

(12) PATENT
(19) AUSTRALIAN PATENT OFFICE

(11) Application No. **AU 199675661 B2**
(10) Patent No. **711483**

(54) Title
Biphenyl ether oxazolines and their use as pest-control agents

(51)⁶ International Patent Classification(s)
C07D 263/14 C07D 413/12
A01N 043/76

(21) Application No: **199675661** (22) Application Date: **1996.11.06**

(87) WIPO No: **WO97/19067**

(30) Priority Data

(31) Number	(32) Date	(33) Country
19542934	1995.11.17	DE
19638047	1996.09.18	DE

(43) Publication Date : **1997.06.11**

(43) Publication Journal Date : **1997.08.07**

(44) Accepted Journal Date : **1999.10.14**

(71) Applicant(s)
Bayer Aktiengesellschaft

(72) Inventor(s)
Wolfgang Kramer; Udo Kraatz; Christoph Erdelen; Ulrike Wachendorff-Neumann; Andreas Turberg; Norbert Mencke

(74) Agent/Attorney
DAVIES COLLISON CAVE, GPO Box 3876, SYDNEY NSW 2001

(56) Related Art
AU 28445/95
AU 42437/96

OPI DATE 11/06/97 APPLN. ID 75661/96
 AOJP DATE 07/08/97 PCT NUMBER PCT/EP96/04846



AU9675661

(51) Internationale Patentklassifikation ⁶ : C07D 263/14, A01N 43/76, C07D 413/12		A1	(11) Internationale Veröffentlichungsnummer: WO 97/19067
		(43) Internationales Veröffentlichungsdatum:	29. Mai 1997 (29.05.97)
(21) Internationales Aktenzeichen: PCT/EP96/04846		MENCKE, Norbert [DE/DE]; Grundermühle 2, D-51381 Leverkusen (DE).	
(22) Internationales Anmeldedatum: 6. November 1996 (06.11.96)		(74) Gemeinsamer Vertreter: BAYER AKTIENGESELLSCHAFT; D-51368 Leverkusen (DE).	
(30) Prioritätsdaten: 195 42 934.6 17. November 1995 (17.11.95) DE 196 38 047.2 18. September 1996 (18.09.96) DE		(81) Bestimmungsstaaten: AU, BB, BG, BR, BY, CA, CN, CZ, HU, JP, KR, KZ, LK, MX, NO, NZ, PL, RO, RU, SK, TR, UA, US, europäisches Patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI Patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).	
(71) Anmelder (für alle Bestimmungsstaaten ausser US): BAYER AKTIENGESELLSCHAFT [DE/DE]; D-51368 Leverkusen (DE).		Veröffentlicht Mit internationalem Recherchenbericht.	
(71) Anmelder (nur für CN JP KR): YASHIMA CHEMICAL INDUSTRY CO., LTD. [JP/JP]; YTT Building 757-1, Takatsu-ku, Kawasaki-shi, Kanagawa 213 (JP).			
(72) Erfinder; und (75) Erfinder/Anmelder (nur für US): KRÄMER, Wolfgang [DE/DE]; Rosenkranz 25, D-51399 Burscheid (DE). KRAATZ, Udo [DE/DE]; Andreasstrasse 22a, D-51375 Leverkusen (DE). ERDELEN, Christoph [DE/DE]; Unterbüscherhof 15, D-42799 Leichlingen (DE). WACHENDORFF-NEUMANN, Ulrike [DE/DE]; Oberer Markweg 85, D-56566 Neuwied (DE). TURBERG, Andreas [DE/DE]; Naheweg 19, D-40699 Erkrath (DE).			
(54) Title: BIPHENYL ETHER OXAZOLINES AND THEIR USE AS PEST-CONTROL AGENTS			
(54) Bezeichnung: BIPHENYLETHER-OXAZOLINE UND IHRE VERWENDUNG ALS SCHÄDLINGSBEKÄMPFUNGSMITTEL			
(57) Abstract			
<p>The invention concerns biphenyl ether oxazolines of formula (I), in which X, Y, Z, R¹, R², A, m and n are as defined in the description, methods of preparing them, intermediates used in their preparation and their use in the control of animal pests.</p>		<p style="text-align: right;">(1)</p>	
(57) Zusammenfassung			
<p>Die vorliegende Erfindung betrifft neue Biphenylether-oxazoline der Formel (I), in welcher X, Y, Z, R¹, R², A, m und n die in der Beschreibung angegebene Bedeutung haben, Verfahren zu ihrer Herstellung und ihre Verwendung zur Bekämpfung von tierischen Schädlingen sowie neue Zwischenprodukte.</p>			

ONLY FOR INFORMATION

Codes used to identify the PCT member States on the flyleaves of the brochures in which international applications made under the PCT are published.

AM	Armenia	KZ	Kazakhstan
AT	Austria	LI	Liechtenstein
AU	Australia	LK	Sri Lanka
BB	Barbados	LR	Liberia
BE	Belgium	LK	Lithuania
BF	Burkina Fasso	LU	Luxembourg
BG	Bulgaria	LV	Latvia
BJ	Benin	MC	Monaco
BR	Brazil	MD	Republic of Moldova
BY	Belarus	MG	Madagascar
CA	Canada	ML	Mali
CF	Central African Republic	MN	Mongolia
CG	Congo	MR	Mauritania
CH	Switzerland	MW	Malawi
CI	Ivory Coast	MX	Mexico
CM	Cameroon	NE	Niger
CN	China	NL	Netherlands
CS	Czechoslovakia	NO	Norway
CZ	Czech Republic	NZ	New Zealand
DE	Germany	PL	Poland
DK	Denmark	PT	Portugal
EE	Estonia	RO	Romania
ES	Spain	RU	Russian Federation
FI	Finland	SD	Sudan
FR	France	SE	Sweden
GA	Gabon	SG	Singapore
GB	United Kingdom	SI	Slovenia
GE	Georgia	SK	Slovakia
GN	Guinea	SN	Senegal
GR	Greece	SZ	Swaziland
HU	Hungary	TD	Chad
IE	Ireland	TG	Togo
IT	Italy	TJ	Tajikistan
JP	Japan	TT	Trinidad and Tobago
KE	Kenya	UA	Ukraine
KG	Kyrgyzstan	UG	Uganda
KP	Democratic People's Republic of Korea	US	United States of America
KR	Republic of Korea	UZ	Uzbekistan
		VN	Vietnam

- 1 -

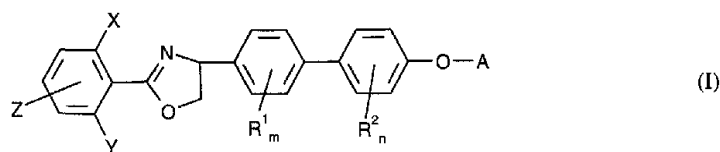
BIPHENYL ETHEROXAZOLINES AND THEIR USE AS PEST CONTROL AGENTS

The present invention relates to novel biphenyl ether-oxazolines, processes for their preparation and their use for controlling animal pests, and to novel intermediates.

- 5 It is already known that certain substituted biphenyl-oxazolines, such as, for example, 2-(2,6-difluorophenyl)-4-(4'-methoxybiphenyl-4-yl)-1,3-oxazoline, have an insecticidal and acaricidal action (cf. EP-A 0 432 661).

However, the level of action and/or duration of action of these known compounds is not completely satisfactory in all fields of use, especially against certain organisms or
 10 if low concentrations are used.

Novel biphenyl ether-oxazolines of the formula (I)



in which

- X represents hydrogen, fluorine or chlorine,
- 15 Y represents fluorine, chlorine or methyl,
- Z represents hydrogen, halogen, alkyl, alkoxy or dialkylamino,
- R¹ and R² independently of one another represent halogen, alkyl, alkoxy, alkylthio, halogenoalkyl, halogenoalkoxy or halogenoalkylthio,
- m and n independently of one another represent 0, 1 or 2 and

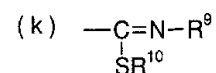
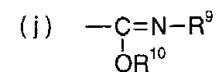
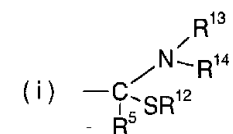
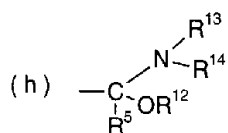
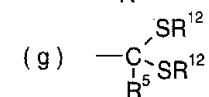
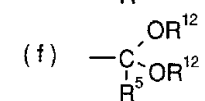
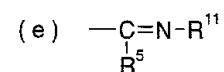
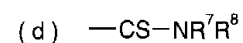
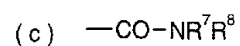
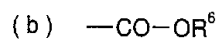
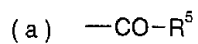


A represents the grouping $-(CH_2)_p-(CR^3R^4)_q-(CH_2)_r-R$ wherein

R^3 and R^4 independently of one another represent hydrogen or alkyl,

p , q and r independently of one another represent 0, 1, 2 or 3, at least one index being other than 0, and

5 R represents cyano; or represents an optionally substituted, saturated, partly saturated or unsaturated 5- or 6-membered heterocyclic radical, or represents one of the following groupings:



wherein

R⁵ represents hydrogen, alkyl, halogenoalkyl, alkenyl, halogenoalkenyl,
optionally substituted cycloalkyl or optionally substituted aryl,

5

R⁶ represents hydrogen, alkyl, alkenyl, halogenoalkyl, halogenoalkenyl,
optionally substituted arylalkyl or in each case optionally substituted
cycloalkyl or cycloalkylalkyl,

10

R⁷ and R⁸ independently of one another represent hydrogen, alkyl, alkoxy,
alkenyl, halogenoalkyl, halogenoalkenyl, in each case optionally substituted
aryl or arylalkyl, or in each case optionally substituted cycloalkyl or
cycloalkylalkyl, or represent -OR⁶ or -NR⁵R⁶,

15

wherein

R⁵ and R⁶ have the abovementioned meaning, or

20

R⁷ and R⁸ together represent a 5- or 6-membered alkylene chain, which optionally
contains an oxygen atom,

R⁹ and R¹⁰ independently of one another represent alkyl,

25

R¹¹ represents -OR⁶, -NR⁵R⁶ or -N(R⁵)-COOR⁶,

wherein

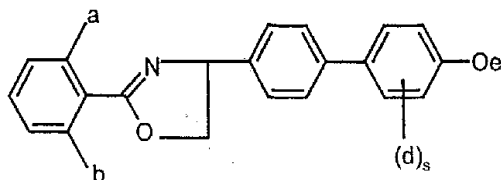
R⁵ and R⁶ have the abovementioned meaning, and

30

R¹², R¹³ and R¹⁴ independently of one another represent alkyl,

provided that compounds of the formula

35



wherein

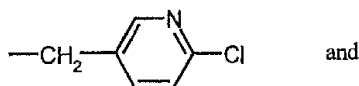
- a represents hydrogen,
- b represents fluorine or chlorine,
- 5 c represents optionally substituted heteroarylalkyl,
- d represents halogen, alkyl or alkoxy and
- s represents 0, 1 or 2

or

10 wherein

- a represents fluorine,
- b represents fluorine

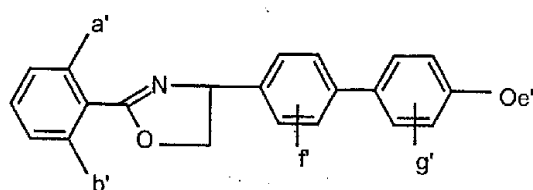
- 15 e represents
- s represents 0,



are specifically excluded

and that the compounds of the formula

20



25

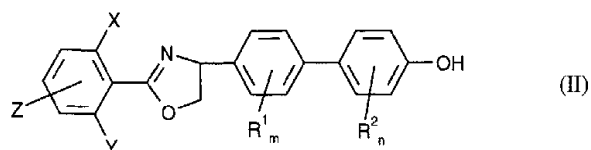
wherein

- a' represents hydrogen, fluorine or chlorine
- b' represents fluorine and chlorine
- e' represents C₂-C₄-cyanoalkyl
- 30 f' represents hydrogen, halogen, alkyl, alkoxy, halogenoalkyl, halogenoalkoxy
- and
- g' represents hydrogen, alkyl, alkoxy, halogenoalkyl, halogenoalkoxy,
- are specifically excluded.



The biphenyl ether-oxazolines of the formula (I) can be obtained as optical and/or geometric isomers, also depending on the substituents. The present invention relates both to the isomer mixtures and to the pure isomers.

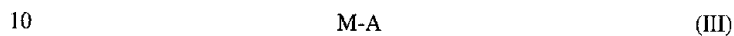
It has furthermore been found that the biphenyl ether-oxazolines of the formula (I) are
5 obtained by a process in which hydroxybiphenyloxazolines of the formula (II)



in which

X, Y, Z, R¹, R², m and n have the abovementioned meaning.

are reacted with a compound of the formula (III)



in which

A has the abovementioned meaning and

M represents a leaving group,

if appropriate in the presence of a base and/or a catalyst and if appropriate in the
15 presence of a diluent.

It has furthermore been found that the novel substituted biphenyl ether-oxazolines of the formula (I) are particularly suitable for controlling animal pests, in particular insects, arachnids and nematodes which occur in agriculture, in forestry, in the preservation of stored products and materials and in the hygiene sector.



Formula (I) provides a general definition of the compounds according to the invention.

Preferred substituents and ranges of the radicals listed in the formulae mentioned above and below are explained in the following.

X preferably represents hydrogen, fluorine or chlorine.

5 Y preferably represents fluorine, chlorine or methyl.

Z preferably represents hydrogen, fluorine, chlorine, C₁-C₄-alkyl, C₁-C₄-alkoxy or di(C₁-C₄)alkylamino.

10 R¹ and R² independently of one another preferably represent fluorine, chlorine, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy or C₁-C₄-halogenoalkylthio.

m and n independently of one another preferably represent 0, 1 or 2.

A preferably represents the grouping (CH₂)_p-(CR³R⁴)_q-(CH₂)_r-R,
wherein

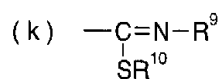
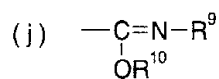
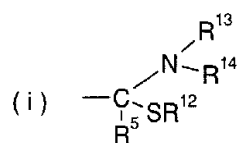
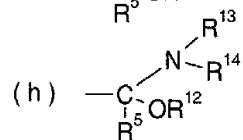
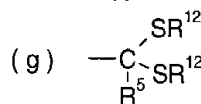
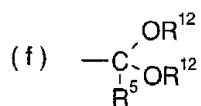
R³ and R⁴ independently of one another represent hydrogen or C₁-C₄-alkyl,

15 p, q and r independently of one another represent 0, 1, 2 or 3, at least one index being other than 0 and the sum of the indices being not greater than 5.

