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Compositions for controlling plant pests

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(56) Related Art

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ABSTRACT

The invention relates to agents for combating plant pests, containing the compound of formula (I) mixed with fungicidal active agents, excluding cyclopropylcarboxamide derivatives and azolylmethylcycloalkanes.

Australian Patents Act 1990 - Regulation 3.2

ORIGINAL COMPLETE SPECIFICATION STANDARD PATENT

Invention Title: Compositions for controlling plant pests

The following statement is a full description of this invention, including the best method of performing it known to me:-

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Compositions for controlling plant pests

The present invention relates to pesticides which comprise an active compound combination of the compound of the formula (I)

$$CI \longrightarrow S \longrightarrow CH_2 \longrightarrow H \longrightarrow N \longrightarrow CH_3 \longrightarrow NO_2$$
 (I)

with fungicides.

Fungicidally active compounds, such as azole derivatives, aryl benzyl ethers, benzamide, morpholine compounds and other heterocycles, are known (cf. K.H. Büchel "Pflanzenschutz und Schädlingsbekämpfung", pages 140 to 153, Georg Thieme-Verlag, Stuttgart 1977, EP published specification 0 040 345, German Offenlegungsschrift 2 324 010, German Offenlegungsschrift 2 201 063, EP published specification 0 112 284, EP published specification 0 304 758 and DD patent specification 140 412).

Mixtures of certain nitromethylene derivatives with fungicidally active compounds and their use as pesticides in crop protection are already known (US-P-4 731 385; JP published specifications 63-68507, 63/68505; 63/72 608; 63/72 609, 63/72 610, WO 96/03 045, Japanese patent specification 08 245 323, Japanese patent specification 04 368 303, Japanese patent specification 05 017 311, WO 92/21 241. WO 97/22 254). Mixtures of certain open-chain nitromethylenes and nitroguanidines with fungicides are already known (Japanese published specification 30 47 106; US-P 5 181 587).

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Mixtures of cyclopropylcarboxamides with certain nitromethylene or nitroguanidine derivatives are already known (Japanese published specification 3 271 207).

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Mixtures of, inter alia, imidacloprid and fungicidally active compounds for use in the protection of materials and against termites, but not for use against plant-damaging pests, are already known (EP published specification 0511541). Mixtures of imidacloprid and azolylmethylcycloalkanes, in particular triticonazole, are known from EP published specification 545 834.

However, hitherto it has not been known that nitroguanidine derivatives and fungicides with the exception of cyclopropylcarboxamides and triticonazole have such a mutually beneficial effect in their activity that they are outstandingly suitable as compositions for controlling plant pests, whilst being tolerated well by plants.

In a first aspect the present invention provides a synergistic composition comprising a compound of the general formula (I)

and Fluquinconazole, Triadimenol, Difenoconazole, Tebuconazole or Azoxystrobin.

In a second aspect the present invention provides a process for controlling fungi and insects, said process comprising allowing a composition of the first aspect to act on the fungi and insects, and/or their habitat.

In a third aspect the present invention provides a process for preparing a composition comprising a compound of the general formula (I) as defined in the first aspect and Fluquinconazole, Triadimenol, Difenoconazole, Tebuconazole or Azoxystrobin, wherein the compound of the general formula (I) and Fluquinconazole, Triadimenol, Difenoconazole, Tebuconazole or Azoxystrobin are mixed with extenders and/or surfactants.

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The present invention relates to compositions against plant pests which contain the compound of the formula (I)

$$CI$$
 CH_2
 H
 H
 H
 H
 CH_3
 N
 NO_2
 NO_2
 N
 NO_2

in a mixture with fungicidally active compounds, except for cyclopropylcarboxamide derivatives and azolylmethylcycloalkanes.

Examples of fungicides in the compositions according to the invention for controlling plant pests which may be mentioned are:

(1) azole derivatives of the formula

$$\begin{array}{c}
R^{2} \\
I \\
R^{1}-C-R^{3} \\
I \\
(CH_{2})_{n} \\
N \\
N
\end{array}$$
(II)

(II-1)
$$R^1 = CI - (CH_2)_2$$
, $R^2 = -C(CH_3)_3$, $R^3 = OH$, $N = 1$, (TEBUCONAZOLE)

5 (II-2)
$$R^1 = CI$$
, R^2 , $R^3 = -OCH_2CH(n-C_3H_7)O$ -, $n = 1$, (PROPICONAZOLE)

(II-3)
$$R^1 = CI$$
 O $n = 1$, R^2 , $R^3 = -OCH_2CH(CH_3)O$ -, (DIFENCONAZOLE)

(II-4)
$$R^1 = CI$$
 $R^2 = CH-CH_3$ $R^3 = -OH, n = 1,$ (CYPROCONAZOLE)

(II-5)
$$R^1 = F$$

$$R^2 = R^3 = OH, n = 1,$$
(FLUTRIAFOL)

(II-6)
$$R^3 = CI$$
, $R^4 = -(CH_2)_3CH_3$, $R^5 = OH$, $n = 1$, (HEXACONAZOLE)

(II-7)
$$R^1 = CI \longrightarrow R^2 = (CH_2)_3 CH_3, R^3 = CN, n = 1,$$
(MYCLOBUTANIL)

