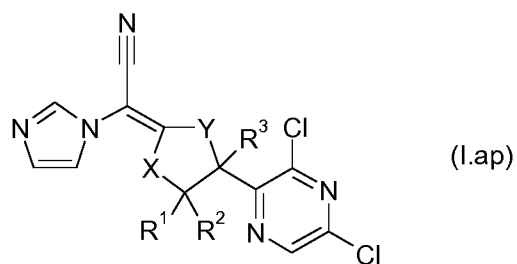
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

ao) 28 compounds of formula (I.ao):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

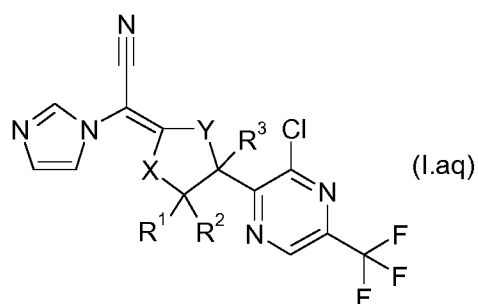
ap) 28 compounds of formula (I.ap):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

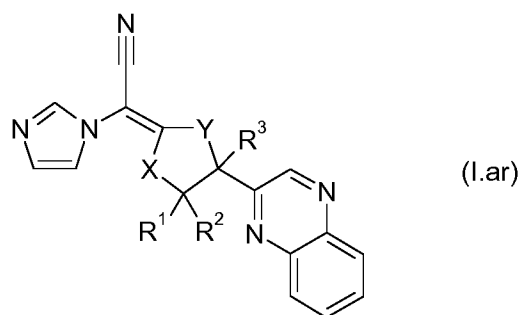
15

aq) 28 compounds of formula (I.aq):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

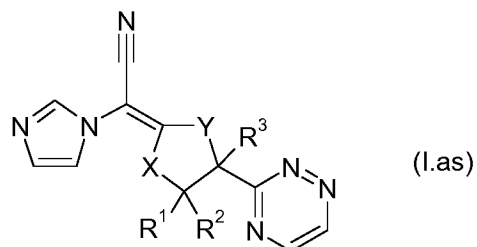
ar) 28 compounds of formula (I.ar):



5

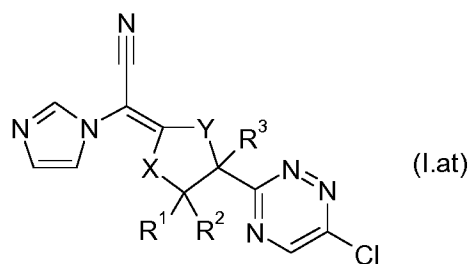
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

as) 28 compounds of formula (I.as):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

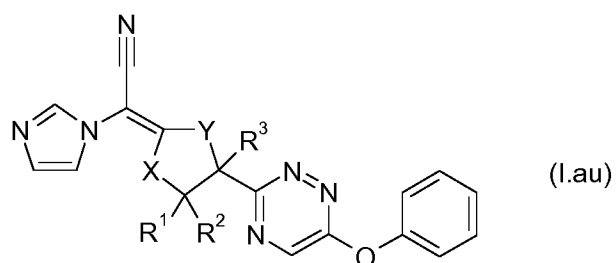
at) 28 compounds of formula (I.at):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

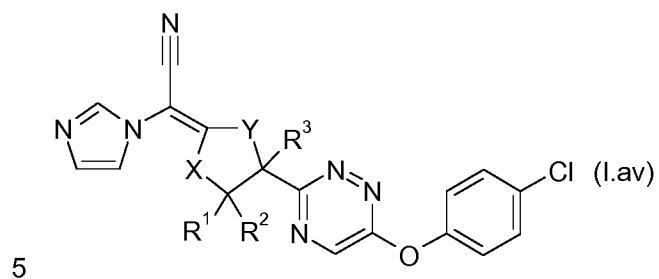
15

au) 28 compounds of formula (I.au):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

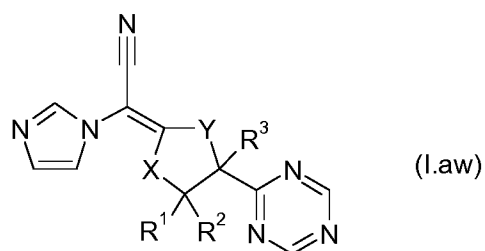
av) 28 compounds of formula (I.av):