20 R preferably represents cyano; or represents a saturated, partly saturated or unsaturated 5- or 6-membered heterocyclic radical which has 1 to 3 identical or different heteroatoms from the series consisting of nitrogen, oxygen and sulfur and is optionally substituted by halogen, C₁-C₄-alkyl or C₁-C₄-halogenoalkyl, or represents one of the following groupings:



- (a) $-\text{CO}-\text{R}^5$
- (b) $-\text{CO}-\text{OR}^6$
- (c) $-\text{CO}-\text{NR}^7\text{R}^8$
- (d) $-\text{CS}-\text{NR}^7\text{R}^8$
- (e) $-\underset{\text{R}^5}{\text{C}}=\text{N}-\text{R}^{11}$



wherein

R^5 represents hydrogen, $\text{C}_1\text{-C}_6\text{-alkyl}$, $\text{C}_1\text{-C}_4\text{-halogenoalkyl}$, $\text{C}_2\text{-C}_6\text{-alkenyl}$ or $\text{C}_2\text{-C}_6\text{-halogenoalkyl}$; or $\text{C}_3\text{-C}_6\text{-cycloalkyl}$ which is optionally substituted once or several times in an identical or different manner by $\text{C}_1\text{-C}_4\text{-alkyl}$, halogen or $\text{C}_1\text{-C}_4\text{-halogenoalkyl}$,
or represents phenyl which is optionally substituted once to five times in an identical or different manner by halogen, $\text{C}_1\text{-C}_4\text{-alkyl}$, $\text{C}_1\text{-C}_4\text{-alkoxy}$, $\text{C}_1\text{-C}_4\text{-alkylthio}$, $\text{C}_1\text{-C}_4\text{-halogenoalkyl}$, $\text{C}_1\text{-C}_4\text{-halogenoalkoxy}$,



C₁-C₄-halogenoalkylthio, di(C₁-C₄)alkylamino, C₁-C₄-alkylcarbonyl-amino, C₁-C₄-alkylcarbonyl-C₁-C₄-alkylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷,

5 R⁶ represents hydrogen, C₁-C₆-alkyl, C₁-C₆-halogenoalkyl, C₂-C₆-alkenyl or C₂-C₆-halogenoalkenyl; or C₃-C₆-cycloalkyl or C₃-C₆-cycloalkyl-C₁-C₄-alkyl, in each case optionally substituted once or several times in an identical or different manner by C₁-C₄-alkyl, halogen or C₁-C₄-halogenoalkyl,

10 or represents C₆-C₁₀-aryl-C₁-C₆-alkyl which is optionally substituted once to five times in an identical or different manner by halogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy, C₁-C₄-halogenoalkylthio, di(C₁-C₄)alkylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷,

15 R⁷ and R⁸ independently of one another represent hydrogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-halogenoalkyl, C₂-C₆-alkenyl or C₂-C₆-halogenoalkenyl; or C₃-C₆-cycloalkyl or C₃-C₆-cycloalkyl-C₁-C₄-alkyl, in each case optionally substituted once or several times in an identical or different manner by C₁-C₄-alkyl, halogen or C₁-C₄-halogenoalkyl,

20 or represent phenyl or phenyl-C₁-C₄-alkyl, in each case optionally substituted once to five times in an identical or different manner by halogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy, C₁-C₄-halogenoalkylthio, di(C₁-C₄)alkylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷, or represent -OR⁶ or -NR⁵R⁶,

25 or

R⁷ and R⁸ together represent -(CH₂)₅-, -(CH₂)₆- or -(CH₂)₂-O-(CH₂)₂-;

R⁹ and R¹⁰ independently of one another represent C₁-C₄-alkyl;



R¹¹ represents -OR⁶-, -NR⁵R⁶ or -N(R⁵)-COOR⁶,

and

R¹², R¹³ and R¹⁴ independently of one another represent C₁-C₄-alkyl.

5 R¹⁵ preferably represents hydrogen, C₁-C₄-alkyl or C₁-C₄-halogenoalkyl; or C₃-C₆-cycloalkyl which is optionally substituted once or several times in an identical or different manner by C₁-C₄-alkyl, halogen or C₁-C₄-halogenoalkyl, or represents phenyl which is optionally substituted once to five times in an identical or different manner by halogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy, C₁-C₄-halogenoalkylthio, 10 di(C₁-C₄)alkylamino, cyano, nitro, C₁-C₆-alkoxycarbonyl or di(C₁-C₆)alkylaminocarbonyl.

R¹⁶ and R¹⁷ independently of one another preferably represent hydrogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-halogenoalkyl, C₂-C₆-alkenyl or C₂-C₆-halogenoalkenyl; or C₃-C₆-cycloalkyl or C₃-C₆-cycloalkyl-C₁-C₄-alkyl, in each case optionally 15 substituted once or several times in an identical or different manner by C₁-C₄-alkyl, halogen or C₁-C₄-halogenoalkyl, or represent phenyl or phenyl-C₁-C₄-alkyl, in each case optionally substituted once to five times in an identical or different manner by halogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy, 20 C₁-C₄-halogenoalkylthio, di(C₁-C₄)alkylamino, cyano, nitro, C₁-C₆-alkoxycarbonyl or di(C₁-C₆)alkylaminocarbonyl, or represent -OR⁶ or -NR⁵R⁶,

or

R¹⁶ and R¹⁷ together represent -(CH₂)₅-, -(CH₂)₆- or -(CH₂)₂-O-(CH₂)₂.

25 X particularly preferably represents hydrogen, fluorine or chlorine.



Y particularly preferably represents fluorine or chlorine.

Z particularly preferably represents hydrogen, fluorine, chlorine, methyl, ethyl, methoxy, dimethylamino or diethylamino.

5 R¹ and R² independently of one another particularly preferably represent fluorine, chlorine, methyl, ethyl, methoxy, methylthio, trifluoromethyl, trifluoromethoxy or trifluoromethylthio.

m and n independently of one another particularly preferably represent 0, 1 or 2.

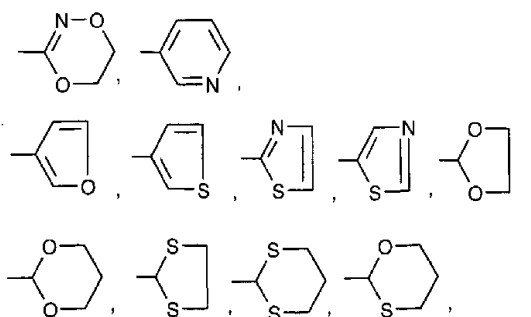
A particularly preferably represents the grouping (CH₂)_p-(CR³R⁴)_q-(CH₂)_r-R,

wherein

10 R³ and R⁴ independently of one another represent hydrogen, methyl, ethyl, n- or i-propyl, n-, i-, s- or t-butyl;

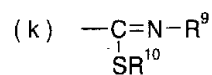
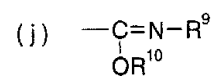
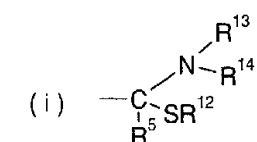
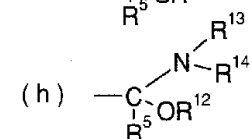
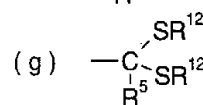
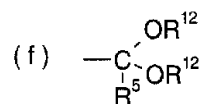
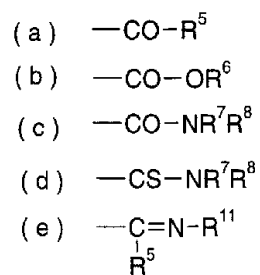
p, q and r independently of one another represent 0, 1, 2 or 3, at least one index being other than 0 and the sum of the indices being not greater than 5, in particular not greater than 3.

15 R particularly preferably represents cyano; or represents one of the following heterocyclic radicals



which are optionally substituted once to three times in an identical or different manner by fluorine, chlorine, bromine, methyl, ethyl, n- or i-propyl or trifluoromethyl;

or represents one of the following groupings:



wherein

R^5 represents hydrogen; methyl, ethyl, n- or i-propyl, the isomeric butyls, the



5 isomeric pentyls or the isomeric hexyls; or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms, C₃-C₆-alkenyl or C₃-C₆-halogenoalkenyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or cyclopropyl, cyclopentyl or cyclohexyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms,

10 or represents phenyl which is optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, methylcarbonylamino, ethylcarbonylamino, methylcarbonyl-methylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷,

15 R⁶ represents hydrogen, methyl, ethyl, n- or i-propyl or n-, i-, s- or t-butyl; or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or allyl which is optionally substituted once or several times by fluorine and/or chlorine; or cyclopropyl, cyclopentyl, cyclohexyl, cyclopropyl-C₁-C₂-alkyl, cyclopentyl-C₁-C₂-alkyl or cyclohexyl-C₁-C₂-alkyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms,

20 or represents phenyl-C₁-C₂-alkyl or naphthyl-C₁-C₂-alkyl, in each case optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷,

25 R⁷ and R⁸ independently of one another represent hydrogen, C₁-C₄-alkyl, methoxy, ethoxy or C₁-C₅-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or allyl which is optionally



substituted once or several times by fluorine and/or chlorine; or cyclopropyl, cyclopentyl, cyclohexyl, cyclopropylmethyl, cyclopentylmethyl or cyclohexylmethyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n-
5 or i-propyl or trifluoromethyl,
or represent phenyl or phenyl-C₁-C₂-alkyl, in each case optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, cyano,
10 nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷,
or represent -OR⁶ or -NR⁵R⁶,

R⁹ and R¹⁰ independently of one another represent methyl, ethyl or n- or i-propyl;

R¹¹ represents -OR⁶, -NR⁵R⁶ or -N(R⁵)-COOR⁶,

15 and

R¹², R¹³ and R¹⁴ independently of one another represent methyl, ethyl or n- or i-propyl.

R¹⁵ particularly preferably represents hydrogen, methyl, ethyl, or n- or i-propyl; or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as
20 F and Cl atoms; or cyclopropyl, cyclopentyl or cyclohexyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms,
or represents phenyl which is optionally substituted once or twice in an
25 identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, cyano, nitro, C₁-C₄-alkoxycarbonyl or



di(C₁-C₄)alkylaminocarbonyl.

- R¹⁶ and R¹⁷ independently of one another particularly preferably represent hydrogen, C₁-C₄-alkyl, methoxy, ethoxy or C₁-C₃-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or allyl which is optionally substituted once or several times by fluorine and/or chlorine; or cyclopropyl, cyclopentyl, cyclohexyl, cyclopropylmethyl, cyclopentylmethyl or cyclohexylmethyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or trifluoromethyl,
- or represent phenyl or phenyl-C₁-C₂-alkyl, in each case optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, cyano, nitro, C₁-C₄-alkoxycarbonyl or di(C₁-C₄)alkylaminocarbonyl,
- or represent -OR⁶ or -NR⁵R⁶.

In each case, if m = 2 and/or n = 2, the substituents R¹ and R² can be identical or different.

- The radical definitions and explanations listed above generally or listed in preferred ranges apply accordingly to the end products and to the starting substances and intermediate products. These definitions of radicals can be combined as desired with one another, that is to say also between the particularly preferred ranges.

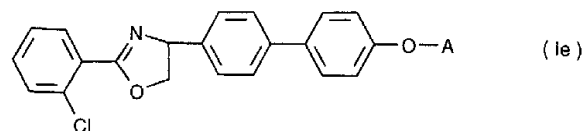
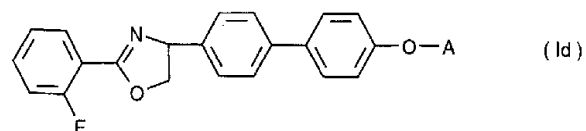
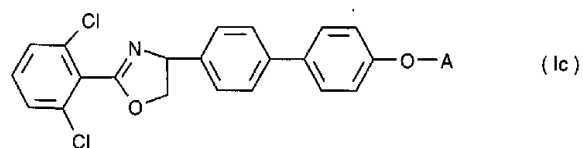
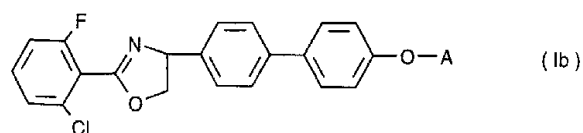
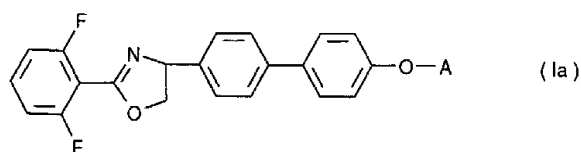
The compounds of the general formula (I) in which a combination of the meanings listed above as preferred (preferably) is present are preferred according to the invention.

- The compounds of the general formula (I) in which a combination of the meanings listed above as particularly preferred is present are particularly preferred according to the invention.



In the radical definitions listed above and below, hydrocarbon radicals, such as alkyl or alkenyl - including in combinations with heteroatoms, such as alkoxy or alkylthio - are, where possible, in each case straight-chain or branched.

Preferred compounds according to the invention are substances of the formulae (Ia) to (Ie):

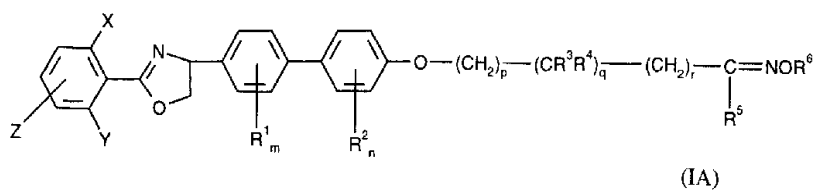


in which

A has the meanings given in the definition of the invention.

10 Preferred compounds according to the invention are also substances of the formula (IA):

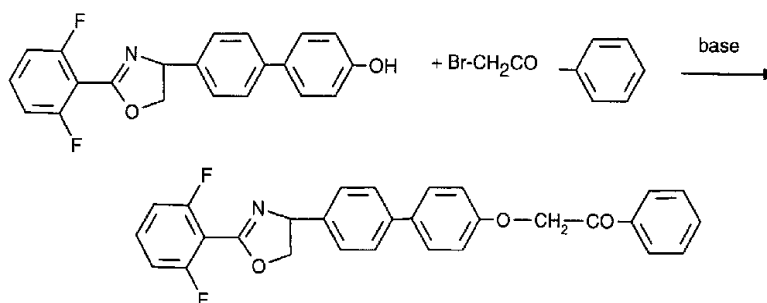




in which

- 5 $R^1, R^2, R^3, R^4, R^5, R^6, X, Y, Z, m, n, p, q$ and r have the meanings given in the definition of the invention, and among these, in turn, those where $m = 0$ and $Z = \text{hydrogen}$ are preferred, particularly preferably $m = 0, n = 0$ and $Z = \text{hydrogen}$.

- 10 If 2-(2,6-difluorophenyl)-4-(4'-hydroxybiphenyl-4-yl)-1,3-oxazoline and benzoylmethyl bromide are used as starting substances for carrying out the process according to the invention, the course of the reaction can be represented by the following equation:

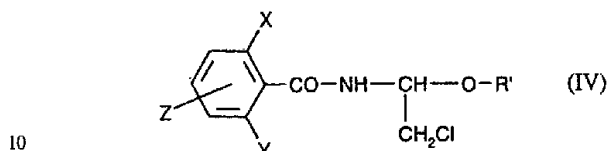


- 15 Formula (II) provides a general definition of the hydroxybiphenyloxazolines to be used as starting substances in the process according to the invention. In the formula (II), X, Y, Z, R^1, R^2, m and n preferably or particularly preferably have those meanings which have already been mentioned above as preferred or particularly preferred for these substituents and indices in connection with the description of the compounds of the formula (I) according to the invention.

- 20 The hydroxybiphenyloxazolines of the formula (II) are not yet known. However, some



are the subject matter of earlier applications filed by the Applicant which do not yet belong to the prior art (cf., for example, German Patent Application No. 44 44 108.1 of 12.12.1994 which is a priority document for Australian Patent Application No. 28445/95), and/or can be obtained by generally known processes (cf., for example, EP-A-0 432 661), by a procedure in which amide derivatives of the formula (IV)

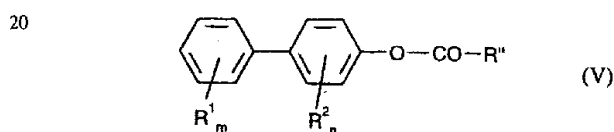


in which

X, Y and Z have the abovementioned meaning and

15 R' represents C₁-C₄-alkyl, preferably methyl or ethyl,

are reacted with biphenyl derivatives of the formula (V)



25 in which

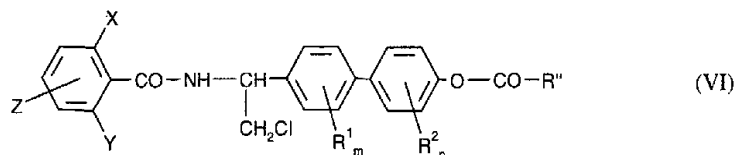
R¹, R², m and n have the abovementioned meanings and

R'' represents C₁-C₄-alkyl or C₁-C₄-alkoxy, preferably methyl, ethyl, methoxy or ethoxy,

30 in the presence of an acid catalyst such as, for example, sulfuric acid, acetic acid or aluminum chloride, and in the presence of a diluent, such as, for example, methylene chloride or acetonitrile, at temperatures between 0°C and 80°C;



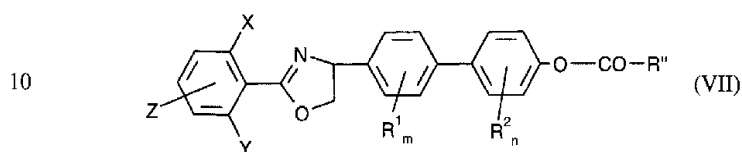
the compounds thus obtained, of the formula (VI)



in which

R^1 , R^2 , m , n , R'' , X , Y and Z have the abovementioned meaning,

- 5 are cyclized in the presence of a base, such as, for example, sodium hydroxide solution, if appropriate in the presence of a catalyst, such as, for example, ammonia compounds, and if appropriate in the presence of a diluent, such as, for example, dimethylformamide, at temperatures between 0°C and 100°C to give biphenyloxazolines of the formula (VII)



in which

R^1 , R^2 , m , n , R'' , X , Y and Z have the abovementioned meaning,

- 15 and these are hydrolyzed in the customary manner in the presence of a diluent, such as, for example, methanol, at room temperature, such as, for example, by means of aqueous ammonia solution, it also being possible for the cyclization and hydrolysis to be carried out in a one-pot reaction, if appropriate.