(II-8)
$$R^1 = CI \longrightarrow R^2 = -(CH_2)_2CH_3, R^3 = H, n = 1,$$
 (PENCONAZOLE)

(II-9)
$$R^1 = CI$$
, R^2 , $R^3 = -OCHCH_2CH_2$, $n = 1$, OCH_2CF_3

(II-10)
$$R^{1} = CI$$
, R^{2} , $R^{3} = -CCHCH_{2}C$, $n = 1$, $C_{2}H_{5}$

(ETACONAZOLE)

(II-11)
$$R^1 = CI$$
, R^2 , $R^3 = -OCH_2CHCH_2$, $n = 1$, Br (BROMUCONAZOLE)

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(II-12)
$$R^1 = CH_2$$
, $R^2 = CI$, $R^3 = OH$, $n = 1$,

(II-13)
$$R^1 = F - CI$$
, R^2 , $R^3 = CI$, $n = 1$,

(II-14)
$$R^1 =$$
 $R^2 = -CH_2-CH_2$ CI , $n = 1$, $R^3 = CN$, (FENBUCONAZOLE)

(II-15)
$$R^1 = CI$$
 $R^2 = CH_2OCF_2CHF_2$, $R^3 = H$, $n = 1$, (TETRACONAZOLE)

(II-16)
$$R^1 = CI \longrightarrow O$$
, $R^2 = -CH(OH) - C(CH_3)_3$, $n = 0$, $R^3 = H$, (TRIADIMENOL)

(II-17)
$$R^1 = CI$$
 O, $R^2 = -CO-C(CH_3)_3$, $n = 0$, $R^3 = H$, (TRIADIMEFON)

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(II-19)
$$R^1 = CI \longrightarrow CH_2$$
-, $R^2 = -CH(OH)-C(CH_3)_3$, $n = 0$, $R^3 = H$, (DICLOBUTRAZOLE)

(II-20)
$$R^1$$
 and $R^2 = CI$ $CH=$, $R^3 = -CH$ -tButyl, $n=0$, (DINICONAZOLE)

(2) azole derivatives of the formula

$$F \xrightarrow{CH_3} N = N$$

$$Si - CH_2 - N = N$$

$$F = N$$

$$(III)$$

(FLUSILAZOLE)

(3) the azole derivative of the formula

$$CI \xrightarrow{CI} O \cdot (CH_2)_2 \stackrel{N}{\downarrow} \stackrel{O}{\downarrow} N \stackrel{= N}{=} N$$

$$CI \xrightarrow{CI} O \cdot (CH_2)_2 \stackrel{N}{\downarrow} \stackrel{O}{\downarrow} N \stackrel{= N}{=} N$$

$$(IV)$$

(PROCHLORAZ)

(4) of the compound

 S_{χ} (V)

(5) the azole derivative of the formula

$$F \xrightarrow{O} N \xrightarrow{N-N} CI$$

$$VI)$$

(FLUQUINCONAZOLE)

5 (6) heterocycles of the formula

$$(VII-1)X = 0$$
, $R^8 = CH_3$, $R^9 = H$, $R^{10} = C_{10}H_{21}$

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(TRIDEMORPH)

$$(VII-2)X = 0$$
, $R^8 = CH_3$, $R^9 = H$, $R^{10} = C_9H_{19}$

(ALDIMORPH)

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(VII-3)
$$X = 0$$
, $R^8 = CH_3$, $R^9 = CH_3$, $R^{10} = CH_3$

(FENPROPIMORPH)

$$(VII-4)X = CH_2, R^8 = H, R^9 = CH_3, R^{10} = CH_3$$

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(FENPROPIDIN)

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(7) the compound of the formula

(8) the compound of the formula

$$CH_3$$
 O N OCH_3 O

(9) the compound of the formula

$$F_3C$$
 CH_2 -NH CI CI (X)

(10) the compound of the formula

$$CI \longrightarrow CH_2 \longrightarrow N$$
 OCH_3
 (XI)
 $(PYRIFENOX)$

the compound of the formula (11)

(FENARIMOL)

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(12)the compound of the formula

(TRIFLUMIZOLE)

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compounds of the formula (13)

(XIV-1)
$$R^{11} = \begin{array}{c} O \\ > O \\ > O \\ > CH_3 \\ O \\ CH = CH_2 \end{array}$$

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$$(XIV-2) R11 = -N CH3$$

(PROCYMIDONE)

(XIV-3)
$$R^{11} = -N$$
 N-CO-NH-CH(CH₃)₂ (IPRODIONE)

(14) compounds of the formula

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$$(XV-1)$$
 $R^{12} = CH_3$ (PYRIMETHANIL)

(XV-2)
$$R^{12} = C C - CH_3$$
 (MEPANIPYRIM)

(15) compounds of the formula

$$(CH_3)_2N-SO_2-N-S-CCI_2F$$

$$(XVI)$$

(XVI-1) $R^{13} = H$

(DICHLOFLUANID)

(XVI-2) $R^{13} = CH_3$

(TOLYLFLUANID)

(16) the compound of the formula

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(DITHIANON)

(17) the compound of the formula

$$H_3C-(CH_2)_{11}-NH$$
 NH_2 CH_3CO_2H $(XVIII)$

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(DODINE)