5

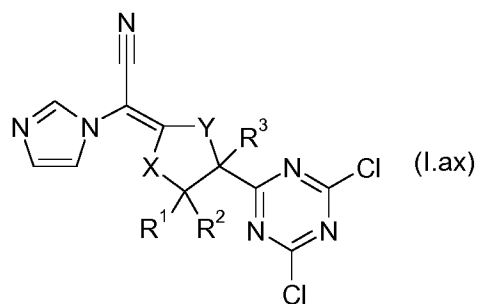
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

aw) 28 compounds of formula (I.aw):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

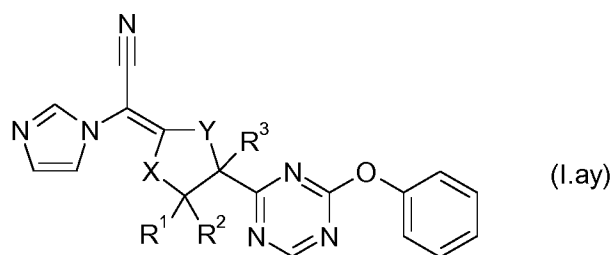
ax) 28 compounds of formula (I.ax):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

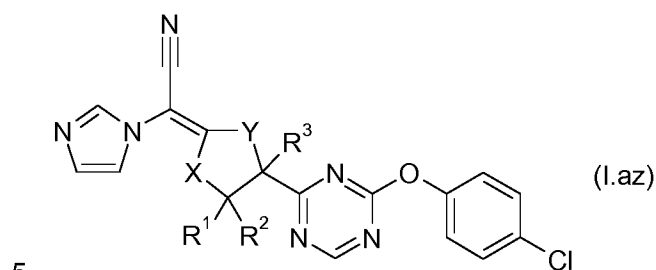
15

ay) 28 compounds of formula (I.ay):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

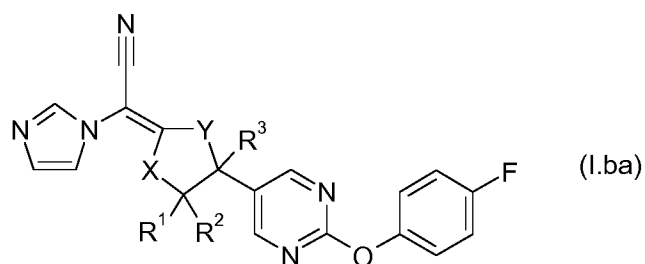
az) 28 compounds of formula (I.az):



5

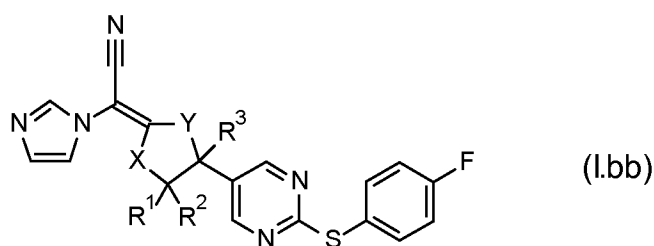
wherein X, Y, R¹, R² and R³ are as defined in Table 1.

ba) 28 compounds of formula (I.ba):



10 wherein X, Y, R¹, R² and R³ are as defined in Table 1.

bb) 28 compounds of formula (I.bb):



wherein X, Y, R¹, R² and R³ are as defined in Table 1.

15

Throughout this description, temperatures are given in degrees Celsius and "m.p." means melting point. LC/MS means Liquid Chromatography Mass Spectroscopy and the description of the apparatus and the method is: (ACQUITY UPLC from Waters, Phenomenex Gemini C18, 3 μ m particle size (3 micrometer particle size), 110 Angström, 30 x 3 mm column, 1.7mL/min., 60 °C, H₂O + 0.05% HCOOH (95%) / CH₃CN/MeOH 4:1 + 0.04% HCOOH (5%) – 2 min. – CH₃CN/MeOH 4:1 + 0.04%

20

HCOOH (5%) – 0.8 min., ACQUITY SQD Mass Spectrometer from Waters, ionization method: electrospray (ESI), Polarity: positive ions, Capillary (kV) 3.00, Cone (V) 20.00, Extractor (V) 3.00, Source Temperature (°C) 150, Desolvation Temperature (°C) 400, Cone Gas Flow (L/Hr) 60, Desolvation Gas Flow (L/Hr) 700).