The amide derivatives of the formula (IV) are known (cf., for example, EP-A-0 594 179) and/or can be obtained by the methods mentioned therein.



Formula (III) provides a general definition of the compounds also to be used as starting substances in the process according to the invention. In the formula (III), A preferably or particularly preferably has those meanings which have already been mentioned as preferred or particularly preferred for this substituent in connection with the description
5 of the compounds of the formula (I) according to the invention.

M represents a customary leaving group, preferably halogen, in particular chlorine or bromine; alkylsulfonyloxy, in particular methylsulfonyloxy; or optionally substituted arylsulfonyloxy, in particular phenylsulfonyloxy, p-chlorophenylsulfonyloxy or tolylsulfonyloxy.

10 The compounds of the formula (III) are generally known compounds of organic chemistry.

Possible diluents for carrying out the process according to the invention are all the customary solvents. Solvents which can preferably be used are optionally halogenated, aromatic or aliphatic hydrocarbons, ketones, nitriles and amides. Examples which may
15 be mentioned are toluene, acetone, acetonitrile dimethylformamide and dimethylacetamide.

All the customary inorganic and organic bases are possible as the base for carrying out the process according to the invention. Examples which may be mentioned are tertiary amines, such as triethylamine, DBN (diazabicyclononene), DBU
20 (diazabicycloundecene), DABCO (diazabicyclooctane), alkali metal and alkaline earth metal hydroxides, such as, for example, sodium hydroxide, potassium hydroxide and calcium hydroxide, and alkali metal and alkaline earth metal carbonates, such as, for example, sodium carbonate or potassium carbonate.

If appropriate, the process according to the invention is carried out in the presence of
25 a phase transfer catalyst. Examples which may be mentioned are quaternary ammonium salts, such as tetraoctylammonium bromide or benzyltriethylammonium chloride, as well as tris(3,6-dioxaheptyl)amine (TDA).



The reaction temperatures can be varied within a substantial range in carrying out the process according to the invention. The reaction is in general carried out at temperatures between -20°C and 100°C, preferably between 0°C and 60°C.

5 In general, approximate equimolar amounts are used in carrying out the process according to the invention. However, it is also possible to use an excess of the compound of the formula (III).

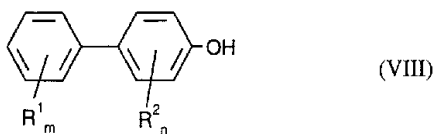
Working up and isolation are carried out in the generally customary manner.

10 In some cases, it proves advantageous to obtain compounds of the formula (I) in which R represents one of the groupings (e) to (k) by generally customary and known derivatizations of the corresponding keto derivatives, carboxylic acid derivatives and nitriles, i.e. compounds of the formula (I) in which R represents cyano or one of the groupings (a) to (d).

The derivatizations which lead to ketals, thioketals, oximes, oxime ethers or hydrazones are particularly preferred here.

15 The compounds of the formula (I) according to the invention can also be obtained by a process in which

A) in a first stage, compounds of the formula (VIII)



in which

20 R^1 , R^2 , m and n have the abovementioned meaning,



are reacted with a compound of the formula (III)

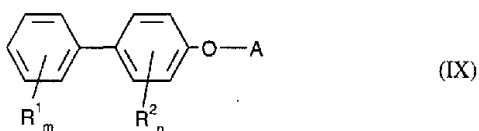


in which

M and A have the abovementioned meaning,

5 under the conditions of the process according to the invention;

B) in a second stage, the compounds thus obtained, of the formula (IX)

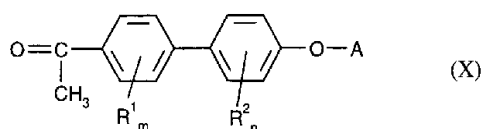


in which

A, R¹, R², m and n have the abovementioned meaning,

10 are reacted with acetyl chloride in the presence of a diluent, such as, for example, methylene chloride or dichloroethane, and in the presence of an acid or Lewis acid suitable for Friedel-Crafts reactions, such as, for example, tetrafluoroboric acid or aluminum chloride, at temperatures between -20°C and +50°C;

15 C) in a third stage, the compounds thus obtained, of the formula (X)

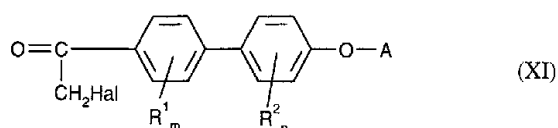


in which

A, R¹, R², m and n have the abovementioned meaning,

are chlorinated or brominated, if appropriate in the presence of a diluent, such as, for example, methylene chloride or carbon tetrachloride, at temperatures
5 between -10°C and 25°C;

D) in a fourth stage, the compounds thus obtained, of the formula (XI)



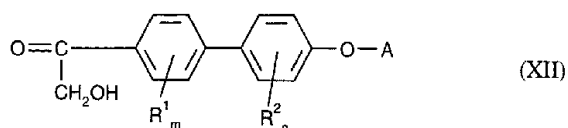
in which

A, R¹, R², m and n have the abovementioned meaning and

10 Hal represents chlorine or bromine,

are reacted with a salt of formic acid, such as, for example, sodium formate; in the presence of a diluent, if appropriate as a mixture with water, such as, for example, ethanol/water, and if appropriate in the presence of a phase transfer catalyst, such as, for example, quaternary ammonium salts, at temperatures
15 between 50°C and 150°C;

E) in a fifth stage, the compounds thus obtained, of the formula (XII)

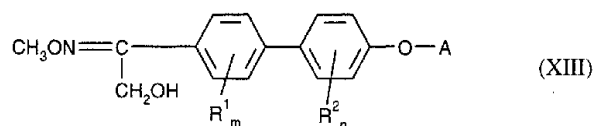


in which

A, R¹, R², m and n have the abovementioned meaning,

are reacted with O-methylhydroxylamine, if appropriate in the form of a salt, for example the hydrochloride, in the presence of a diluent, for example alcohols or ethers, and if appropriate in the presence of a base, for example sodium acetate, at temperatures between 0°C and 60°C;

F) in a sixth stage, the compounds thus obtained, of the formula (XIII)

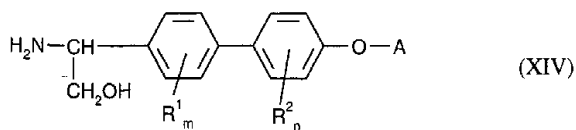


in which

A, R¹, R², m and n have the abovementioned meaning,

are reacted with a reducing agent, such as, for example, sodium boronate, in the presence of an acid, such as, for example, trifluoroacetic acid, and if appropriate in the presence of a diluent, such as, for example, tetrahydrofuran, at temperatures between 0°C and 120°C;

G) in a seventh stage, the compounds thus obtained, of the formula (XIV)

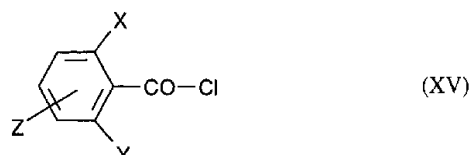


in which



A, R¹, R², m and n have the abovementioned meaning,

are reacted with benzoyl chlorides of the formula (XV)

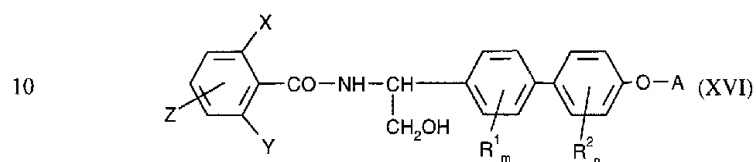


in which

5 X, Y and Z have the abovementioned meaning,

if appropriate in the presence of a base, such as, for example, triethylamine, and if appropriate in the presence of a diluent, such as, for example, tetrahydrofuran, at temperatures between 0°C and 100°C;

H) in an eighth stage, the compounds thus obtained, of the formula (XVI)



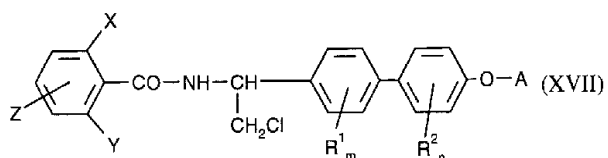
in which

A, R¹, R², m, n, X, Y and Z have the abovementioned meaning,

are reacted with a chlorinating agent, such as, for example, thionyl chloride, if appropriate in the presence of a diluent, such as, for example, toluene, at temperatures between 20°C and 100°C;

I) in a ninth stage, the compounds thus obtained, of the formula (XVII)





in which

A, R¹, R², m, n, X, Y and Z have the abovementioned meaning,

5 are cyclized in the presence of a base, such as, for example, sodium hydroxide solution; if appropriate in the presence of a phase transfer catalyst, such as, for example, ammonium compounds, and if appropriate in the presence of a diluent, such as, for example, dimethylformamide, at temperatures between 0°C and 100°C, to give the compounds of the formula (I) according to the invention.

10 The compounds of the formula (XVII) can also be obtained directly by reacting amide derivatives of the formula (IV) with compounds of the formula (IX) in the presence of an acid catalyst, such as, for example, hydrogen fluoride, boron trifluoride or aluminum chloride, and in the presence of a diluent, such as, for example, methylene chloride or acetonitrile, at temperatures between 0°C and 80°C.

15 The starting substances of the formula (VIII) are known and/or can be prepared in a simple manner by known methods.

The compounds of the formula (VIII) are obtained, for example, by sulfonating optionally substituted biphenyls and then reacting the products with alkali metal hydroxides to give the hydroxybiphenyls, or diazotizing aminobiphenyls and boiling the products. (cf., for example, Houben-Weyl, Volume VI/1c (1976), page 216 and 251).

20 The benzoyl chlorides of the formula (XV) are generally known compounds of organic chemistry.

The compounds of the formulae (X), (XI), (XII), (XIII), (XIV), (XVI) and (XVII)



which occur as intermediate products are not yet known and the invention also relates to these.

5 The active compounds are suitable for controlling animal pests, in particular insects, arachnids and nematodes which are encountered in agriculture, in forestry, in the preservation of stored products and of materials, and in the hygiene sector, while being well tolerated by crops and of favorable toxicity to warm-blooded animals. They can preferably be used as crop protection products. They are active against normally sensitive and resistant species and against all or some stages of development. The abovementioned pests include:

10 From the order of the Isopoda, for example, *Oniscus asellus*, *Armadillidium vulgare* and *Porcellio scaber*.

From the order of the Diplopoda, for example, *Blaniulus guttulatus*.

From the order of the Chilopoda, for example, *Geophilus carpophagus* and *Scutigera spec.*

15 From the order of the Symphyla, for example, *Scutigera immaculata*.

From the order of the Thysanura, for example, *Lepisma saccharina*.

From the order of the Collembola, for example, *Onychiurus armatus*.

20 From the order of the Orthoptera, for example, *Blatta orientalis*, *Periplaneta americana*, *Leucophaea maderae*, *Blattella germanica*, *Acheta domesticus*, *Gryllotalpa* spp., *Locusta migratoria migratorioides*, *Melanoplus differentialis* and *Schistocerca gregaria*.

From the order of the Dermaptera, for example, *Forficula auricularia*.

From the order of the Isoptera, for example, *Reticulitermes* spp.



From the order of the Anoplura, for example, *Pediculus humanus corporis*, *Haematopinus* spp. and *Linognathus* spp.

From the order of the Mallophaga, for example, *Trichodectes* spp. and *Damalinea* spp.

5 From the order of the Thysanoptera, for example, *Hercinothrips femoralis* and *Thrips tabaci*.

From the order of the Heteroptera, for example, *Eurygaster* spp., *Dysdercus intermedius*, *Piesma quadrata*, *Cimex lectularius*, *Rhodnius prolixus* and *Triatoma* spp.

10 From the order of the Homoptera, for example, *Aleurodes brassicae*, *Bemisia tabaci*, *Trialeurodes vaporariorum*, *Aphis gossypii*, *Brevicoryne brassicae*, *Cryptomyzus ribis*, *Aphis fabae*, *Aphis pomi*, *Eriosoma lanigerum*, *Hyalopterus arundinis*, *Phylloxera vastatrix*, *Pemphigus* spp., *Macrosiphum avenae*, *Myzus* spp., *Phorodon humuli*, *Rhopalosiphum padi*, *Empoasca* spp., *Euscelis bilobatus*, *Nephotettix cincticeps*, *Lecanium corni*, *Saissetia oleae*, *Laodelphax striatellus*, *Nilaparvata lugens*, *Aonidiella aurantii*, *Aspidiotus hederae*, *Pseudococcus* spp. and *Psylla* spp.

15 From the order of the Lepidoptera, for example, *Pectinophora gossypiella*, *Bupalus piniarius*, *Cheimatobia brumata*, *Lithocolletis blancardella*, *Hyponomeuta padella*, *Plutella maculipennis*, *Malacosoma neustria*, *Euproctis chrysorrhoea*, *Lymantria* spp., *Bucculatrix thurberiella*, *Phyllocnistis citrella*, *Agrotis* spp., *Euxoa* spp., *Feltia* spp., *Earias insulana*, *Heliothis* spp., *Spodoptera exigua*, *Mamestra brassicae*, *Panolis*
20 *flammea*, *Spodoptera litura*, *Spodoptera* spp., *Trichoplusia ni*, *Carpocapsa pomonella*, *Pieris* spp., *Chilo* spp., *Pyrausta nubilalis*, *Ephestia kuehniella*, *Galleria mellonella*, *Tineola bisselliella*, *Tinea pellionella*, *Hofmannophila pseudospretella*, *Cacoecia podana*, *Capua reticulana*, *Choristoneura fumiferana*, *Clysia ambiguella*, *Homona magnanima* and *Tortrix viridana*.

25 From the order of the Coleoptera, for example, *Anobium punctatum*, *Rhizopertha dominica*, *Bruchidius obtectus*, *Acanthoscelides obtectus*, *Hylotrupes bajulus*, *Agelastica*



alni, *Leptinotarsa decemlineata*, *Phaedon cochleariae*, *Diabrotica* spp., *Psylliodes chrysocephala*, *Epilachna varivestis*, *Atomaria* spp., *Oryzaephilus surinamensis*, *Anthonomus* spp., *Sitophilus* spp., *Otiorrhynchus sulcatus*, *Cosmopolites sordidus*, *Ceuthorrhynchus assimilis*, *Hypera postica*, *Dermestes* spp., *Trogoderma* spp.,
5 *Anthrenus* spp., *Attagenus* spp., *Lyctus* spp., *Meligethes aeneus*, *Ptinus* spp., *Niptus hololeucus*, *Gibbium psylloides*, *Tribolium* spp., *Tenebrio molitor*, *Agriotes* spp., *Conoderus* spp., *Melolontha melolontha*, *Amphimallon solstitialis* and *Costelytra zealandica*.

From the order of the Hymenoptera, for example, *Diprion* spp., *Hoplocampa* spp.,
10 *Lasius* spp., *Monomorium pharaonis* and *Vespa* spp.

From the order of the Diptera, for example, *Aedes* spp., *Anopheles* spp., *Culex* spp., *Drosophila melanogaster*, *Musca* spp., *Fannia* spp., *Calliphora erythrocephala*, *Lucilia* spp., *Chrysomyia* spp., *Cuterebra* spp., *Gastrophilus* spp., *Hyppobosca* spp., *Stomoxys* spp., *Oestrus* spp., *Hypoderma* spp., *Tabanus* spp., *Tannia* spp., *Bibio hortulanus*,
15 *Oscinella frit*, *Phorbia* spp., *Pegomyia hyoscyami*, *Ceratitis capitata*, *Dacus oleae* and *Tipula paludosa*.

From the order of the Siphonaptera, for example, *Xenopsylla cheopis* and *Ceratophyllus* spp.