(18) the compound of the formula

$$\begin{array}{c} CI \\ CI \\ CI \\ CI \end{array} \qquad (XIX)$$

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(CHLOROTHALONIL)

(19) the compound of the formula

$$CH_3O$$
 CH_3O
 CH_3O
 CH_3O
 CH_3O
 CH_3O

(DIMETHOMORPH)

5 (20) the compound of the formula

$$CH_3$$
-O- CH_2 - C
 CH_3
 CH - CO_2CH_3
 CH_3
 CH_3

10 (21) the compound of the formula

$$H_5C_2$$
-NH-CO-NH-CO-C=N-OCH $_3$ (XXII)

(CYMOXANIL)

(METALAXYL)

(22) the compound of the formula

$$F_3C \xrightarrow{CI} O_2N \xrightarrow{CI} CF_3$$
 (XXIII)

(FLUAZINAM)

(23) the compound of the formula

$$H_3C$$
 N S O $(XXIV)$

10 (24) compounds of the formula

$$(XXV-1) R14 = -N$$

(CAPTAN)

(XXV-2)
$$R^{14} = -N$$
 (FOLPET)

(25) the compound of the formula

$$\begin{array}{c|c}
CI & CI \\
N & NH \\
\end{array}$$
CI
$$\begin{array}{c}
CI & CI \\
N & NH \\
\end{array}$$
(XXVI)

(ANILAZIN)

5 (26) the compound of the formula

$$CH_{3}$$

$$CH_{2}$$

$$CH_{3}$$

$$C=0$$

$$CH_{3}$$

$$C=0$$

$$CH_{3}$$

$$C=0$$

$$CH_{3}$$

$$C=0$$

$$CH_{3}$$

$$C=0$$

$$CH_{3}$$

(OXADIXYL)

(27) the compound of the formula

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$$\begin{bmatrix} H_5C_2O \\ H \end{bmatrix} P = \begin{bmatrix} O \\ O \end{bmatrix}_3 AI$$
 (XXVIII)

(FOSETYL AL)

(28) the compound of the formula

$$O_{2}N \xrightarrow{O-CO-CH=CH-CH_{3}} CH-C_{6}H_{13}$$

$$CH_{3}$$

$$NO_{2}$$

(DINOCAP)

(XXIX)

(29) the compound of the formula

$$CH_{2}$$
- $CO_{2}CH_{3}$
 CH_{2} - $CO_{2}N$
 CH_{3}
 $CH_$

5 (30) the compound of the formula

(31) the compound of the formula

(ANTRACOL)

(32) compounds of the formula

(XXXIII)

$$(XXXIII-1)$$
 $M = Zn$ $(ZINEB)$

$$(XXXIII-2)$$
 $M = Mn$ $(MANEB)$

(33) the compound of the formula

$$(CH3)2N - \langle \begin{array}{c} S & S \\ \\ S - S \\ \\ \end{array}) N(CH3)2$$
(XXXIV)
(THIRAM)

(34) the compound of the formula

$$CI \xrightarrow{CH_2} CI$$

$$CI \xrightarrow{CH_2} N$$

$$CH_2 - N = N$$

$$(XXXV)$$

$$(TYMIBENCONAZOLE)$$

(35) the compound of the formula

$$\begin{bmatrix} [(S-CS-NH-CH_2-CH_2-NH-CS-S-) \cdot Zn(NH_3)_3)] \\ (-S-CS-NH-CH_2-CH_2-NH-CS-S-] \end{bmatrix}_{\Upsilon}$$
 (XXXVI)

(METIRAM)

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(36) the compound of the formula

10 (37) the compound of the formula

(38) compounds of the formula

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$$R^{15}$$
 R^{16}
 R^{17}
 CH_3HNC
 $NOCH_3$
 $(XXXIX)$

in which

 R^{15} and R^{16} independently of one another each represent hydrogen, halogen, methyl or phenyl and

R¹⁷ represents hydrogen or methyl,

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(39) 8-tbutyl-2-(N-ethyl-N-n-propylamino)-methyl-1,4-dioxaspiro[4.5]decane of the formula

$$\begin{array}{c} C(CH_3)_3 \\ O \\ O \\ CH_2 - N \\ CH_2CH_2CH_3 \end{array} (XL)$$

10 (40) the compound of the formula

$$\mathsf{F} = \mathsf{CH_3} \\ \mathsf{CH_2} - \mathsf{CH_2OSO_2CH_3)_2}$$

(41) the compound of the formula

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(42) the compound of the formula

$$A = \bigcap_{CI}^{CI} fenpiclonil$$

$$A = \bigcup_{CF_3}^{F}$$

(43) the compound of the formula

tBu
$$C - CH_3 - CH_2 - N$$

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benzimidazoles of the formula (44)

$$\mathbb{R}^9$$

 $R^9 = CONHtBu; R^6 = -NHCOOMe$

benomyl

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$$R^9 = H; R^6 = -C$$

thiabendazole

$$R^9$$
 = H; R^6 = -NHCOOMe

carbendazim

(45) the compound of the formula

5 (46) the compound of the formula

$$HN = \begin{bmatrix} NH \\ I \\ (CH_2)_8 NH - C - NH_2 \end{bmatrix}$$

(47) the compound of the formula

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(48) the compound of the formula

(49) the compound of the formula

5 (50) the compound of the formula

(51) the compound of the formula

$$\sim$$

IPBC

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(52) the compound of the formula

OIT

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(53) the compound of the formula

DCOIT

(54) the compound of the formula

butylbenzisothiazolinone

(55) the compound of the formula

benzothiophene-2-cyclohexyl-

carboxamide S,S-dioxide

The active compound of the formula (I) is known from EP published specification 0 375 907.