5

Table 2: Melting point and LC/MS data for compounds of Table 1

Compound No.	Melting point (°C)	LC/MS
I.k.25	185 - 186	
I.r.25		Rt = 0.50 min; MS: m/z = 306 (M+1)
I.v.25	54 - 59	
I.ao.25		Rt = 0.66 min; MS: m/z = 366 (M+1)
I.aa.25		Rt = 0.87 min; MS: m/z = 416 (M+1)
I.ae.25		Rt = 0.27 min; MS: m/z = 288 (M+1)
I.bb.25		Rt = 1.19 min; MS: m/z = 414 (M+1)

10 Biological examples

Alternaria solani / tomato / leaf disc (early blight)

Tomato leaf disks cv. Baby are placed on agar in multiwell plates (24-well format) and sprayed with the formulated test compound diluted in water. The leaf disks are inoculated with a spore suspension of the fungus 2 days after application. The inoculated leaf disks are incubated at 23 °C / 21°C (day/night) and 80% rh under a light regime of 12/12 h (light/dark) in a climate cabinet and the activity of a compound is assessed as percent disease control compared to untreated when an appropriate level of disease damage appears on untreated check disk leaf disks (5 – 7 days after application).

Compounds I.k.025, I.r.25, I.ao.25, I.aa.25 and I.bb.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

Blumeria graminis f. sp. tritici (Erysiphe graminis f. sp. tritici) / wheat / leaf disc preventative (Powdery mildew on wheat)

Wheat leaf segments cv. Kanzler are placed on agar in a multiwell plate (24-well format) and sprayed with the formulated test compound diluted in water. The leaf disks are inoculated by shaking powdery mildew infected plants above the test plates 1 day after application. The inoculated leaf disks are incubated at 20°C and 60% rh under a light regime of 24 h darkness followed by 12 h light / 12 h darkness in a climate chamber and the activity of a compound is assessed as percent disease control compared to untreated when an appropriate level of disease damage appears on untreated check leaf segments (6 – 8 days after application).

Compounds I.k.025, I.r.25, I.v.25, I.a.o.25, I.a.a.25, I.a.e.25 and I.b.b.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

5 *Botryotinia fuckeliana (Botrytis cinerea)* / liquid culture (Gray mould)

Conidia of the fungus from cryogenic storage are directly mixed into nutrient broth (Vogels broth). After placing a (DMSO) solution of test compound into a microtiter plate (96-well format), the nutrient broth containing the fungal spores is added. The test plates are incubated at 24°C and the inhibition of growth is determined photometrically 3-4 days after application.

- 10 Compounds I.k.025, I.r.25, I.v.25, I.a.o.25, I.a.a.25 and I.b.b.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

Fusarium culmorum / liquid culture (Head blight)

- 15 Conidia of the fungus from cryogenic storage are directly mixed into nutrient broth (PDB potato dextrose broth). After placing a (DMSO) solution of test compound into a microtiter plate (96-well format), the nutrient broth containing the fungal spores is added. The test plates are incubated at 24°C and the inhibition of growth is determined photometrically 3-4 days after application.

- 20 Compounds I.k.25, I.r.25, I.v.25, I.a.o.25, I.a.a.25 and I.b.b.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

Gaeumannomyces graminis / liquid culture (Take-all of cereals)

- 25 Mycelial fragments of the fungus from cryogenic storage were directly mixed into nutrient broth (PDB potato dextrose broth). After placing a (DMSO) solution of test compound into a microtiter plate (96-well format), the nutrient broth containing the fungal spores is added. The test plates are incubated at 24°C and the inhibition of growth is determined photometrically 4-5 days after application.

- 30 Compounds I.k.25, I.r.25 and I.a.a.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

Glomerella lagenarium (Colletotrichum lagenarium) / liquid culture (Anthracnose)

- 35 Conidia of the fungus from cryogenic storage are directly mixed into nutrient broth (PDB potato dextrose broth). After placing a (DMSO) solution of test compound into a microtiter plate (96-well format), the nutrient broth containing the fungal spores is added. The test plates are incubated at 24°C and the inhibition of growth is measured photometrically 3-4 days after application.