From the order of the Arachnida, for example, *Scorpio maurus*² and *Latrodectus mactans*.
20

From the order of the Acarina, for example, *Acarus siro*, *Argas* spp., *Ornithodoros* spp., *Dermanyssus gallinae*, *Eriophyes ribis*, *Phyllocoptruta oleivora*, *Boophilus* spp., *Rhipicephalus* spp., *Amblyomma* spp., *Hyalomma* spp., *Ixodes* spp., *Psoroptes* spp., *Chorioptes* spp., *Sarcoptes* spp., *Tarsonemus* spp., *Bryobia praetiosa*, *Panonychus* spp.,
25 and *Tetranychus* spp..

The phytoparasitic nematodes include *Pratylenchus* spp., *Radopholus similis*,



Ditylenchus dipsaci, Tylenchulus semipenetrans, Heterodera spp., Globodera spp., Meloidogyne spp., Aphelenchoides spp., Longidorus spp., Xiphinema spp. and Trichodorus spp..

5 The compounds of the formula (I) according to the invention are distinguished in particular by a high insecticidal and acaricidal activity.

10 They can be used with particularly good success for controlling phytopathogenic insects, such as, for example, against the mustard beetle larvae (Phaedon cochlaeriae), caterpillars of the diamond-back moth (Plutella maculipennis), the peach aphid (Myzus persicae) and caterpillars of the owlet moth (Spodoptera frugiperda), or for controlling phytopathogenic mites, such as, for example, against the common spider mite (Tetranychus urticae).

15 The active compounds can be converted to the customary formulations, such as solutions, emulsions, wettable powders, suspensions, powders, dusts, pastes, soluble powders, granules, suspension-emulsion concentrates, natural and synthetic materials impregnated with active compound and very fine capsules in polymeric substances.

These formulations are produced in a known manner, for example by mixing the active compounds with extenders, that is, liquid solvents and/or solid carriers, optionally with the use of surface-active agents, that is emulsifying agents and/or dispersing agents, and/or foam-forming agents.

20 In the case of the use of water as an extender, organic solvents can, for example, also be used as auxiliary solvents. As liquid solvents, there are suitable in the main: aromatics, such as xylene, toluene or alkylnaphthalenes, chlorinated aromatics or chlorinated aliphatic hydrocarbons, such as chlorobenzenes, chloroethylenes or methylene chloride, aliphatic hydrocarbons, such as cyclohexane or paraffins, for
25 example mineral oil fractions, mineral and vegetable oils, alcohols, such as butanol or glycol as well as their ethers and esters, ketones, such as acetone, methyl ethyl ketone, methyl isobutyl ketone or cyclohexanone, strongly polar solvents, such as



dimethylformamide and dimethyl sulfoxide, as well as water.

As solid carriers there are suitable:

for example ammonium salts and ground natural minerals, such as kaolins, clays, talc, chalk, quartz, attapulgite, montmorillonite or diatomaceous earth, and ground synthetic

- 5 minerals, such as highly disperse silica, alumina and silicates; as solid carriers for granules there are suitable: for example crushed and fractionated natural rocks such as calcite, marble, pumice, sepiolite and dolomite, as well as synthetic granules of inorganic and organic meals, and granules of organic material such as sawdust, coconut shells, maize cobs and tobacco stalks; as emulsifying and/or foam-forming agents there
10 are suitable: for example nonionic and anionic emulsifiers, such as polyoxyethylene fatty acid esters, polyoxyethylene fatty alcohol ethers, for example alkylaryl polyglycol ethers, alkylsulfonates, alkyl sulfates, arylsulfonates as well as albumen hydrolysis products; as dispersing agents there are suitable: for example lignin-sulfite waste liquors and methylcellulose.

- 15 Adhesives such as carboxymethylcellulose and natural and synthetic polymers in the form of powders, granules or latices, such as gum arabic, polyvinyl alcohol and polyvinyl acetate, as well as natural phospholipids, such as cephalins and lecithins, and synthetic phospholipids, can be used in the formulations. Other additives can be mineral and vegetable oils.

- 20 It is possible to use colorants such as inorganic pigments, for example iron oxide, titanium oxide and Prussian Blue, and organic dyestuffs, such as alizarin dyestuffs, azo dyestuffs and metal phthalocyanine dyestuffs, and trace nutrients such as salts of iron, manganese, boron, copper, cobalt, molybdenum and zinc.

- The formulations in general contain between 0.1 and 95 percent by weight of active
25 compound, preferably between 0.5 and 90%.

The active compound according to the invention can be present in its commercially available formulations and in the use forms, prepared from these formulations, as a



- mixture with other active compounds, such as insecticides, attractants, sterilizing agents, bactericides, acaricides, nematocides, fungicides, growth-regulating substances or herbicides. The insecticides include, for example, phosphates, carbamates, carboxylates, chlorinated hydrocarbons, phenylureas and substances produced by
- 5 microorganisms, inter alia.

Particularly favorable mixing partners are, for example, the following:

Fungicides:

- 2-aminobutane; 2-anilino-4-methyl-6-cyclopropyl-pyrimidine; 2',6'-dibromo-2-methyl-4'-trifluoromethoxy-4'-trifluoro-methyl-1,3-thiazole-5-carboxanilide; 2,6-dichloro-N-
- 10 (4-trifluoromethylbenzyl)-benzamide; (E)-2-methoxyimino-N-methyl-2-(2-phenoxyphenyl)-acetamide; 8-hydroxyquinoline sulfate; methyl (E)-2-{2-[6-(2-cyano-phenoxy)-pyrimidin-4-yl]oxy}-phenyl}-3-methoxyacrylate; methyl (E)-methoximino[alpha-(o-tolyloxy)-o-tolyl]acetate; 2-phenylphenol (OPP), aldimorph, ampropylfos, anilazine, azaconazole,
- 15 benalaxyl, benodanil, benomyl, binapacryl, biphenyl, bitertanol, blasticidin-S, bromuconazole, bupirimate, buthiobate,
- calcium polysulfide, captafol, captan, carbendazime, carboxin, chinomethionat (quinomethionate), chloroneb, chloropicrin, chlorothalonil, chlozolate, cufraneb, cymoxanil, cyproconazole, cyprofuram,
- 20 dichlorophen, diclobutrazole, diclofluanid, diclomezin, dicloran, diethofencarb, difenoconazole, dimethirimol, dimethomorph, diniconazole, dinocap, diphenylamine, dipyrithione, ditalimfos, dithianon, dodine, drazoxolon,
- edifenphos, epoxyconazole, ethirimol, etridiazole,
- fenarimol, fenbuconazole, fenfuram, fenitropan, fenpiclonil, fenpropidin,
- 25 fenpropimorph, fentin acetate, fentin hydroxide, ferbam, ferimzone, fluazinam,



fludioxonil, fluoromide, fluquinconazole, flusilazole, flusulfamide, flutolanil, flutriafol, folpet, fosetyl-aluminum, fthalide, fuberidazole, furalaxyl, furmecyclox,

guazatine,

hexachlorobenzene, hexaconazole, hymexazole,

5 imazalil, imibenconazole, iminoctadine, iprobenfos (IBP), iprodione, isoprothiolane,

kasugamycin, copper formulations, such as: copper hydroxide, copper naphthenate, copper oxychloride, copper sulfate, copper oxide, oxine-copper and Bordeaux mixture,

mancopper, mancozeb, maneb, mepanipyrim, mepronil, metalaxyl, metconazole, methasulfocarb, methfuroxam, metiram, metsulfovax, myclobutanil,

10 nickel dimethyldithiocarbamate, nitrothal-isopropyl, nuarimol,

ofurace, oxadixyl, oxamocarb, oxycarboxin,

pefurazate, penconazole, pencycuron, phosdiphen, phthalide, pimaricin, piperalin, polycarbamate, polyoxin, probenazole, prochloraz, procymidone, propamocarb, propiconazole, propineb, pyrazophos, pyrifenox, pyrimethanil, pyroquilon,

15 quintozone (PCNB),

sulfur and sulfur formulations,

tebuconazole, tecloftalam, tecnazene, tetraconazole, thiabendazole, thicyofen, thiophanate-methyl, thiram, tolclophos-methyl, tolylfluanid, triadimefon, triadimenol, triazoxide, trichlamide, tricyclazole, tridemorph, triflumizole, triforine, triticonazole,

20 validamycin A, vinclozolin,



zineb, ziram.

Bactericides:

- bronopol, dichlorophen, nitrapyrin, nickel dimethyldithiocarbamate, kasugamycin, oethilinone, furanecarboxylic acid, oxytetracycline, probenazole, streptomycin,
5 tecloftalam, copper sulfate and other copper formulations.

Insecticides/Acaricides/Nematicides:

- abamectin, AC 303 630, acephate, acrinathrin, alanycarb, aldicarb, alphamethrin, amitraz, avermectin, AZ 60541, azadirachtin, azinphos A, azinphos M, azocyclotin, Bacillus thuringiensis, bendiocarb, benfuracarb, bensultap, betacyfluthrin, bifenthrin,
10 BPMC, brofenprox, bromophos A, bufencarb, buprofezin, butocarboxin, butylpyridaben, cadusafos, carbaryl, carbofuran, carbophenothion, carbosulfan, cartap, CGA 157 419, CGA 184 699, chloethocarb, chlorethoxyfos, chlorfenvinphos, chlorfluazuron, chlormephos, chlorpyrifos, chlorpyrifos M, cis-resmethrin, clocythrion, clofentezine, cyanophos, cycloprothrin, cyfluthrin, cyhalothrin, cyhexatin, cypermethrin, cyromazine,
15 deltamethrin, demeton M, demeton S, demeton-S-methyl, diafenthiuron, diazinon, dichlofenthion, dichlorvos, dicliphos, dicrotophos, diethion, diflubenzuron, dimethoate, dimethylvinphos, dioxathion, disulfoton, edifenphos, emamectin, esfenvalerate, ethiofencarb, ethion, ethofenprox, ethoprophos, etrimphos,
20 fenamiphos, fenazaquin, fenbutatin oxide, fenitrothion, fenobucarb, fenothiocarb, fenoxycarb, fenpropathrin, fenpyrad, fenpyroximate, fenthion, fenvalerate, fipronil, fluazinam, flucycloxuron, flucythrinate, flufenoxuron, flufenprox, fluvalinate, fonophos, formothion, fosthiazate, fubfenprox, furathiocarb, HCH, heptenophos, hexaflumuron, hexythiazox,



imidacloprid, iprobenfos, isazophos, isofenphos, isoprocab, isoxathion, ivermectin,
lambda-cyhalothrin, lufenuron,

malathion, mecarbam, mervinphos, mesulfenphos, metaldehyde, methacrifos,
methamidophos, methidathion, methiocarb, methomyl, metolcarb, milbemectin,
5 monocrotophos, moxidectin,

naled, NC 184, NI 25, nitenpyram,

omethoate, oxamyl, oxydemethon M, oxydeprofos,

parathion A, parathion M, permethrin, phenthoate, phorate, phosalone, phosmet,
phosphamidon, phoxim, pirimicarb, pirimiphos M, pirimiphos A, profenofos,
10 promecarb, propaphos, propoxur, prothiofos, prothoate, pymetrozin, pyrachlophos,
pyradaphenthion, pyresmethrin, pyrethrum, pyridaben, pyrimidifen, pyriproxifen,
quinalphos,

RH 5992,

salithion, sebufos, silafluofen, sulfotep, sulprofos,

15 tebufenozide, tebufenpyrad, tebupirimiphos, teflubenzuron, tefluthrin, temephos, terbam,
terbufos, tetrachlorvinphos, thiafenox, thiodicarb, thiofanox, thiomethon, thionazin,
thuringiensin, tralomethrin, triarathen, triazophos, triazuron, trichlorfon, triflumuron,
trimethacarb,

vamidothion, XMC, xylylcarb, YI 5301/5302, zetamethrin.

20 A mixture with other known active compounds, such as herbicides, or with fertilizers
and growth regulators is also possible.



The active compounds according to the invention can furthermore be present in their commercially available formulations and in the use forms, prepared from these formulations, as a mixture with synergistic agents. Synergistic agents are compounds which increase the action of the active compounds, without it being necessary for the synergistic agent added to be active itself.

The active compound content of the use forms prepared from the commercially available formulations can vary within wide limits. The active compound concentration of the use forms can be from 0.0000001 to 95% by weight of active compound, preferably between 0.0001 and 1% by weight.

10 The compounds are employed in a customary manner appropriate for the use forms.

When used against hygiene pests and pests of stored products, the active compound is distinguished by an excellent residual action on wood and clay as well as a good stability to alkali on limed substrates.

15 The active compounds according to the invention act not only against plant and hygiene pests and pests of stored products, but also in the veterinary medicine sector against animal parasites (ectoparasites), such as hard ticks, soft ticks, mange mites, harvest mites, flies (biting and licking), parasitic fly larvae, lice, biting lice, bird lice and fleas. These parasites include:

20 From the order of the Anoplurida, for example, *Haematopinus* spp., *Linognathus* spp., *Pediculus* spp., *Phthirus* spp. and *Solenopotes* spp.

From the order of the Mallophagida and the suborders Amblycerina and Ischnocerina, for example, *Trimenopon* spp., *Menopon* spp., *Trinoton* spp., *Bovicola* spp., *Werneckiella* spp., *Lepikentron* spp., *Damalina* spp., *Trichodectes* spp. and *Felicola* spp..

25 From the order Diptera and the suborders Nematocerina and Brachycerina, for example,



- Aedes spp., Anopheles spp., Culex spp., Simulium spp., Eusimulium spp., Phlebotomus spp., Lutzomyia spp., Culicoides spp., Chrysops spp., Hybomitra spp., Atylotus spp., Tabanus spp., Haematopota spp., Philipomyia spp., Braula spp., Musca spp., Hydrotæa spp., Stomoxys spp., Haematobia spp., Morellia spp., Fannia spp., Glossina spp.,
- 5 Calliphora spp., Lucilia spp., Chrysomyia spp., Wohlfahrtia spp., Sarcophaga spp., Oestrus spp., Hypoderma spp., Gasterophilus spp., Hippobosca spp., Lipoptena spp. and Melophagus spp.

From the order of the Siphonaptera, for example, Pulex spp., Ctenocephalides spp., Xenopsylla spp. and Ceratophyllus spp.

- 10 From the order of the Heteroptera, for example, Cimex spp., Triatoma spp., Rhodnius spp. and Panstrongylus spp.

From the order of the Blattaria, for example, Blatta orientalis, Periplaneta americana, Blattella germanica and Supella spp.

- 15 From the subclass of the Acaria (Acarida) and the orders of the Meta- and Mesostigmata, for example, Argas spp., Ornithodoros spp., Otobius spp., Ixodes spp., Amblyomma spp., Boophilus spp., Dermacentor spp., Haemophysalis spp., Hyalomma spp., Rhipicephalus spp., Dermanyssus spp., Raillietia spp., Pneumonyssus spp., Sternostoma spp. and Varroa spp.

- 20 From the order of the Actiniedida (Prostigmata) and Acaridida (Astigmata), for example, Acarapis spp., Cheyletiella spp., Ornithocheyletia spp., Myobia spp., Psorergates spp., Demodex spp., Trombicula spp., Listrophorus spp., Acarus spp., Tyrophagus spp., Caloglyphus spp., Hypodectes spp., Pterolichus spp., Psoroptes spp., Choriopetes spp., Otodectes spp., Sarcoptes spp., Notoedres spp., Knemidocoptes spp., Cytodites spp. and Laminosioptes spp.

- 25 For example, they show a good development-inhibiting action against fly larvae of Lucilla cuprina.



The active compounds of the formula (I) according to the invention are also suitable for controlling arthropods which infest agricultural productive livestock, such as, for example, cattle, sheep, goats, horses, pigs, donkeys, camels, buffalo, rabbits, chickens, turkeys, ducks, geese and bees, other pets, such as, for example, dogs, cats, cage birds
5 and aquarium fish, and so-called experimental animals, such as, for example, hamsters, guinea pigs, rats and mice. By controlling these arthropods, cases of death and reductions in productivity (for meat, milk, wool, hides, eggs, honey and the like) should be diminished, so that more economical and easier animal husbandry is possible by use of the active compounds according to the invention.