The fungicidal active compounds are also known.

- 15 Thus, for example, the following publications describe:
 - (1) compounds of the formula (II)

DE published specification 2 201 063

DE published specification 2 324 010

DE published specification 2 737 489

DE published specification 3 018 866

DE published specification 2 551 560

EP 47 594

DE 2735872

(2)	the compound of the formula (III)
	EP 68 813
	US 4 496 551

- 5 (3) the compound of the formula (IV)

 DE published specification 2 429 523

 DE published specification 2 856 974

 US 4 108 411
- 10 (6) compounds of the formula (VII)
 DL 140 041
 - (7) the compound of the formula (VIII) EP 382 375

(8) the compound of the formula (IX) EP 515 901

- (9) the compound of the formula (X)20 EP 314 422
 - (10) the compound of the formula (XI) EP 49 854
- 25 (11) the compound of the formula (XII)

 DE published specification 1 770 288

 US 3 869 456

(13) compounds of the formula (XIV)

DE 2 207 576

US 3 903 090

US 3 755 350

5 US 3 823 240

- (14) compounds of the formula (XV) EP 270 111
- 10 (19) the compound of the formula (XX) EP 219 756
 - (34) the compound of the formula (XXXV)
 US 4 512 989

(38) compounds of the formula (XXXIX) EP 398 692

- (48) compound from WO 97/27189
 - (49) compound from

 WO 96/16048, this compound can be present in 2 tautomeric forms (A) and

 (B)

[A]
$$CI$$
 OH CH_2 CH_2

Compounds from the groups (15), (16), (17), (18), (23), (34), (25), (28), (31), (32), (33) and (38) to (47) are described, for example, in K.H. Büchel, "Pflanzenschutz und Schädlingsbekämpfung", pages 121-153, Georg Thieme-Verlag, Stuttgart, 1977. The compound of group (39) is known from EP published specification 281 842.

5 The compound of group (50) is known from WO 97/06 171.

The compound of group (51) is known from DE-24 33 410.

The compounds of groups (52) to (54) are known from W. Paulus, "Microbicides for the Protection of Materials", Chapman & Hall 1993.

The compound of group (55) is known from EP-0 512 349.

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In addition to the active compound of the formula (I), the active compound combinations according to the invention contain at least one fungicidal active compound, for example selected from the compounds of groups (1) to (55). In addition, they may also contain other active compounds and customary auxiliaries and additives and also diluents.

Preferred fungicidally active compounds in the compositions according to the invention are:

kresoxim-methyl, tebuconazole, metalaxyl, azoxystrobin, triadimenol, bitertanol, fenpicolonil, cyproconazole, propiconazole, fludioxonil and triazoxides.

The mixtures show a clear synergistic effect when the active compounds in the active compound combinations according to the invention are present in certain weight ratios. However, the weight ratios of the active compounds in the active compound combinations can be varied within a relatively wide range. In general,

0.1 to 10 parts by weight, preferably

0.3 to 3 parts by weight of at least one fungicidal active compound, for example from groups (1) to (55),

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are present per part by weight of active compound of the formula (I).

The active compound combinations according to the invention have very good fungicidal properties. They can be employed in particular for controlling phytopathogenic fungi, such as Plasmodiophoromycetes, Oomycetes, Chytridiomycetes, Zygomycetes, Ascomycetes, Basidiomycetes, Deuteromycetes, etc.

The active compound combinations according to the invention are particularly suitable for controlling cereal diseases, such as Erysiphe, Cochliobolus, Septoria, Pyrenophora and Leptosphaeria, and against fungal attack on vegetables, grapevines and fruits, for example against Venturia or Podosphaera on apples, Uncinula on grapevines or Sphaerotheca on cucumbers.

The active compound combinations are also highly suitable for controlling animal pests, preferably arthropods, in particular insects, which are encountered in agriculture, in forestry, in the protection of stored products and of materials, and in the hygiene sector. They are active against normally sensitive and resistant species and against all or some stages of development. The abovementioned pests include:

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.

From the order of the Symphyla, for example, Scutigerella immaculata.

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

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

From the order of the Orthoptera, for example, Blatta orientalis, Periplaneta americana, Leucophaea maderae, Blattella germanica, Acheta domesticus,

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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.

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.

From the order of the Homoptera, for example, Aleurodes brassicae, Bemisia tabaci,

Trialeurodes vaporariorum, Aphis gossypii, Brevicoryne brassicae, Cryptomyzus ribis, Doralis fabae, Doralis pomi, Eriosoma lanigerum, Hyalopterus arundinis, Macrosiphum avenae, Myzus spp., Phorodon humuli, Rhopalosiphum padi, Phylloxera vastatrix, Pemphigus spp., Empoasca spp., Euscelis bilobatus, Nephotettix cincticeps, Lecanium corni, Saissetia oleae, Laodelphax striatellus,

Nilaparvata lugens, Aonidiella aurantii, Aspidiotus hederae, Pseudococcus spp. and Psylla spp.