Compounds I.k.025, I.r.25, I.a.o.25, I.a.a.25 and I.b.b.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

40

Magnaporthe grisea (Pyricularia oryzae) / rice / leaf disc preventative (Rice Blast)

Rice leaf segments cv. Ballila are placed on agar in a multiwell plate (24-well format) and sprayed with the formulated test compound diluted in water. The leaf segments are inoculated with a spore suspension of the fungus 2 days after application. The inoculated leaf segments are incubated at 22°C and 80% rh under a light regime of 24 h darkness followed by 12 h light / 12 h darkness in a climate cabinet and the activity of a compound is assessed as percent disease control compared to untreated when an appropriate level of disease damage appears in untreated check leaf segments (5 – 7 days after application).

Compounds I.k.025 and I.bb.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

Monographella nivalis (*Microdochium nivale*) / liquid culture (foot rot cereals)

Conidia of the fungus from cryogenic storage are directly mixed into nutrient broth (PDB potato dextrose broth). After placing a (DMSO) solution of test compound into a microtiter plate (96-well format), the nutrient broth containing the fungal spores is added. The test plates are incubated at 24°C and the inhibition of growth is determined photometrically 4-5 days after application.

Compounds I.v.25, I.ao.25 and I.bb.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

20

Mycosphaerella arachidis (*Cercospora arachidicola*) / liquid culture (early leaf spot)

Conidia of the fungus from cryogenic storage are directly mixed into nutrient broth (PDB potato dextrose broth). After placing a (DMSO) solution of test compound into a microtiter plate (96-well format), the nutrient broth containing the fungal spores is added. The test plates are incubated at 24°C and the inhibition of growth is determined photometrically 4-5 days after application.

Compounds I.k.025, I.r.25, I.v.25, I.ao.25, I.aa.25, I.ae.25 and I.bb.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

30 *Mycosphaerella graminicola* (*Septoria tritici*) / liquid culture (Septoria blotch)

Conidia of the fungus from cryogenic storage are directly mixed into nutrient broth (PDB potato dextrose broth). After placing a (DMSO) solution of test compound into a microtiter plate (96-well format), the nutrient broth containing the fungal spores is added. The test plates are incubated at 24°C and the inhibition of growth is determined photometrically 4-5 days after application.

Compounds I.k.025, I.r.25, I.v.25, I.ao.25, I.aa.25 and I.bb.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

Pyrenophora teres / barley / leaf disc preventative (Net blotch)

Barley leaf segments cv. Hasso are placed on agar in a multiwell plate (24-well format) and sprayed with the formulated test compound diluted in water. The leaf segments are inoculated with a spore

suspension of the fungus 2 days after application. The inoculated leaf segments are incubated at 20°C and 65% rh under a light regime of 12 h light / 12 h darkness in a climate cabinet and the activity of a compound is assessed as disease control compared to untreated when an appropriate level of disease damage appears in untreated check leaf segments (5 – 7 days after application).

- 5 Compounds I.k.025, I.r.25, I.v.25, I.ao.25, I.aa.25 and I.bb.25 at 200 ppm give at least 80% disease control in this test when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

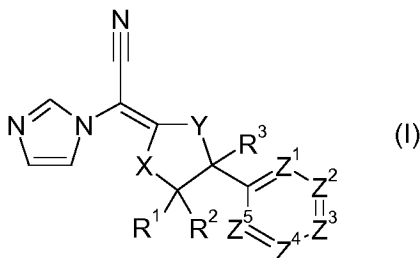
Thanatephorus cucumeris (Rhizoctonia solani) / liquid culture (foot rot, damping-off)

- 10 Mycelia fragments of a newly grown liquid culture of the fungus are directly mixed into nutrient broth (PDB potato dextrose broth). After placing a (DMSO) solution of the test compounds into a microtiter plate (96-well format), the nutrient broth containing the fungal material is added. The test plates are incubated at 24°C and the inhibition of growth is determined photometrically 3-4 days after application.

- Compounds I.r.25, I.v.25 and I.aa.25 at 200 ppm give at least 80% disease control in this test
15 when compared to untreated control leaf disks under the same conditions, which show extensive disease development.