10 The active compounds according to the invention are used in the veterinary sector in a known manner by enteral administration in the form of, for example, tablets, capsules, potions, drenches, granules, pastes, boli, the feed-through process and suppositories, by parenteral administration, such as, for example, by injections (intramuscular, subcutaneous, intravenous, intraperitoneal and the like), implants, by
15 nasal administration, by dermal use in the form, for example, of dipping or bathing, spraying, pouring on and spotting on, washing and powdering, and also with the aid of molded articles containing the active compound, such as collars, ear tags, tail tags, limb bands, halters, marking devices and the like.

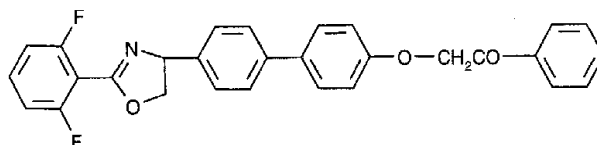
When used for livestock, poultry, pets and the like, the active compounds of the
20 formula (I) can be used as formulations (for example powders, emulsions, free-flowing compositions) which comprise the active compounds in an amount of 1 to 80 % by weight, directly or after 100- to 10,000-fold dilution, or they can be used as a chemical bath.

The preparation and the use of the active compounds according to the invention can be
25 seen from the following examples.



Preparation examples:

Example 1

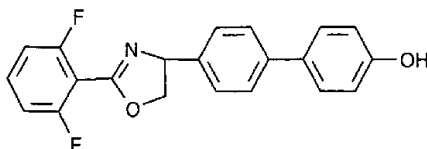


3.51 g (0.01 mol) of 2-(2,6-difluorophenyl)-4-(4'-hydroxy-biphenyl-4-yl)-1,3-oxazoline,
5 2.0 g (0.01 mol) of benzoylmethyl bromide and 1.38 g (0.01 mol) of potassium carbonate are suspended in 50 ml of acetonitrile. After addition of three drops of tris(3,6-dioxahexyl)amine (TDA), the reaction mixture is stirred at 80°C for 18 hours and then concentrated. The residue is taken up in 100 ml of methylene chloride, the mixture is washed with water, dried over sodium sulfate and concentrated and the
10 residue is purified by column chromatography with methylene chloride as the mobile phase.

1.7 g (36.2 % of theory) of 2-(2,6-difluorophenyl)-4-(4'-benzoylmethoxy-biphenyl-4-yl)-1,3-oxazoline of melting point m.p.: 78-82°C are obtained.

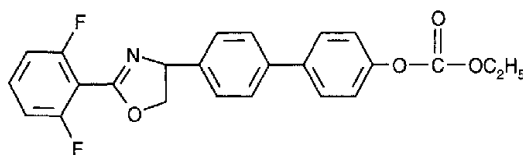


Preparation of the starting substance

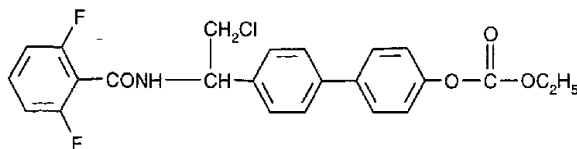


- 5 10.9 g (0.026 mol) of 2-(2,6-difluorophenyl)-4-(4'-ethoxycarbonyloxybiphenyl-4-yl)-1,3-oxazoline are suspended in 50 ml of methanol, and 35.4 ml (0.52 mol) of 25 % strength aqueous ammonia solution are then added dropwise at room temperature. After 28 hours at room temperature, the precipitate is filtered off with suction and washed with a little methanol.

Yield 7.6 g (97.4 % of theory), m.p.: 180 to 182°C.



- 10 4 ml (0.045 mol) of 50 % strength sodium hydroxide solution are added dropwise to 19.8 g (0.043 mol) of 2-(2,6-difluorobenzoylamido)-2-(4'-ethoxycarbonyloxybiphenyl-4-yl)-1-chloroethane in 120 ml of dimethyl formamide. After 2 hours at room temperature, the mixture is stirred into 500 ml of ice-water and the crystals which have precipitated out are filtered off with suction.
- 15 Yield: 15.1 g (83 % of theory), m.p.: 98 to 100°C.

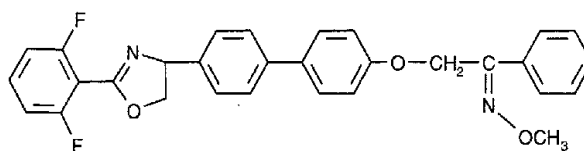


130.4 g (0.98 mol) of aluminum chloride are introduced in portions into a mixture of



- 53 g (0.213 mol) of 2-(2,6-difluorobenzoylamido)-2-methoxy-1-chloroethane, 48.4 g (0.2 mol) of 4-ethoxycarbonyloxybiphenyl and 12 ml of glacial acetic acid in 200 ml of methylene chloride at 5°C in the course of 30 minutes. During this operation, the batch changes color via blue to red-violet. The reaction mixture is stirred at 5°C for one hour and at 10°C for 1 hour and poured cautiously onto ice, the suspension is decanted cautiously from the water and concentrated on a rotary evaporator, 50 ml of acetonitrile are added to the residue and the crystals which have precipitated out are filtered off with suction.
- Yield 42.8 g (47 % of theory), m.p.: 209°C.

10 **Example 2**

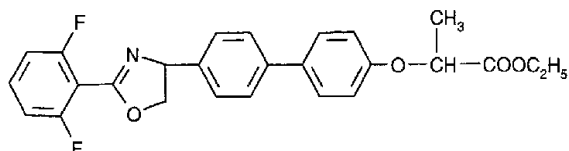


- 0.167 g (0.002 mol) of O-methylhydroxylamine hydrochloride and 0.164 g (0.002 mol) of sodium acetate are added to 0.96 g (0.002 mol) of 2-(2,6-difluorophenyl)-4-(4'-benzoylmethoxy-biphenyl-4-yl)-1,3-oxazoline (Example 1) in 20 ml of methanol.
- 15 The reaction mixture is stirred at 65°C for 6 hours and then concentrated. The residue is taken up in 20 ml of methylene chloride and the mixture is washed with water, dried over sodium sulfate and concentrated. The oily residue is purified by column chromatography (mobile phase: toluene/ether: 8/2).
- 0.7 g (70 % of theory) of 2-(2,6-difluorophenyl)-4-[4'-(2-methoximino-2-phenyl)-ethoxy-biphenyl-4-yl]-1,3-oxazoline with a log p = 5.01 is obtained.
- 20

[log p = logarithm of the partition coefficient p of the substance between the solvents octanol and water, determined experimentally from reversed phase HPLC]



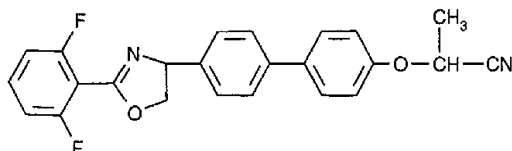
Example 3



3.51 g (0.01 mol) of 2-(2,6-difluorophenyl)-4-(4'-hydroxybiphenyl-4-yl)-1,3-oxazoline and 1.38 g (0.01 mol) of potassium carbonate are suspended in 50 ml of acetonitrile and, after addition of three drops of tris(3,6-dioxahexyl)amine (TDA), 1.8 g (0.01 mol) of ethyl 2-bromopropionate are added. The reaction mixture is stirred at 65°C for 18 hours and then concentrated. The residue is taken up in 100 ml of methylene chloride, and the mixture is washed with water, dried over sodium sulfate and concentrated. The oily residue is purified by column chromatography with methylene chloride as the mobile phase.

2.5 g (55.4 % of theory) of 2-(2,6-difluorophenyl)-4-[4'-(1-ethoxycarbonyl-ethyl-oxy)-biphenyl-4-yl]-1,3-oxazoline with a log p = 4.12 are obtained.

Example 4

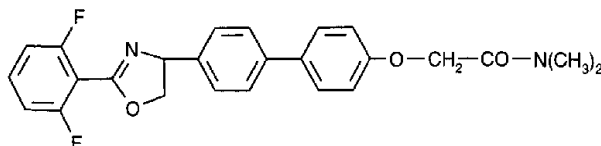


3.51 g (0.01 mol) of 2-(2,6-difluorophenyl)-4-(4'-hydroxybiphenyl-4-yl)-1,3-oxazoline and 1.38 g (0.01 mol) of potassium carbonate are suspended in 50 ml of acetonitrile and, after addition of three drops of tris(3,6-dioxahexyl)amine (TDA), 0.9 g (0.01 mol) of 2-chloropropionitrile is added. The reaction mixture is stirred at 80°C for 20 hours and then concentrated. The residue is taken up in 100 ml of methylene chloride and the mixture is washed with water, dried over sodium sulfate and concentrated. The oily residue is purified by column chromatography with methylene chloride as the mobile phase.



2.6 g (64.4 % of theory) of 2-(2,6-difluorophenyl)-4-[4'-(2-propionitriloxy)-biphenyl-4-yl]-1,3-oxazoline of melting point 100 - 120°C are obtained.

Example 5



5 3.51 g (0.01 mol) of 2-(2,6-difluorophenyl)-4-(4'-hydroxybiphenyl-4-yl)-1,3-oxazoline and 1.38 g (0.01 mol) of potassium carbonate are suspended in 50 ml of acetonitrile and, after addition of three drops of tris(3,6-dioxaheptyl)amine (TDA), 1.2 g (0.01 mol) of 2-chloro-N,N-dimethyl-acetamide are added. The reaction mixture is stirred at 80°C for 20 hours and then concentrated. The residue is taken up in 100 ml of methylene chloride and the mixture is washed with water, dried over sodium sulfate and concentrated. The residue is stirred with diisopropyl ether and the mixture is concentrated.

10 2.4 g (55 % of theory) of 2-(2,6-difluorophenyl)-4-(4'-dimethylaminocarbonylmethoxy-biphenyl-4-yl)-1,3-oxazoline of melting point 15 78-80°C are obtained.

The compounds of the formula (I) shown in the following table are obtained analogously to Examples 1 to 5 and in accordance with the general preparation instructions:

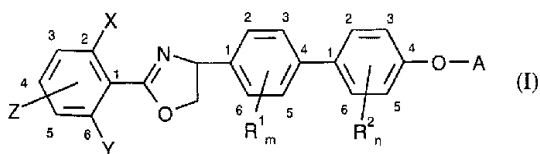


Table 1:

Ex. No.	X	Y	Z	R _m ¹	R _n ²	A	m.p. (°C ¹) or log p
6	F	F	H	H	H	-CH ₂ -COOCH ₃	125-28 ¹⁾
7	F	F	H	H	H	-CH ₂ -COCH ₃	120-25 ¹⁾
8	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}-\text{COCH}_3 \end{array}$	120-22 ¹⁾
9	F	F	H	H	H	-CH ₂ -COOC ₂ H ₅	115-18 ¹⁾
10	F	F	H	H	H	-CH ₂ -COOC ₄ H ₉ -t	4.43
11	F	F	H	H	H	$\begin{array}{c} \text{C}_2\text{H}_5 \\ \\ -\text{CH}-\text{COOC}_2\text{H}_5 \end{array}$	4.53
12	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}-\text{COOCH}_3 \end{array}$	3.80
13	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_3 \end{array}$	70 ¹⁾
14	F	F	H	H	H	-CH ₂ -CH ₂ -CH ₂ -CN	95-98 ¹⁾
15	F	F	H	H	H	-CH ₂ -CO-N(C ₂ H ₅) ₂	3.35
16	F	F	H	H	H	-CH ₂ -CH ₂ -CH(OC ₂ H ₅) ₂	4.74
17	F	F	H	H	H	$\begin{array}{c} \text{C}_3\text{H}_7\text{-i} \\ \\ -\text{CH}-\text{CN} \end{array}$	4.45
18	F	F	H	H	H	$\begin{array}{c} \text{C}_3\text{H}_7\text{-i} \\ \\ -\text{CH}-\text{COOCH}_3 \end{array}$	4.56
19	F	F	H	H	H	$\begin{array}{c} \text{C}_2\text{H}_5 \\ \\ -\text{CH}-\text{COOCH}_3 \end{array}$	4.18



Table 1 (continued)

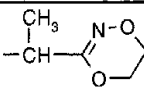

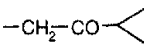
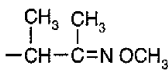
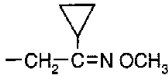
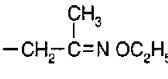
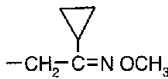
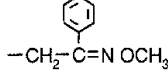
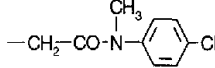
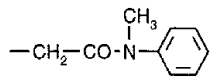
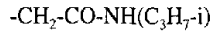
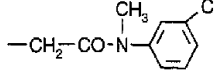
Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
20	F	F	H	H	H		70 ¹⁾
21	F	F	H	H	H		110-14 ¹⁾
22	F	F	H	H	H		120-22 ¹⁾
23	F	F	H	H	H		4.62 (A isomer)* 4.71 (B isomer)*
24	F	F	H	H	H		4.90 (E isomers)
25	F	F	H	H	H		4.70 (A isomer)* 4.76 (B isomer)*
26	F	F	H	H	H		4.90 (A isomer)* 4.56 (B isomer)*
27	F	F	H	H	H		4.74 (A isomer)*
28	F	F	H	H	H		60-65 ¹⁾
29	F	F	H	H	H		155-60 ¹⁾
30	F	F	H	H	H		156 ¹⁾
31	F	F	H	H	H		110-14 ¹⁾



Table 1 (continued)

Ex. No.	X	Y	Z	R _m ¹	R _n ²	A	m.p. (°C) ¹⁾ or log p
32	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{N}(\text{CH}_3)-\text{C}_6\text{H}_4-\text{CF}_3$	$n_D^{20} = 1.5644$
33	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{CH}_2-\text{C}_6\text{H}_5$	142-44 ¹⁾
34	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{N}(\text{CH}_3)-\text{OCH}_3$	3.07
35	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}_2$	185-88 ¹⁾
36	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_6\text{H}_4-\text{C}_2\text{H}_5$	110-12 ¹⁾
37	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_6\text{H}_4-\text{F}$	150-55 ¹⁾
38	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_4\text{H}_9-t$	110 ¹⁾
39	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_6\text{H}_5$	155-58 ¹⁾
40	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_6\text{H}_4-\text{Cl}$	164-66 ¹⁾
41	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_6\text{H}_3(\text{Cl})_2$	157-60 ¹⁾
42	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_6\text{H}_4-\text{OC}_2\text{H}_5$	168-70 ¹⁾
43	F	F	H	H	H	$-\text{CH}_2-\text{CH}(\text{OC}_2\text{H}_5)_2$	4.44
44	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_6\text{H}_3(\text{Cl})_2$	202 ¹⁾



Table 1 (continued)

Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
45	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}_6\text{H}_3(\text{CN})_2$	232-34 ¹⁾
46	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{NH}-\text{C}(\text{CH}_3)(\text{CF}_3)$	148-50 ¹⁾
47	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{N}(\text{CH}_2\text{CH}=\text{CH}_2)_2$	3.76
48	F	F	H	H	H	$-\text{CH}(\text{CH}_3)-\text{C}(\text{CH}_3)=\text{N}-\text{OC}_2\text{H}_5$	5.02 (A isomer)* 5.13 (B isomer)*
49	H	Cl	H	H	H	$-\text{CH}_2-\text{CO}-\text{C}_6\text{H}_5$	4.38
50	H	Cl	H	H	H	$-\text{CH}_2-\text{CO}-\text{CH}_3$	70-72 ¹⁾
51	H	Cl	H	H	H	$-\text{CH}_2-\text{C}_5\text{H}_4\text{N}-\text{Cl}$	135-37 ¹⁾
52	H	Cl	H	H	H	$-\text{CH}_2-\text{C}(\text{NOCH}_3)-\text{C}_6\text{H}_5$	5.40
53	H	Cl	H	H	H	$-\text{CH}_2-\text{C}(\text{CH}_3)=\text{NOCH}_3$	80-85 ¹⁾
54	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{C}(\text{Cl})(\text{CH}_2)_2$	4.04
55	F	F	H	H	H	$-\text{CH}_2-\text{C}(\text{Cl})(\text{NOCH}_3)-\text{C}_6\text{H}_5$	5.09 (A isomer)* 4.78 (B isomer)*
56	F	F	H	H	H	$-\text{CH}_2-\text{C}(\text{NOC}_2\text{H}_5)-\text{C}_6\text{H}_5$	5.28 (A isomer)* 4.92 (B isomer)*