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 flammea, Prodenia litura, 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.

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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., 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., 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, Oscinella frit, Phorbia spp., Pegomyia hyoscyami, Ceratitis capitata, Dacus oleae and Tipula paludosa.

The fact that the active compound combinations are well tolerated by plants at the concentrations required for controlling plant diseases permits the treatment of aboveground parts of plants, of propagation stock and seeds, and of the soil.

The active compound combinations according to the invention can be converted into the customary formulations, such as solutions, emulsions, suspensions, powders, foams, pastes, granules, aerosols, microencapsulations in polymeric compounds and in coating compositions for seeds, and ULV formulations.

These formulations are produced in a known manner, for example by mixing the active compounds with extenders, that is liquid solvents, liquefied gases under pressure, and/or solid carriers, optionally with the use of surfactants, that is

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emulsifiers and/or dispersants, and/or foam formers. If the extender used is water, it is also possible to use, for example, organic solvents as auxiliary solvents. Essentially, suitable liquid solvents include 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 example petroleum fractions, alcohols, such as butanol or glycol and also their ethers and esters, ketones, such as acetone, methyl ethyl ketone, methyl isobutyl ketone or cyclohexanone, strongly polar solvents, such as dimethylformamide and dimethyl sulphoxide, and also water. Liquefied gaseous extenders or carriers refer to those liquids which are gaseous at normal temperature and under atmospheric pressure, for example aerosol propellants, such as halogenated hydrocarbons, and also butane, propane, nitrogen and carbon monoxide. Suitable solid carriers are: for example ground natural minerals, such as kaolins, clays, talc, chalk, quartz, attapulgite, montmorillonite or diatomaceous earth, and ground synthetic minerals, such as finely divided silica, alumina and silicates. Suitable solid carriers for granules are: for example crushed and fractionated natural rocks, such as calcite, marble, pumice, sepiolite and dolomite, or else synthetic granules of inorganic and organic meals, and granules of organic materials such as sawdust, coconut shells, maize cobs and tobacco stalks. Suitable emulsifiers and/or foam formers are: for example nonionic and anionic emulsifiers, such as polyoxyethylene fatty acid esters, polyoxyethylene fatty alcohol ethers, for example alkylaryl polyglycol ethers, alkylsulphonates, alkyl sulphates, arylsulphonates, or else protein hydrolysates. Suitable dispersants are: for example lignin-sulphite waste liquors and methylcellulose.

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Tackifiers, 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, or else natural phospholipids, such as cephalins and lecithins and synthetic phospholipids, can be used in the formulations. Other possible additives are mineral and vegetable oils.

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.

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The formulations generally comprise between 0.1 and 95 per cent by weight of active compound, preferably between 0.5 and 90%.

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In the formulations, the active compound combinations according to the invention can be present as a mixture with other known active compounds, such as fungicides, insecticides, acaricides and herbicides, and as mixtures with fertilizers or plant growth regulators.

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The active compound combinations can be employed as such, in the form of their formulations or of the use forms prepared therefrom, such as ready-to-use solutions, emulsifiable concentrates, emulsions, suspensions, wettable powders, soluble powders and granules.

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The application is carried out in a customary manner, for example by watering, spraying, atomizing, scattering, spreading, dry dressing, wet dressing, liquid dressing, slurry treatment of seeds or incrustation.

In the treatment of parts of plants, the active compound concentrations in the use forms can be varied within a relatively wide range. In general, they are between 1 and 0.0001% by weight, preferably between 0.5 and 0.001%.

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In the treatment of seed, the amounts of active compound which are generally required are from 0.001 to 50 g per kilogram of seed, preferably from 0.01 to 10 g. In the treatment of the soil, active compound concentrations of from 0.00001 to 0.1% by weight, preferably from 0.0001 to 0.02% by weight, are required at the site of action.

Furthermore, it has been found that the active compound combinations according to the invention have a potent insecticidal action against insects which destroy industrial materials.

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The following insects may be mentioned by way of example and as being preferred, but without any limitation:

Beetles, such as

Hylotrupes bajulus, Chlorophorus pilosis, Anobium punctatum, Xestobium rufovillosum, Ptilinus pecticornis, Dendrobium pertinex, Ernobius mollis, Priobium carpini, Lyctus brunneus, Lyctus africanus, Lyctus planicollis, Lyctus linearis, Lyctus pubescens, Trogoxylon aequale, Minthes rugicollis, Xyleborus spec., Tryptodendron spec., Apate monachus, Bostrychus capucins, Heterobostrychus brunneus, Sinoxylon spec., Dinoderus minutus.

Dermapterans, such as

Sirex juvencus, Urocerus gigas, Urocerus gigas taignus, Urocerus augur.

20 Termites, such as

Kalotermes flavicollis, Cryptotermes brevis, Heterotermes indicola, Reticulitermes flavipes, Reticulitermes santonensis, Reticulitermes lucifugus, Mastotermes darwiniensis, Zootermopsis nevadensis, Coptotermes formosanus.

25 Bristletails, such as Lepisma saccharina.

Industrial materials are to be understood as meaning, in the present context, non-live materials such as, preferably, plastics, glues, sizes, paper and board, leather, wood and timber products, and coatings.