Claims

- 5 1. A compound of formula (I):



wherein,

X and Y are independently O, S or NR⁴;

- Z¹, Z², Z³, Z⁴ and Z⁵ are independently CR⁵ or N; optionally Z¹ and Z², Z² and Z³, Z³ and Z⁴ or Z⁴ and
 10 Z⁵ may also belong to a second, annulated saturated, partially unsaturated or aromatic isocyclic or heterocyclic ring to form a bicyclic ring system,
 with the proviso, that at least two out of Z¹, Z², Z³, Z⁴ and Z⁵ are N;
 R¹, R² and R³ are independently hydrogen, halogen, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy;

- 15 R⁵ is hydrogen, halogen, cyano, hydroxyl, amino, nitro, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-C₆cycloalkyl, C₃-C₆halocycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₁-C₆alkylthio, C₁-C₆alkylsulfinyl, C₁-C₆alkylsulfonyl, C₁-C₆alkylcarbonyl, C₂-C₆alkenyl, C₂-C₆haloalkenyl, C₂-C₆alkenyloxy, C₂-C₆haloalkenyloxy, C₂-C₆alkynyl, C₂-C₆alkynyloxy, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or heteroaryl substituted by 1 to 3 substituents R⁶,
 20 unsubstituted aryloxy or aryloxy substituted by 1 to 3 substituents R⁶, unsubstituted phenylthio or phenylthio substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryloxy or heteroaryloxy substituted by 1 to 3 substituents R⁶, unsubstituted arylC₁-C₆alkyl or heteroarylC₁-C₆alkyl substituted by 1 to 3 substituents R⁶;
 R⁶ is independently hydroxyl, amino, nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy C₁-C₆haloalkyl, C₁-C₆haloalkoxy, aryl, aryloxy, or arylC₁-C₆alkyl, in which the alkyl, alkoxy and aryl groups can be substituted by halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy;
 25 or a salt or a N-oxide thereof.

2. A compound of formula (I) according to claim 1 wherein X and Y are independently O or S.
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3. A compound of formula (I) according to claim 1 or claim 2 wherein R¹, R² and R³ are independently hydrogen, halogen, C₁-C₆alkyl or C₁-C₆haloalkyl.

4. A compound of formula (I) according to anyone of the claims 1 to 3 wherein
 35 the ring formed by Z¹, Z², Z³, Z⁴ and Z⁵ is selected from pyridazinyl, cinnolinyl, pyrimidinyl, pyrazinyl, triazinyl each of can be substituted by hydrogen, halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₃-

C₆cycloalkyl, C₁-C₆alkoxy, C₁-C₆haloalkoxy, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted aryl or aryl substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryl or heteroaryl substituted by 1 to 3 substituents R⁶, unsubstituted aryloxy or aryloxy substituted by 1 to 3 substituents R⁶, unsubstituted phenylthio or phenylthio substituted by 1 to 3 substituents R⁶, unsubstituted heteroaryloxy or heteroaryloxy substituted by 1 to 3 substituents R⁶, unsubstituted arylC₁-C₆alkyl or heteroarylC₁-C₆alkyl substituted by 1 to 3 substituents R⁶; R⁶ is independently hydroxyl, amino, nitro, cyano, halogen, C₁-C₆alkyl, C₁-C₆alkoxy, C₁-C₆haloalkyl, C₁-C₆haloalkoxy, aryl, aryloxy, or arylC₁-C₆alkyl, in which the alkyl, alkoxy and aryl groups can be substituted by halogen, cyano, C₁-C₆alkyl, C₁-C₆haloalkyl, C₁-C₆alkoxy or C₁-C₆haloalkoxy.

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5. A compound of formula (I) according to anyone of the claims 1 to 4 wherein
X and Y are each S;
R¹, R² and R³ are all hydrogen.