Table 1 (continued)

Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
57	F	F	H	H	H	$\begin{array}{c} \text{Cl} \\ \\ -\text{CH}_2-\text{C}-\text{Cyclopropyl} \\ \\ \text{NOC}_2\text{H}_5 \end{array}$	5.49 (A isomer)* 4.85 (B isomer)*
5	58	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}-\text{CO}-\text{CH}_3 \end{array}$	77-79 ¹⁾
59	Cl	H	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}-\text{C}-\text{CH}_3 \\ \\ \text{NOC}_2\text{H}_5 \end{array}$	5.44 (A isomer)* 5.56 (B isomer)*
60	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}-\text{CO}-\text{C}_6\text{H}_4-\text{NHCOCH}_3 \end{array}$	95-96 ¹⁾
61	F	F	H	H	H	$\begin{array}{c} \text{C}_4\text{H}_9\text{-n} \\ \\ -\text{CH}-\text{COOC}_2\text{H}_5 \end{array}$	5.24
62	Cl	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}-\text{C}-\text{CH}_3 \\ \\ \text{NOC}_2\text{H}_5 \end{array}$	5.31 (A isomer)* 5.42 (B isomer)*
10	63	Cl	F	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}-\text{CO}-\text{CH}_3 \end{array}$	112-15 ¹⁾
64	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}-\text{C}-\text{CH}_3 \\ \\ \text{N-OH} \end{array}$	3.48 (A isomer)* 3.54 (B isomer)*



Table 1 (continued)

	Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
5	65	F	F	H	H	H	-CH ₂ -COC ₂ H ₅	122-24 ¹⁾
	66	F	F	H	H	H	$\begin{array}{c} \text{C}_2\text{H}_5 \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	4.98 (A isomer)*
							4.90 (B isomer)*	
	67	F	F	H	H	H	$\begin{array}{c} \text{C}_2\text{H}_5 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_3 \end{array}$	4.72 (A isomer)*
							4.61 (B isomer)*	
68	F	F	H	H	H	-CH ₂ -CO-C(CH ₃) ₂ C ₃ H ₇ -i	4.55	
69	F	F	H	H	H	$\begin{array}{cc} \text{CH}_3 & \text{CH}_3 \\ & \\ -\text{CH}-\text{C}=\text{NOC}_3\text{H}_7\text{-i} \end{array}$	4.47	
10	70	F	F	H	H	H	$\begin{array}{cc} \text{CH}_3 & \text{CH}_3 \\ & \\ -\text{CH}-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.12
	71	F	F	H	H	H	-CH ₂ -CO-C(CH ₃) ₂ CHCl ₂	4.55
	72	F	F	H	H	H	-CH ₂ -CO-C ₄ H ₉ -t	101-02 ¹⁾
	73	F	F	H	H	H	-CH ₂ -CO-C ₃ H ₇ -i	86-88 ¹⁾
	74	F	F	H	H	H	-C(CH ₃) ₂ -COCH ₃	102-06 ¹⁾
15	75	F	F	H	H	H	-CH ₂ -CO-C(CH ₃) ₂ -CH=CCl ₂	92-94 ¹⁾
	76	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOH} \end{array}$	129-36 ¹⁾
	77	Cl	F	H	H	H	-CH ₂ -COCH ₃	92 ¹⁾
	78	F	F	H	H	H	$\begin{array}{c} \text{C}_3\text{H}_7\text{-i} \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.72 (A isomer)*
							5.49 (B isomer)*	
	79	F	F	H	H	H	$\begin{array}{c} \text{C}_4\text{H}_9\text{-t} \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.95



Table 1 (continued)

Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
5	80	F	F	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_3\text{H}_7\text{-i} \end{array}$	5.10 (A isomer)* 5.17 (B isomer)*
	81	Cl	F	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	4.98
	82	F	F	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_2-\text{C}_6\text{H}_5 \end{array}$	5.15 (A isomer)* 5.22 (B isomer)*
	83	F	F	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.49 (A isomer)* 5.56 (B isomer)*
	84	F	F	H	3-Cl	-CH ₂ -COCH ₃	118-19 ¹⁾
10	85	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \quad \text{CH}_3 \\ \quad \\ -\text{CH}-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.40
	86	F	F	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_3 \end{array}$	4.79 (A isomer)* 4.70 (B isomer)*
	87	F	F	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.39 (A isomer)* 5.19 (B isomer)*
	88	F	F	H	H	$\begin{array}{c} \text{CH}_3 \quad \text{CH}_3 \\ \quad \\ -\text{CH}-\text{C}=\text{NOCH}_2\text{-CH=CH}_2 \end{array}$	5.04 (A isomer)* 5.13 (B isomer)*
	89	F	F	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_2\text{-CH=CH}_2 \end{array}$	4.78 (A isomer)* 4.72 (B isomer)*



Table 1 (continued)

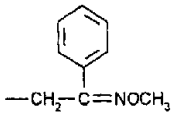
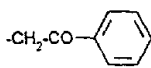
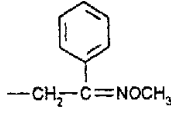

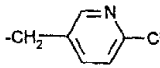
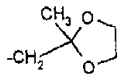
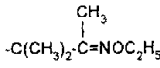
Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
90	F	F	H	H	3-Cl		95 ¹⁾
91	F	F	H	H	3-Cl	-CH(CN)C ₃ H _{7-n}	4.87
92	F	F	H	H	3-Cl	-(CH ₂) ₃ -CN	108-11 ¹⁾
93	F	Cl	H	H	H		4.28
94	F	Cl	H	H	H		5.25
95	F	F	H	H	3-Cl		4.06
96	F	F	H	H	3-Cl		107-109 ¹⁾
97	F	F	H	H	3-Cl	-CH ₂ CH ₂ CH(OC ₂ H ₅) ₂	5.12
98	F	F	H	H	H		3.77
99	F	F	H	H	H		5.57



Table 1 (continued)

Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
100	F	F	H	H	H	$\text{-C(CH}_3)_2\text{-C(=NOCH}_3\text{)-CH}_3$	5.14
101	F	F	H	H	3-Cl	$\text{-CH(CH}_3\text{)-COCH}_3$	5.14
102	F	F	H	H	3-Cl	$\text{-CH(C}_3\text{H}_7\text{-i)-CN}$	4.84
103	F	F	H	H	H	$\text{-CH}_2\text{-CO-C}_6\text{H}_4\text{-Br}$	121-23 ¹⁾
104	F	F	H	H	H	$\text{-CH}_2\text{-CO-C}_6\text{H}_4\text{-Cl}$	154-55 ¹⁾
105	F	F	H	H	H	$\text{-CH}_2\text{-CO-C}_6\text{H}_4\text{-CH}_3$	124-25 ¹⁾
106	F	F	H	H	H	$\text{-CH(CH}_3\text{)-C(=NOCH}_3\text{)-C}_6\text{H}_4\text{-CF}_3$	5.50
107	F	F	H	H	H	$\text{-CH}_2\text{-CO-C}_6\text{H}_3\text{(Cl)-CH}_3$	82-84 ¹⁾
108	F	F	H	H	H	$\text{-CH(CH}_3\text{)-C(=NOCH}_3\text{)-C}_6\text{H}_4\text{-F}$	5.01



Table 1 (continued)

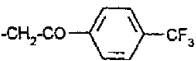
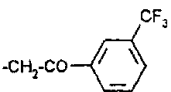
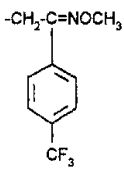
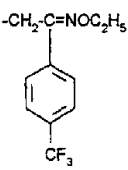
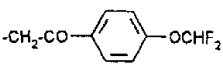
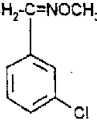
Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
109	F	F	H	H	H		142-44 ¹⁾
110	F	F	H	H	H		116-18 ¹⁾
111	F	F	H	H	H		5.53
112	F	F	H	H	H		5.87 (A isomer)* 5.57 (B isomer)*
113	H	Cl	H	H	3-Cl	-CH(CH ₃)-COCH ₃	4.51
114	F	F	H	H	H		4.16
115	F	F	H	H	H		5.52 (A isomer)* 5.19 (B isomer)*



Table 1 (continued)


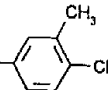

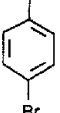

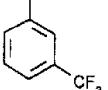
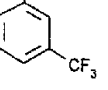
Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
116	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	5.89 (A isomer)* 5.54 (B isomer)*
117	F	F	H	H	H	$-\text{CH}_2-\text{CO}-$ 	4.86
118	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOCH}_3$ 	5.65 (A isomer)* 5.57 (B isomer)*
119	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	6.03 (A isomer)* 5.72 (B isomer)*
120	F	Cl	H	H	H	$-\text{CH}(\text{CH}_3)-\text{C}=\text{NOCH}_2\text{CH}=\text{CH}_2$ 	5.70 (A isomer)* 5.81 (B isomer)*
121	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOCH}_3$ 	5.80 (A isomer)* 4.49 (B isomer)*
122	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	6.18 (A isomer)* 5.87 (B isomer)*



Table 1 (continued)

Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
123	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}(\text{CH}_3)-\text{C}=\text{NOC}_3\text{H}_7-i \end{array}$	6.15 (A isomer)* 6.30 (B isomer)*
124	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_3\text{H}_7-i \end{array}$	5.78 (A isomer)* 5.88 (B isomer)*
125	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_2\text{CH}=\text{CH}_2 \end{array}$	5.04 (A isomer)* 5.11 (B isomer)*
126	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_2-\text{C}_6\text{H}_5 \end{array}$	5.45 (A isomer)* 5.53 (B isomer)*
127	F	F	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}(\text{CH}_3)-\text{C}=\text{NOCH}_3 \end{array}$	5.06 (A isomer)* 5.15 (B isomer)*
128	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{C}_6\text{H}_4-\text{OCH}_3$	135-37 ¹⁾
129	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{C}_6\text{H}_4-\text{CH}_2\text{C}_3\text{H}_7-i$	118-20 ¹⁾
130	F	F	H	H	H	$\begin{array}{c} -\text{CH}_2-\text{C}=\text{NOCH}_3 \\ \\ \text{C}_6\text{H}_4 \\ \\ \text{Cl} \end{array}$	5.27 (A isomer)* 5.54 (B isomer)*



Table 1 (continued)

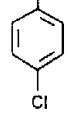
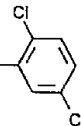
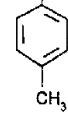


Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
131	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	5.61 (A isomer)* 5.54 (B isomer)*.
132	F	F	H	H	H	 $-\text{CH}_2-\text{CO}-$	4.78
133	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOCH}_3$ 	5.07 (A isomer)* 5.35 (B isomer)*
134	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	5.42 (A isomer)* 5.71 (B isomer)*
135	H	Cl	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_3\text{H}_7-i$ 	5.53 (A isomer)* 5.61 (B isomer)*



Table 1 (continued)

Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
136	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}(\text{CH}_3)-\text{C}=\text{NOC}_3\text{H}_7\text{-i} \end{array}$	5.88 (A isomer)* 6.02 (B isomer)*
137	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_2-\text{C}_6\text{H}_5 \end{array}$	5.55 (A isomer)* 5.62 (B isomer)*
138	F	F	H	H	H	$\begin{array}{c} -\text{CH}_2-\text{C}=\text{NOCH}_3 \\ \\ \text{C}_6\text{H}_4 \end{array}$	4.78 (A isomer)* 4.98 (B isomer)*
139	F	F	H	H	H	$\begin{array}{c} \text{CHF}_2 \\ \\ -\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5 \\ \\ \text{C}_6\text{H}_4 \end{array}$	5.07 (A isomer) 5.31 (B isomer)
140	F	F	H	H	H	$\begin{array}{c} \text{CHF}_2 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_3 \\ \\ \text{C}_6\text{H}_3(\text{CH}_3)(\text{Cl}) \end{array}$	5.57 (A isomer)* 5.88 (B isomer)*
141	F	F	H	H	H	$\begin{array}{c} -\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5 \\ \\ \text{C}_6\text{H}_3(\text{CH}_3)(\text{Cl}) \end{array}$	5.93 (A isomer)* 6.23 (B isomer)*
142	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}_2-\text{C}=\text{NOCH}_2-\text{CH}=\text{CH}_2 \end{array}$	5.12 (A isomer)* 5.20 (B isomer)*



Table 1 (continued)

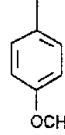
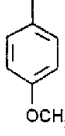
Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
143	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}(\text{CH}_3)-\text{C}=\text{NOCH}_2-\text{CH}=\text{CH}_2 \end{array}$	5.45 (A isomer)* 5.57 (B isomer)*
144	H	Cl	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}(\text{CH}_3)-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.83 (A isomer)* 6.01 (B isomer)*
145	H	Cl	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}(\text{CH}_3)-\text{C}=\text{NOC}_3\text{H}_7-i \end{array}$	6.30 (A isomer)* 6.44 (B isomer)*
146	F	F	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}(\text{CH}_3)-\text{C}=\text{NOC}_3\text{H}_7-i \end{array}$	5.87 (A isomer)* 6.00 (B isomer)*
147	F	F	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{CH}(\text{CH}_3)-\text{C}=\text{NOCH}_2-\text{CH}=\text{CH}_2 \end{array}$	5.49 (A isomer)* 5.58 (B isomer)*
148	F	F	H	H	H	$-\text{CH}_2-\text{CO}-\text{C}_6\text{H}_3(\text{CH}_3)_2$	4.62
149	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOCH}_3$ 	4.70 (A isomer)* 4.91 (B isomer)*
150	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	5.04 (A isomer)* 5.27 (B isomer)*



Table 1 (continued)


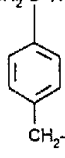


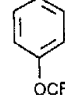
Ex. No.	X	Y	Z	R _m ¹	R _n ²	A	m.p. (°C) ¹⁾ or log p
151	F	Cl	H	H	H	-C(CH ₃) ₂ -COCH ₃	4.48
152	H	Cl	H	H	H	-C(CH ₃) ₂ -COCH ₃	4.56
153	F	F	H	H	3-Cl	-C(CH ₃) ₂ -COCH ₃	4.61
154	F	F	H	H	H	-CH ₂ -C=NOCH ₃ 	6.34 (A isomer)* 6.05 (B isomer)*
155	F	F	H	H	H	-CH ₂ -C=NOCH ₂ C ₆ H ₄ -CH ₂ -C ₃ H ₇ -i 	6.70 (A isomer)* 6.36 (B isomer)*
156	F	F	H	H	H	-CH ₂ -CO-  -OCF ₂ -CHF ₂	84-85 ¹⁾
157	F	F	H	H	H	-CH ₂ -C=NOCH ₃  OCF ₂ -CHF ₂	5.28 (A isomer)* 5.04 (B isomer)*
158	F	F	H	H	H	-CH ₂ -C=NOCH ₂ C ₆ H ₄ -CH ₂ -C ₃ H ₇ -i 	5.59 (A isomer)* 5.35 (B isomer)*



Table 1 (continued)

Ex. No.	X	Y	Z	R _m ¹	R _n ²	A	m.p. (°C) ¹⁾ or log p
159	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOH} \end{array}$	
160	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOCH}_3 \end{array}$	5.45
161	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.88
162	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_3\text{H}_7\text{-n} \end{array}$	
163	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOH} \end{array}$	
164	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOCH}_3 \end{array}$	
165	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	
166	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_3\text{H}_7\text{-n} \end{array}$	
167	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOH} \end{array}$	3.82 (A isomer)* 3.39 (B isomer)*



Table 1 (continued)

Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
168	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_3\text{H}_7-n \end{array}$	
169	F	F	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOH} \end{array}$	203-205 ¹⁾
170	F	F	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOCH}_3 \end{array}$	5.61
171	F	F	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	5.05
172	F	F	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_3\text{H}_7-n \end{array}$	
173	H	Cl	H	H	3-Cl	$-\text{C}(\text{CH}_3)_2-\text{CO}-\text{CH}_3$	5.02
174	H	Cl	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOH} \end{array}$	
175	H	Cl	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOCH}_3 \end{array}$	6.06
176	H	Cl	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_2\text{H}_5 \end{array}$	6.49
177	H	Cl	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ -\text{C}(\text{CH}_3)_2-\text{C}=\text{NOC}_3\text{H}_7-n \end{array}$	



Table 1 (continued)

Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
178	F	F	H	H	H	$\begin{array}{c} \text{---CH}_2\text{---C=NOC}_2\text{H}_5 \\ \\ \text{C}_6\text{H}_3\text{Cl}_2 \end{array}$	
179	F	F	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \\ \\ \text{---C(CH}_3)_2\text{---C=NOC}_3\text{H}_7\text{---n} \end{array}$	
180	H	Cl	H	H	3-Cl	$\begin{array}{c} \text{CH}_3 \quad \text{CH}_3 \\ \quad \quad \\ \text{---CH---C=NOC}_3\text{H}_7\text{---n} \end{array}$	
181	Cl	F	H	H	H	$\begin{array}{c} \text{CH}_3 \quad \text{CH}_3 \\ \quad \quad \\ \text{---CH---C=NOC}_3\text{H}_7\text{---n} \end{array}$	
182	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ \text{---CH}_2\text{---C=NOC}_3\text{H}_7\text{---n} \end{array}$	
183	H	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ \text{---CH(CH}_3\text{)---C=NOC}_3\text{H}_7\text{---n} \end{array}$	
184	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ \text{---CH(CH}_3\text{)---C=NOC}_3\text{H}_7\text{---n} \end{array}$	
185	F	Cl	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ \text{---CH}_2\text{---C=NOC}_3\text{H}_7\text{---n} \end{array}$	
186	F	F	H	H	H	$\begin{array}{c} \text{CH}_3 \\ \\ \text{---CH}_2\text{---C=NOC}_3\text{H}_7\text{---n} \end{array}$	



Table 1 (continued)



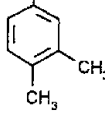
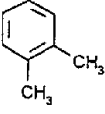
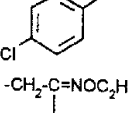
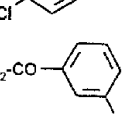
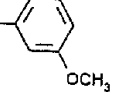
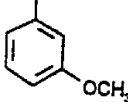
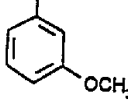
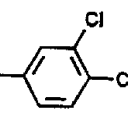
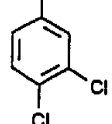
Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
187	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_3\text{H}_7-n$ 	
188	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_3\text{H}_7-n$ 	
189	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOCH}_3$ 	5.58 (A isomer)* 5.29 (B isomer)*
190	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	5.94 (A isomer)* 5.62 (B isomer)*
191	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOCH}_3$ 	5.62 (A isomer)* 5.44 (B isomer)*
192	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	6.00 (A isomer)* 5.79 (B isomer)*
193	F	F	H	H	H	$-\text{CH}_2-\text{CO}-$ 	



Table 1 (continued)

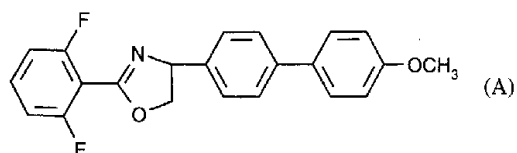
Ex. No.	X	Y	Z	R ¹ _m	R ² _n	A	m.p. (°C) ¹⁾ or log p
194	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOCH}_3$ 	
195	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOC}_2\text{H}_5$ 	
196	F	F	H	H	H	$-\text{CH}_2-\text{CO}-$ 	
197	F	F	H	H	H	$-\text{CH}_2-\text{C}=\text{NOCH}_3$ 	

* A isomer and B isomer: possible geometric



Use examples

The compound of the formula (A)



known from EP-A 0 432 661 was employed as the comparison substance in the
5 following use examples.



Example A

Phaedon larvae test

Solvent: 7 parts by weight of dimethylformamide

Emulsifier: 1 part by weight of alkylaryl polyglycol ether

- 5 To produce a suitable preparation of active compound, 1 part by weight of active compound is mixed with the stated amount of solvent and the stated amount of emulsifier, and the concentrate is diluted with water to the desired concentration.

- Cabbage leaves (*Brassica oleracea*) are treated by being dipped into the preparation of the active compound of the desired concentration and are infested with mustard
10 beetle larvae (*Phaedon cochleariae*) while the leaves are still moist.

After the desired time, the destruction in % is determined. 100% means that all the beetle larvae have been killed; 0% means that none of the beetle larvae have been killed.

- In this test, for example, the compounds of Preparation Examples 1, 2, 4, 13, 14, 16,
15 20 and 21 caused a destruction of 100 % after 7 days at an active compound concentration of, for example, 0.01 %, while the known compound (A) showed no destruction.



Example: B

Plutella test

Solvent: 7 parts by weight of dimethylformamide

Emulsifier: 1 part by weight of alkylaryl polyglycol ether

- 5 To produce a suitable preparation of active compound, 1 part by weight of active compound is mixed with the stated amount of solvent and the stated amount of emulsifier, and the concentrate is diluted with water to the desired concentration.

- 10 Cabbage leaves (*Brassica oleracea*) are treated by being dipped into the preparation of active compound of the desired concentration and are infested with caterpillars of the diamond-back moth (*Plutella maculipennis*) while the leaves are still moist.

After the desired time, the destruction in % is determined. 100% means that all the caterpillars have been killed; 0% means that none of the caterpillars have been killed.

- 15 In this test, for example, the compounds of Preparation Examples 2 and 4 caused a destruction of 100 % after 7 days at an active compound concentration of, for example, 0.0001 %, while the known compound (A) caused a destruction of only 5 %.



Example C

Myzus test

Solvent: 7 parts by weight of dimethylformamide

Emulsifier: 1 part by weight of alkylaryl polyglycol ether

- 5 To produce a suitable preparation of active compound, 1 part by weight of active compound is mixed with the stated amount of solvent and the stated amount of emulsifier and the concentrate is diluted with water to the desired concentration.

- Seedlings of broad beans (*Vicia faba*) infested by the green peach aphid (*Myzus persicae*) are dipped into the preparation of active compound of the desired
10 concentration and placed in a plastic container.

After the desired time, the destruction in % is determined. 100% means that all the aphids have been killed; 0 % means that none of the aphids has been killed.

- In this test, for example, the compounds of Preparation Examples 4, 17 and 21 caused a destruction of 95 % to 98 % after 6 days at an active compound
15 concentration of, for example, 0.01 %, while the known compound (A) caused a destruction of only 5 %.



Example D

Spodoptera Frugiperda test

Solvent: 7 parts by weight of dimethylformamide

Emulsifier: 1 part by weight of alkylaryl polyglycol ether

- 5 To produce a suitable preparation of active compound, 1 part by weight of active compound is mixed with the stated amount of solvent and the stated amount of emulsifier, and the concentrate is diluted with water to the desired concentration.

- 10 Cabbage leaves (*Brassica oleracea*) are treated by being dipped into the preparation of active compound of the desired concentration and are infested with caterpillars of the owl moth (*Spodoptera frugiperda*) while the leaves are still moist.

After the desired time, the destruction in % is determined. 100% means that all the caterpillars have been killed; 0% means that none of the caterpillars have been killed.

- 15 In this test, for example, the compounds of Preparation Examples 1, 2, 4, 10, 14, 16, 18, 20 and 21 caused a destruction of 100 % after 7 days at an active compound concentration of, for example, 0.1 %, while the known compound (A) showed no destruction.



Example E

Tetranychus test (OP-resistant/dipping treatment)

Solvent: 7 parts by weight of dimethylformamide

Emulsifier: 1 part by weight of alkylaryl polyglycol ether

- 5 To produce a suitable preparation of active compound, 1 part by weight of active compound is mixed with the stated amount of solvent and the stated amount of emulsifier, and the concentrate is diluted with emulsifier-containing water to the desired concentrations.

- 10 Bean plants (*Phaseolus vulgaris*) which are heavily infested with all stages of the two-spotted spider mite (*Tetranychus urticae*) are treated by being dipped into a preparation of the active compound of the desired concentration.

After the desired time, the effect in % is determined. 100% means that all the spider mites have been killed; 0% means that none of the spider mites have been killed.

- 15 In this test, for example, the compounds of Preparation Examples 1 and 2 caused a destruction of 98 % after 13 days at an active compound concentration of, for example, 0.001 %, while the known compound (A) showed a destruction of only 45 %.



Example F

Test with fly larvae/development-inhibiting action

- Test animals: All larval stages of *Lucilia cuprina* (OP-resistant)
[Pupae and adults (without contact with the active compound)]
- 5 Solvent: 35 parts by weight of ethylene glycol monomethyl ether
Emulsifier: 35 parts by weight of nonylphenol polyglycol ether

To produce a suitable formulation, three parts by weight of active compound are mixed with seven parts by weight of the abovementioned solvent/emulsifier mixture and the emulsion concentrate thus obtained is diluted with water to the particular
10 desired concentration.

30 - 50 larvae per concentration are placed on horse meat (1 cm³) in glass tubes, onto which 500 µl of the dilution to be tested are pipetted. The glass tubes are placed in a plastic beaker, the base of which is covered with sea sand, and are kept in a climatically controlled room (26°C ± 1.5°C, 70 % relative humidity ± 10 %).

15 The action is monitored after 24 hours and 48 hours (larvicidal action). After the larvae have migrated (approx. 72 hours), the glass tubes are removed and perforated plastic lids are placed on the beakers. After 1½ times the duration of development (hatching of the control flies), the flies which have hatched and the pupae/pupal shells are counted.

- 20 The criterion for action is the occurrence of death in the treated larvae after 48 hours (larvicidal effect) or the inhibition of the hatching of adults from the pupae or the inhibition of pupation. The criterion for the in vitro action of a substance is the inhibition of flea development or a stop in development before the adult stage. 100 % larvicidal action here means that all the larvae have died after 48 hours.
- 25 100 % development-inhibiting action means that no adult flies have hatched.



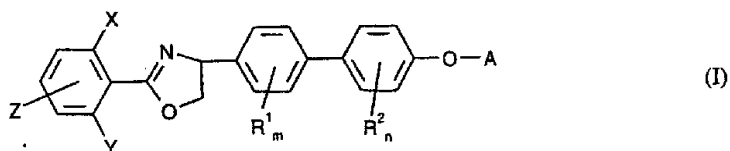
- 70 -

In this test, for example, the compounds of Preparation Examples 4, 13, 15, 16, 17, 18, 20, 21, 23, 24, 25 and 26 caused a destruction of 100 % at an active compound concentration of, for example, 1000 ppm.



The claims defining the invention are as follows:

1. Compounds of the formula (I)



in which

X represents hydrogen, fluorine or chlorine,

Y represents fluorine, chlorine or methyl,

Z represents hydrogen, halogen, alkyl, alkoxy or dialkylamino,

R¹ and R² independently of one another represent halogen, alkyl, alkoxy, alkylthio, halogenoalkyl, halogenoalkoxy or halogenoalkylthio,

m and n independently of one another represent 0, 1 or 2 and

A represents the grouping $-(CH_2)_p-CR^3R^4-(CH_2)_r-R$ wherein

R³ and R⁴ independently of one another represent hydrogen or alkyl,

p, q and r independently of one another represent 0, 1, 2 or 3, at least one index being other than 0, and

R represents cyano; or represents an optionally substituted, saturated, partly saturated or unsaturated 5- or 6-membered heterocyclic radical, or represents one of the following groupings:



- (a) ---CO-R^5
- (b) ---CO-OR^6
- (c) $\text{---CO-NR}^7\text{R}^8$
- (d) $\text{---CS-NR}^7\text{R}^8$
- (e) $\begin{array}{c} \text{---C=N-R}^{11} \\ | \\ \text{R}^5 \end{array}$
- (f) $\begin{array}{c} \text{OR}^{12} \\ \diagup \\ \text{---C} \\ \diagdown \text{OR}^{12} \\ | \\ \text{R}^5 \end{array}$
- (g) $\begin{array}{c} \text{SR}^{12} \\ \diagup \\ \text{---C} \\ \diagdown \text{SR}^{12} \\ | \\ \text{R}^5 \end{array}$
- (h) $\begin{array}{c} \text{R}^{13} \\ \diagup \\ \text{N} \\ \diagdown \text{R}^{14} \\ | \\ \text{---C---OR}^{12} \\ | \\ \text{R}^5 \end{array}$
- (i) $\begin{array}{c} \text{R}^{13} \\ \diagup \\ \text{N} \\ \diagdown \text{R}^{14} \\ | \\ \text{---C---SR}^{12} \\ | \\ \text{R}^5 \end{array}$
- (j) $\begin{array}{c} \text{---C=N-R}^9 \\ | \\ \text{OR}^{10} \end{array}$
- (k) $\begin{array}{c} \text{---C=N-R}^9 \\ | \\ \text{SR}^{10} \end{array}$

wherein

5 R^5 represents hydrogen, alkyl, halogenoalkyl, alkenyl, halogenoalkenyl, optionally substituted cycloalkyl or optionally substituted aryl,

R^6 represents hydrogen, alkyl, alkenyl, halogenoalkyl, halogenoalkenyl, optionally substituted arylalkyl or in each case optionally substituted cycloalkyl or cycloalkylalkyl,



R^7 and R^8 independently of one another represent hydrogen, alkyl, alkoxy, alkenyl, halogenoalkyl, halogenoalkenyl, in each case optionally substituted aryl or arylalkyl, or in each case optionally substituted cycloalkyl or cycloalkylalkyl, or represent $-OR^6$ or $-NR^5R^6$,

wherein

R^5 and R^6 have the abovementioned meaning,

or

R^7 and R^8 together represent a 5- or 6-membered alkylene chain, which optionally contains an oxygen atom,

R^9 and R^{10} independently of one another represent alkyl,

R^{11} represents $-OR^6$, $-NR^5R^6$ or $-N(R^5)-COOR^6$,

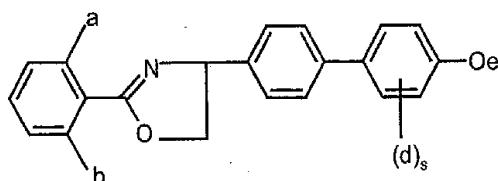
wherein

R^5 and R^6 have the abovementioned meaning,

and

R^{12} , R^{13} and R^{14} independently of one another represent alkyl;

provided that compounds of the formula



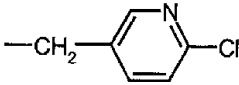
wherein

- a represents hydrogen,
- b represents fluorine or chlorine,
- e represents optionally substituted heteroarylalkyl,
- d represents halogen, alkyl or alkoxy and
- s represents 0, 1 or 2

or

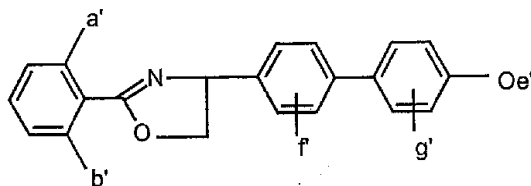
wherein

- a represents fluorine,
- b represents fluorine

- c represents  and
s represents 0,

are specifically excluded

and that the compounds of the formula



wherein

- a' represents hydrogen, fluorine or chlorine
 - b' represents fluorine and chlorine
 - e' represents C₂-C₄-cyanoalkyl
 - f' represents hydrogen, halogen, alkyl, alkoxy, halogenoalkyl, halogenoalkoxy
- and
- g' represents hydrogen, alkyl, alkoxy, halogenoalkyl, halogenoalkoxy,
- are specifically excluded.