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The materials to be protected against attack by insects are very particularly wood and timber products.

Wood and timber products which can be protected by the composition according to the invention or mixtures comprising such a composition are to be understood as meaning, for example, construction timber, wooden beams, railway sleepers, bridge components, jetties, wooden vehicles, boxes, pallets, containers, telephone poles, wood panelling, windows and doors made of wood, plywood, particle board, joiner's articles, or wood products which, quite generally, are used in the construction of houses or in joinery.

The active compound combinations can be used as such, in the form of concentrates or generally customary formulations, such as powders, granules, solutions, suspensions, emulsions or pastes.

The formulations mentioned can be prepared in a manner known per se, for example by mixing the active compounds with at least one solvent or diluent, emulsifier, dispersant and/or binder or fixative, water repellent, if appropriate desiccants and UV

stabilizers and, if appropriate, dyes and pigments and other processing auxiliaries.

The insecticidal compositions or concentrates used for the protection of wood and wooden materials comprise the active compound according to the invention at a concentration of 0.0001 to 95% by weight, in particular 0.001 to 60% by weight.

The amount of the compositions or concentrates employed depends on the species and the occurrence of the insects and on the medium. The optimum rate of application can be determined upon use in each case by test series. However, in general, it suffices to employ 0.0001 to 20% by weight, preferably 0.001 to 10% by weight, of the active compound, based on the material to be protected.

The solvent and/or diluent used is an organochemical solvent or solvent mixture and/or an oily or oil-type organochemical solvent or solvent mixture of low volatility and/or a polar organochemical solvent or solvent mixture and/or water and, if appropriate, an emulsifier and/or wetting agent.

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Organochemical solvents which are preferably employed are oily or oil-type solvents having an evaporation number of above 35 and a flash point of above 30°C, preferably above 45°C. Substances which are used as such oily and oil-type solvents which have low volatility and are insoluble in water are suitable mineral oils or their aromatic fractions, or mineral-oil-containing solvent mixtures, preferably white spirit, petroleum and/or alkylbenzene.

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Substances which are advantageously used are mineral oils with a boiling range of 170 to 220°C, white spirit with a boiling range of 170 to 220°C, spindle oil with a boiling range of 250 to 350°C, petroleum or aromatics of the boiling range 160 to 280°C, essence of turpentine and the like.

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In a preferred embodiment, liquid aliphatic hydrocarbons with a boiling range of 180 to 210° C or high-boiling mixtures of aromatic and aliphatic hydrocarbons with a boiling range of 180 to 220° C and/or spindle oil and/or monochloronaphthalene, preferably α -monochloronaphthalene, are used.

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The organic oily or oil-type solvents of low volatility having an evaporation number of above 35 and a flash point of above 30°C, preferably above 45°C, can be partially replaced by organochemical solvents of high or medium volatility, with the proviso that the solvent mixture also has an evaporation number of above 35 and a flash point of above 30°C, preferably above 45°C, and that the insecticide/fungicide mixture is soluble or emulsifiable in this solvent mixture.

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In a preferred embodiment, part of the organochemical solvent or solvent mixture is replaced by an aliphatic polar organochemical solvent or solvent mixture. Substances

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which are preferably used are aliphatic organochemical solvents having hydroxyl and/or ester and/or ether groups, such as, for example, glycol ethers, esters or the like.

The organochemical binders used within the scope of the present invention are the synthetic resins and/or binding drying oils which are known per se and can be diluted with water and/or are soluble or dispersible or emulsifiable in the organochemical solvents employed, in particular binders composed of, or comprising, an acrylate resin, a vinyl resin, for example polyvinyl acetate, polyester resin, polycondensation or polyaddition, resin, polyurethane resin, alkyd resin or modified alkyd resin, phenolic resin, hydrocarbon resin, such as indene-coumarone resin, silicone resin, drying vegetable oils and/or drying oils and/or physically drying binders based on a natural and/or synthetic resin.

The synthetic resin used as the binder can be employed in the form of an emulsion, dispersion or solution. Up to 10% by weight of bitumen or bituminous substances can also be used as binders. In addition, dyes, pigments, water repellents, odourmasking substances and inhibitors or anticorrosives known per se and the like can also be employed.

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The composition or the concentrate preferably comprises, in accordance with the invention, at least one alkyd resin or modified alkyd resin and/or a drying vegetable oil as the organochemical binders. Preferably used according to the invention are alkyd resins with an oil content of over 45% by weight, preferably 50 to 68% by weight.

All or some of the abovementioned binder can be replaced by a fixative (mixture) or a plasticizer (mixture). These additives are intended to prevent volatilization of the active compounds and crystallization or precipitation. They preferably replace 0.01 to 30% of the binder (based on 100% of binder employed).

The plasticizers are from the chemical classes of the phthalic esters, such as dibutyl phthalate, dioctyl phthalate or benzyl butyl phthalate, the phosphoric esters, such as tributyl phosphate, the adipic esters, such as di-(2-ethylhexyl) adipate, the stearates, such as butyl stearate or amyl stearate, the oleates, such as butyl oleate, the glycerol ethers or relatively high-molecular-weight glycol ethers, glycerol esters and p-toluene-sulphonic esters.