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6. A compound of formula (I) according to to anyone of the claims 1 to 5 wherein
the ring formed by Z¹, Z², Z³, Z⁴ and Z⁵ are selected from pyridazinyl, cinnolinyl, pyrimidinyl, pyrazinyl, triazinyl each of the pyridazinyl, cinnolinyl, pyrimidinyl, pyrazinyl, triazinyl can be substituted by F, Cl, Br, C₁-C₆haloalkyl, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted phenox or substituted phenoxy substituted by F, Cl, unsubstituted phenylthio or substituted phenylthio substituted by F

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7. A compound of formula (I) according to anyone of the claims 1 to 6 wherein
the ring formed by Z¹, Z², Z³, Z⁴ and Z⁵ are selected from pyrimidinyl, pyrazinyl each of the from pyrimidinyl, pyrazinyl can be substituted by F, Cl, Br, C₁-C₆haloalkyl, C₃-C₆cycloalkyl-C₂-C₆alkynyl, unsubstituted phenoxy or phenoxy substituted by F, Cl, unsubstituted phenylthio or substituted

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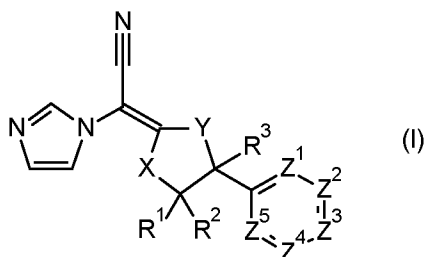
phenylthio substituted by F.

8. A method for controlling or preventing infestation of plants or plant propagation material and/or harvested crops susceptible to microbial attack, characterized in that an effective amount a compound of formula (I) as defined in anyone of the claims 1 to 7 is applied to the plants or plant propagation material and/or harvested food, or to parts thereof or the locus thereof.

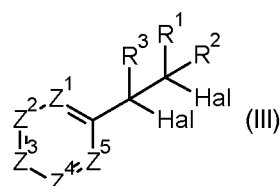
9. A method of controlling phytopathogenic diseases on plants or plant propagation material and/or harvested crops susceptible to microbial attack, characterized in that an effective amount a compound of formula (I) as defined in anyone of the claims 1 to 7 is applied to the plants or plant propagation material and/or harvested food, or to parts thereof or the locus thereof.

10. A composition comprising a fungicidally effective amount of a compound of formula (I) as defined in anyone of the claims 1 to 7.

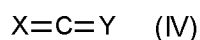
11. A process for the preparation of a compound of formula (I)



comprising reacting imidazol-1-yl-acetonitrile with compound of formula (III)



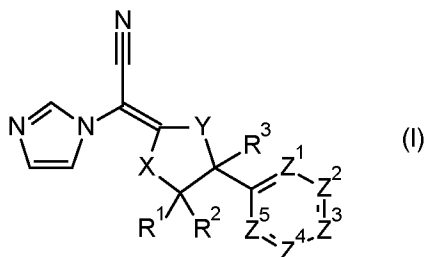
and with a compound of formula (IV)



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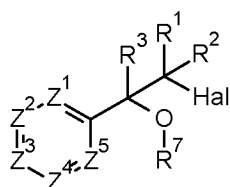
In the presence of a base, wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 are as defined as in any of the claims 1 to 7 and Hal is halogen.

12. A process for the preparation of a compound of formula (I)

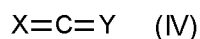


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comprising reacting imidazol-1-yl-acetonitrile with compound of formula (XI)



and with a compound of formula (IV)



15 In the presence of a base, wherein Z^1 , Z^2 , Z^3 , Z^4 , Z^5 , R^1 , R^2 and R^3 are as defined as in any of the claims 1 to 7, R^7 is a sulfonyl group, and Hal is halogen.

13. Compounds obtainable by a process according to claim 11 or claim 12.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2015/058965

A. CLASSIFICATION OF SUBJECT MATTER

INV. C07D405/14 C07D413/14 A01N43/00 C07D403/14 C07D409/14
C07D417/14

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

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, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 218 736 A1 (NIHON NOHYAKU CO LTD [JP]) 22 April 1987 (1987-04-22) claim 1 and test example 1 -----	1-13
A	US 4 738 976 A (SEO AKIRA [JP] ET AL) 19 April 1988 (1988-04-19) claim 1 -----	1-13
A,P	WO 2015/003991 A1 (SYNGENTA PARTICIPATIONS AG [CH]) 15 January 2015 (2015-01-15) claims 1 and -----	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

20 May 2015

Date of mailing of the international search report

27/05/2015

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2015/058965

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0218736	A1	22-04-1987	NONE
US 4738976	A	19-04-1988	NONE
WO 2015003991	A1	15-01-2015	NONE