2. Compounds of the formula (I) according to Claim 1, in which

- X represents hydrogen, fluorine or chlorine,
- Y represents fluorine, chlorine or methyl,



Z represents hydrogen, fluorine, chlorine, C₁-C₄-alkyl, C₁-C₄-alkoxy or di(C₁-C₄)alkylamino,

R¹ and R² independently of one another represent fluorine, chlorine, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy or C₁-C₄-halogenoalkylthio,

m and n independently of one another represent 0, 1 or 2,

A represents the grouping (CH₂)_p-(CR³R⁴)_q-(CH₂)_r-R,
wherein

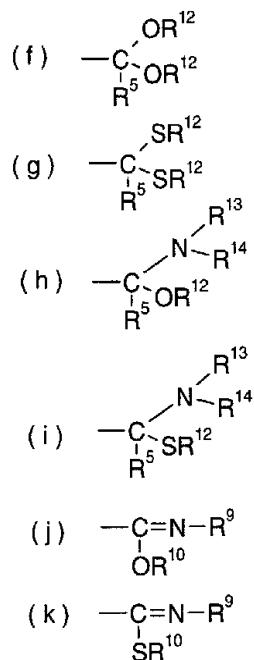
R³ and R⁴ independently of one another represent hydrogen or C₁-C₄-alkyl,

p, q and r independently of one another represent 0, 1, 2 or 3, at least one index being other than 0 and the sum of the indices being not greater than 5,

R represents cyano; or represents a saturated, partly saturated or unsaturated 5- or 6-membered heterocyclic radical which has 1 to 3 identical or different heteroatoms from the series consisting of nitrogen, oxygen and sulfur and is optionally substituted by halogen, C₁-C₄-alkyl or C₁-C₄-halogenoalkyl,
or represents one of the following groupings:

- (a) —CO—R⁵
- (b) —CO—OR⁶
- (c) —CO—NR⁷R⁸
- (d) —CS—NR⁷R⁸
- (e) —C=N—R¹¹
|
R⁵





wherein

5

10

R^5 represents hydrogen, $\text{C}_1\text{-C}_6\text{-alkyl}$, $\text{C}_1\text{-C}_4\text{-halogenoalkyl}$, $\text{C}_2\text{-C}_6\text{-alkenyl}$ or $\text{C}_2\text{-C}_6\text{-halogenoalkenyl}$; or $\text{C}_3\text{-C}_6\text{-cycloalkyl}$ which is optionally substituted once or several times in an identical or different manner by $\text{C}_1\text{-C}_4\text{-alkyl}$, halogen or $\text{C}_1\text{-C}_4\text{-halogenoalkyl}$, or represents phenyl which is optionally substituted once to five times in an identical or different manner by halogen, $\text{C}_1\text{-C}_4\text{-alkyl}$, $\text{C}_1\text{-C}_4\text{-alkoxy}$, $\text{C}_1\text{-C}_4\text{-alkylthio}$, $\text{C}_1\text{-C}_4\text{-halogenoalkyl}$, $\text{C}_1\text{-C}_4\text{-halogenoalkoxy}$, $\text{C}_1\text{-C}_4\text{-halogenoalkylthio}$, $\text{di}(\text{C}_1\text{-C}_4\text{-alkylamino})$, $\text{C}_1\text{-C}_4\text{-alkylcarbonylamino}$, $\text{C}_1\text{-C}_4\text{-alkylcarbonyl-C}_1\text{-C}_4\text{-alkylamino}$, cyano, nitro, $-\text{COOR}^{15}$ or $-\text{CONR}^{16}\text{R}^{17}$,

R^5 represents hydrogen, $\text{C}_1\text{-C}_6\text{-alkyl}$, $\text{C}_1\text{-C}_6\text{-halogenoalkyl}$, $\text{C}_2\text{-C}_6\text{-alkenyl}$ or $\text{C}_2\text{-C}_6\text{-halogenoalkenyl}$; or $\text{C}_3\text{-C}_6\text{-cycloalkyl}$ or $\text{C}_3\text{-C}_6\text{-cycloalkyl-C}_1\text{-C}_4\text{-alkyl}$, in each case optionally substituted once or several times



in an identical or different manner by C₁-C₄-alkyl, halogen or C₁-C₄-halogenoalkyl,

or represents C₆-C₁₀-aryl-C₁-C₆-alkyl which is optionally substituted once to five times in an identical or different manner by halogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy, C₁-C₄-halogenoalkylthio, di(C₁-C₄)alkylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷,

R⁷ and R⁸ independently of one another represent hydrogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-halogenoalkyl, C₂-C₆-alkenyl or C₂-C₆-halogenoalkenyl; or C₃-C₆-cycloalkyl or C₃-C₆-cycloalkyl-C₁-C₄-alkyl, in each case optionally substituted once or several times in an identical or different manner by C₁-C₄-alkyl, halogen or C₁-C₄-halogenoalkyl,

or represent phenyl or phenyl-C₁-C₄-alkyl, in each case optionally substituted once to five times in an identical or different manner by halogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy, C₁-C₄-halogenoalkylthio, di(C₁-C₄)alkylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷, or represent -OR⁶ or -NR⁵R⁶,

or

R⁷ and R⁸ together represent -(CH₂)₅-, -(CH₂)₆- or -(CH₂)₂-O-(CH₂)₂-;

R⁹ and R¹⁰ independently of one another represent C₁-C₄-alkyl;

R¹¹ represents -OR⁶-, -NR⁵R⁶ or -N(R⁵)-COOR⁶,

and

R¹², R¹³ and R¹⁴ independently of one another represent C₁-C₄-alkyl,



R¹⁵ represents hydrogen, C₁-C₄-alkyl or C₁-C₄-halogenoalkyl; or C₃-C₆-cycloalkyl which is optionally substituted once or several times in an identical or different manner by C₁-C₄-alkyl, halogen or C₁-C₄-halogenoalkyl,

5 or represents phenyl which is optionally substituted once to five times in an identical or different manner by halogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy, C₁-C₄-halogenoalkylthio, di(C₁-C₄)alkylamino, cyano, nitro, C₁-C₆-alkoxycarbonyl or di(C₁-C₆)alkylaminocarbonyl,

10 R¹⁶ and R¹⁷ independently of one another represent hydrogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-halogenoalkyl, C₂-C₆-alkenyl or C₂-C₆-halogenoalkenyl; or C₃-C₆-cycloalkyl or C₃-C₆-cycloalkyl-C₁-C₄-alkyl, in each case optionally substituted once or several times in an identical or different manner by C₁-C₄-alkyl, halogen or C₁-C₄-halogenoalkyl,

15 or represent phenyl or phenyl-C₁-C₄-alkyl, in each case optionally substituted once to five times in an identical or different manner by halogen, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-halogenoalkyl, C₁-C₄-halogenoalkoxy, C₁-C₄-halogenoalkylthio, di(C₁-C₄)alkylamino, cyano, nitro, C₁-C₆-alkoxycarbonyl or di(C₁-C₆)alkylaminocarbonyl,

20 or represent -OR⁶ or -NR⁶,

or

R¹⁶ and R¹⁷ together represent -(CH₂)₅-, -(CH₂)₆- or -(CH₂)₂-O-(CH₂)₂.

3. Compounds of the formula (I) according to Claim 1, in which

X represents hydrogen, fluorine or chlorine,

25 Y represents fluorine or chlorine,



Z represents hydrogen, fluorine, chlorine, methyl, ethyl, methoxy, dimethylamino or diethylamino,

R¹ and R² independently of one another represent fluorine, chlorine, methyl, ethyl, methoxy, methylthio, trifluoromethyl, trifluoromethoxy or trifluoromethylthio,

m and n independently of one another represent 0, 1 or 2.,

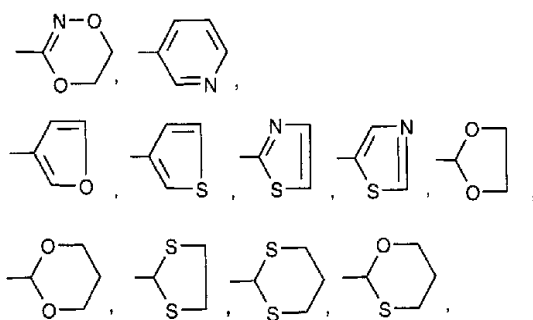
A represents the grouping (CH₂)_p-(CR³R⁴)_q-(CH₂)_r-R,

wherein

R³ and R⁴ independently of one another represent hydrogen, methyl, ethyl, n- or i-propyl, n-, i-, s- or t-butyl;

p, q and r independently of one another represent 0, 1, 2 or 3, at least one index being other than 0,

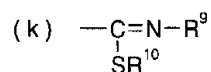
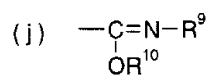
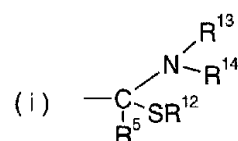
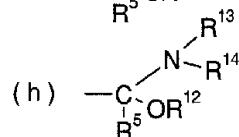
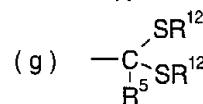
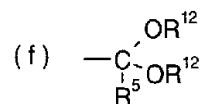
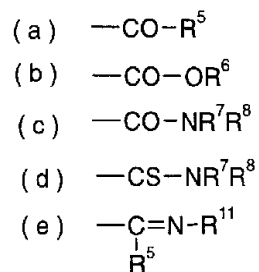
R represents cyano; or represents one of the following heterocyclic radicals



which are optionally substituted once to three times in an identical or different manner by fluorine, chlorine, bromine, methyl, ethyl, n- or i-propyl or trifluoromethyl;



or represents one of the following groupings:



5

wherein

R⁵ represents hydrogen; methyl, ethyl, n- or i-propyl, the isomeric butyls, the isomeric pentyls or the isomeric hexyls; or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms, C₃-C₆-alkenyl or C₃-C₆-halogenoalkenyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or cyclopropyl,



cyclopentyl or cyclohexyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms,
5 or represents phenyl which is optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, methylcarbonylamino, ethylcarbonylamino, methylcarbonyl-
10 methylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷,

R⁶ represents hydrogen, methyl, ethyl, n- or i-propyl or n-, i-, s- or t-butyl; or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or allyl which is optionally substituted once or several times by fluorine and/or chlorine; or
15 cyclopropyl, cyclopentyl, cyclohexyl, cyclopropyl-C₁-C₂-alkyl, cyclopentyl-C₁-C₂-alkyl or cyclohexyl-C₁-C₂-alkyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such
20 as F and Cl atoms,
or represents phenyl-C₁-C₂-alkyl or naphthyl-C₁-C₂-alkyl, in each case optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio,
25 dimethylamino, diethylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷,

R⁷ and R⁸ independently of one another represent hydrogen, C₁-C₄-alkyl, methoxy, ethoxy or C₁-C₃-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or allyl which is optionally substituted once or several times by fluorine and/or chlorine; or cyclopropyl, cyclopentyl, cyclohexyl, cyclopropylmethyl,



5 cyclopentylmethyl or cyclohexylmethyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or trifluoromethyl, or represent phenyl or phenyl-C₁-C₂-alkyl, in each case optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, cyano, nitro, -COOR¹⁵ or -CONR¹⁶R¹⁷, or represent -OR⁶ or -NR⁵R⁶,

10 R⁹ and R¹⁰ independently of one another represent methyl, ethyl or n- or i-propyl;

R¹¹ represents -OR⁶, -NR⁵R⁶ or -N(R⁵)-COOR⁶,

and

15 R¹², R¹³ and R¹⁴ independently of one another represent methyl, ethyl or n- or i-propyl.

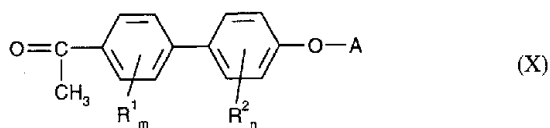
20 R¹⁵ represents hydrogen, methyl, ethyl, or n- or i-propyl; or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or cyclopropyl, cyclopentyl or cyclohexyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or C₁-C₂-halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms,

25 or represents phenyl which is optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, cyano, nitro, C₁-C₄-alkoxycarbonyl or di(C₁-C₄)alkylaminocarbonyl.



R^{16} and R^{17} independently of one another represent hydrogen, C_1 - C_4 -alkyl, methoxy, ethoxy or C_1 - C_3 -halogenoalkyl having 1 to 5 identical or different halogen atoms, such as F and Cl atoms; or allyl which is optionally substituted once or several times by fluorine and/or chlorine; or cyclopropyl, cyclopentyl, cyclohexyl, cyclopropylmethyl, cyclopentylmethyl or cyclohexylmethyl, in each case optionally substituted once to three times in an identical or different manner by fluorine, chlorine, methyl, ethyl, n- or i-propyl or trifluoromethyl, or represent phenyl or phenyl- C_1 - C_2 -alkyl, in each case optionally substituted once or twice in an identical or different manner by fluorine, chlorine, methyl, ethyl, methoxy, ethoxy, methylthio, trifluoromethyl, trifluoromethoxy, trifluoromethylthio, dimethylamino, diethylamino, cyano, nitro, C_1 - C_4 -alkoxycarbonyl or di(C_1 - C_4)alkylaminocarbonyl, or represent $-OR^6$ or $-NR^5R^6$.

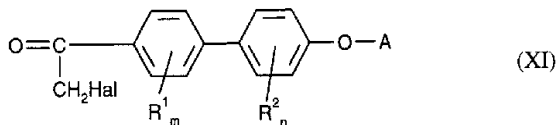
4. Compounds of the formula (X)



in which

A, R^1 , R^2 , m and n have the meaning given in Claim 1.

5. Compounds of the formula (XI)



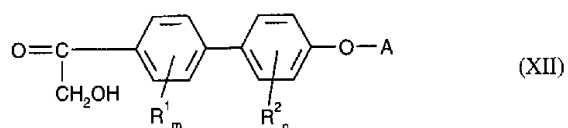
in which



A, R¹, R², m and n have the meaning given in Claim 1 and

Hal represents chlorine or bromine.

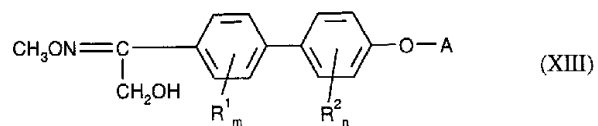
6. Compounds of the formula (XII)



5 in which

A, R¹, R², m and n have the meaning given in Claim 1.

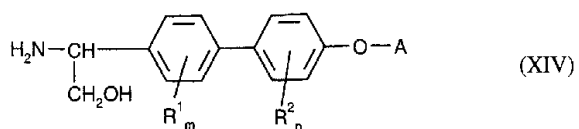
7. Compounds of the formula (XIII)



in which

10 A, R¹, R², m and n have the meaning given in Claim 1.

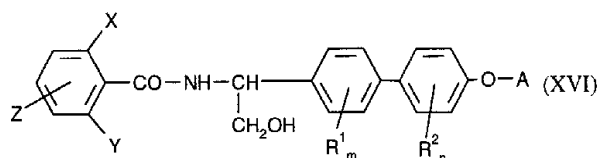
8. Compounds of the formula (XIV)



in which

A, R¹, R², m and n have the meaning given in Claim 1.

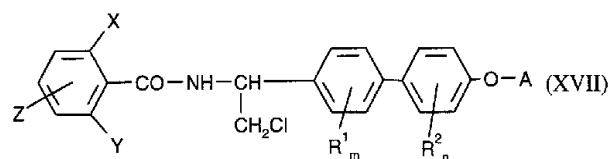
9. Compounds of the formula (XVI)



in which

A, R¹, R², m, n, X, Y and Z have the meaning given in Claim 1.

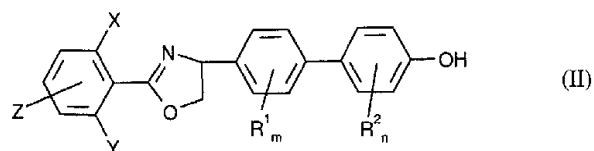
10. Compounds of the formula (XVII)



in which

A, R¹, R², m, n, X, Y and Z have the meaning given in Claim 1.

11. Process for the preparation of compounds of the formula (I) according to Claim 1, characterized in that hydroxybiphenyloxazolines of the formula (II)



in which

X, Y, Z, R¹, R², m and n have the meaning given in Claim 1,

are reacted with a compound of the formula (III)



5 in which

A has the meaning given in Claim 1 and

M represents a leaving group,

if appropriate in the presence of a base and/or a catalyst and if appropriate in the presence of a diluent.

- 10 12. Pest control composition, characterized by a content of at least one compound of the formula (I) according to Claim 1.
13. Use of compounds of the formula (I) according to Claim 1 for controlling pests.
14. Method of controlling pests, characterized in that compounds of the formula (I) according to Claim 1 are allowed to act on pests and/or their environment.
- 15 15. Process for the preparation of pest control compositions, characterized in that compounds of the formula (I) according to Claim 1 are mixed with extenders and/or surface-active agents.
16. Use of compounds of the formula (I) according to Claim 1 for the preparation of pest control compositions.



17. Compounds of formula (I) as defined in claim 1, compositions containing same, process for their preparation and use of the compounds and compositions for controlling pests substantially as hereinbefore described with reference to the Examples.

DATED this 20th day of August 1999

BAYER AKTIENGESELLSCHAFT

By its Patent Attorneys
DAVIES COLLISON CAVE

3
8
8
9