Fixatives are chemically based on polyvinyl alkyl ethers, such as, for example, polyvinyl methyl ether, or ketones, such as benzophenone or ethylenebenzophenone.

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Also particularly suitable as a solvent or diluent is water, if appropriate as a mixture with one or more of the abovementioned organochemical solvents or diluents, emulsifiers and dispersants.

Particularly effective protection of wood is achieved by large-scale industrial impregnation processes, for example vacuum, double-vacuum or pressure processes.

If appropriate, the ready-to-use compositions can additionally comprise other insecticides.

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Suitable additional components which may be admixed are, preferably, the insecticides mentioned in WO 94/29 268. The compounds mentioned in that document are expressly incorporated into the present application by reference.

- Very particularly preferred components which may be admixed are insecticides, such as chlorpyriphos, phoxim, silafluofen, alphamethrin, cyfluthrin, cypermethrin, deltamethrin, permethrin, imidacloprid, NI-25, flufenoxuron, hexaflumuron and triflumuron.
- The good pesticidal activity of the active compound combinations according to the invention is demonstrated by the examples below. Whereas the individual active

compounds or the known active compound combinations have weaknesses in the pesticidal activity, the tables of the examples below show unambiguously that the activity which was found of the active compound combinations according to the invention is greater than the sum of the activities of the individual active compounds and also greater than the activities of the known active compound combinations.

In the examples below, the active compound of the formula (I)

$$CI \longrightarrow S \longrightarrow CH_2 \longrightarrow H \longrightarrow H \longrightarrow CH_3 \longrightarrow NO_2$$
 (I)

10 is used.

The fungicidally active compounds which are also used are given in the examples.

Example A

Phaedon larvae test

5 Solvent:

7 parts by weight of dimethylformamide

Emulsifier:

1 part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, 1 part by weight of an active compound or an active compound combination is mixed with the stated amounts of solvent and 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 active compound of the desired concentration and are populated with larvae of the mustard beetle (Phaedon cochleariae) while the leaves are still moist.

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

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In this test, for example, the following active compound combinations according to the present invention show a synergistically increased activity compared to the individually applied active compounds:

<u>Table A</u>

(plant-damaging insects)

Phaedon larvae test

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Active compound/active com-	Active compound/active com-	Kill in %
pound mixture	pound mixture concentration in %	
CI S HN H-CH3		
N	0.00016	20
NO ₂	0.000032	0
according to formula (I)		
CH ₃ O O N O O O O O O O O O O O O O O O O	0.1	0
cresoxim-methyl		
ormula (I) + kresoxim-methyl	0.00016 + 0.1	75

Example B

Plutella test

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Solvent:

7 parts by weight of dimethylformamide

Emulsifier:

1 part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of an active compound or an active compound combination is mixed with the stated amounts of solvent and 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 active compound of the desired concentration and are populated with caterpillars of the diamondback moth (Plutella xylostella) while the leaves are still moist.

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

In this test, for example, the following active compound combinations according to the present invention show a synergistically increased activity compared to the individually applied active compounds:

Table B (plant-damaging insects)

Plutella test

Active compound/active com-	Active compound/active com-	Kill in %
pound mixture	pound mixture concentration in %	
CI S HN H T		
HN N-CH ₃	0.0008	0
according to formula (I)		
CI	0.02	0
OH C(CH ₃) ₃		
		·
tebuconazole		
CH ₃ O CH ₃ O CH ₃ O CH ₃	0.1	0
metalaxyl		
formula (I) + tebuconazole	0.0008 + 0.02	100
formula (II) + metalaxyl	0.0008 + 0.1	100

Example C

Heliothis virescens test

5 Solvent:

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7 parts by weight of dimethylformamide

Emulsifier:

I part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of an active compound or an active compound combination is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration.

Soya bean shoots (Glycine max) are treated by being dipped into the active compound preparation of the desired concentration and are populated with Heliothis virescens caterpillars while the leaves are still moist.

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

In this test, for example, the following active compound combinations according to the present invention show a synergistically enhanced activity in comparison to the individually applied active compounds:

Table C

(plant-damaging insects)

Heliothis virescens test

Active compound/active com-	Active compound/active	Kill in %
pound mixture	compound mixture	
	concentration in %	
CI S HN H CH ₃ N NO ₂	0.00016	50
according to formula (I)		
CN H ₃ CO OCH ₃	0.0008	0
azoxystrobin		
formula (I) + azoxystrobin	0.00016 + 0.0008	100

Example D

Nephotettix test

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Solvent:

7 parts by weight of dimethylformamide

Emulsifier:

1 part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of an active compound or an active compound combination is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration.

Rice seedlings (Oryza sativa) are treated by being dipped into the active compound preparation of the desired concentration and are populated with the green rice leaf hopper (Nephotettix cincticeps) while the leaves are still moist.

After the desired period of time, the kill in % is determined. 100% means that all the leaf hoppers have been killed; 0% means that none of the leaf hoppers have been killed.

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In this test, for example, the following active compound combinations according to the present invention show a synergistically enhanced activity in comparison to the individually applied active compounds:

Table D

(plant-damaging insects)

Nephotettix test

Active compound/active com-	Active compound/active	Kill in %
pound mixture	compound mixture	
	concentration in %	
CI S HN H-CH ₃	0.00000128	0
according to formula (I)		
CH ₃ O CH ₃ O CH ₃ H ₃ C O CH ₃	0.1	0
metalaxyl		
formula (I) + metalaxyl	0.00000128 + 0.1	85

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Example E

Critical concentration test/soil insects

5 Test insect:

Spodoptera frugiperda

Solvent:

4 parts by weight of acetone

Emulsifier:

1 part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of the active compound or the active compound mixture is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration.

The active compound preparation is mixed intimately with soil. Here, the concentration of the active compound in the preparation is almost irrelevant; only the amount by weight of active compound or active compound mixture per volume unit of soil, which is stated in ppm (mg/l), matters.

Soil is filled into 0.5 l pots and these are allowed to stand at 20°C. Immediately after preparation, 3 pre-germinated maize coms are placed into each pot. After the maize coms have emerged, boring sleeves are placed onto the pots. 9 days after preparation, the maize is populated with the test insects. After a further 5 days, the kill in % is determined. 100% means that all test insects have been killed; 0% means that the number of insects which are still alive is the same as for the untreated control.

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In this test, for example, the following active compound combinations according to the present invention show a synergistically enhanced activity compared to the individually applied active compounds:

<u>Table E 1/4</u> Spodoptera frugiperda test

Active compound/active com-	Active compound/active compound	Kill in %
pound mixture	mixture concentration in ppm	
CI S HN H CH ₃	1.25 0.60 0.30	100 98 50
according to formula (I)		
OH C(CH ₃) ₃	20.00	0
tebuconazole (H ₃ C) ₃ C		
HO N CI	20.00	0
triadimenol		
(H ₃ C) ₃ C HO N 2	20.00	0
bitertanol		

Table E 2/4 (continued)

Spodoptera frugiperda test

Active compound/active com-	Active compound/active compound	Kill in %
pound mixture	mixture concentration in ppm	
CI	20.00	0
fenpiclonil		
CN H ₃ CO OCH ₃	20.00	0
azoxystrobin		
CH ₃ O CH ₃ O CH ₃ H ₃ C O CH ₃	20.00	0
metalaxyl		
CH ₃ OH CI	20.00	0
cyproconazole		

<u>Table E 3/4</u> (continued) Spodoptera frugiperda test

Active compound/active com-	Active compound/active com-	Kill in %
pound mixture	pound mixture concentration	
	in ppm	
CI	20.00	0
N N N		
propiconazole		
formula (I) + tebuconazole	1.25 + 20.00	100
	0.60 + 20.00	80
	0.30 + 20.00	80
formula (I) + triadimenol	1.25 + 20.00	100
	0.60 + 20.00	80
	0.30 + 20.00	50
formula (I) + bitertanol	1.25 + 20.00	100
	0.60 + 20.00	80
	0.30 + 20.00	80
formula (1) + fenpiclonil	1.25 + 20.00	100
	0.60 + 20.00	90
	0.30 + 20.00	80
formula (I) + azoxystrobin	1.25 + 20.00	100
	0.60 + 20.00	80
	0.30 + 20.00	o
formula (I) + metalaxyl	1.25 + 20.00	100
	0.60 + 20.00	95
	0.30 + 20.00	0

Table E 4/4 (continued)

Spodoptera frugiperda test

Active compound/active compound mixture	Active compound/active compound mixture concentration in ppm	Kill in %
formula (I) + cyproconazol	1.25 + 20.00 0.60 + 20.00 0.30 + 20.00	100 50 0
formula (I) + propiconazole	1.25 + 20.00 0.60 + 20.00 0.30 + 20.00	100 90 70

Example F

Critical concentration test/root-systemic action

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Test insect:

Phaedon cochleariae larvae

Solvent:

4 parts by weight of acetone

Emulsifier:

I part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of the active compound or the active compound mixture is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration.

The active compound preparation is mixed intimately with soil. Here, the concentration of the active compound in the preparation is almost irrelevant; only the amount by weight of active compound or active compound mixture per volume unit of soil, which is stated in ppm (mg/l), matters.

Soil is filled into 250 ml pots and these are planted with cabbage (Brassica oleracea).

The active compound or the active compound combination can thus be taken up by the plant roots from the soil and transported into the leaves.

After 7 days, the leaves are populated with the abovementioned test animals. After a further 3 days, the kill in % is determined. 100% means that all test insects have been killed; 0% means that the number of insects which are still alive is the same as in the untreated control.

In this test, for example, the following active compound combinations according to the present invention show a synergistically enhanced activity compared to the individually applied active compounds:

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<u>Table F 1/3</u> Phaedon cochleariae test

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Active compound/active com-	Active compound/active compound	Kill in %
pound mixture	mixture concentration in ppm	
		1
CI S H	1.25	100
HN N-CH ₃	0.60	80
	0.30	50
NO ₂		
according to formula (I)		
ÇI		
ОН	20.00	0
C(CH ₃) ₃		
\sim \uparrow		
Ň		
\\		
tebuconazole (H ₃ C) ₃ C		
(1.130730)		
но		
	20.00	0
N CI		
\ <u>\</u> /		
triadimenol		
(H ₃ C) ₃ C		
HO		
	30.00	_
N N	20.00	0
\\		
~		
-:41		
oitertanol		

Table F 2/3 (continued)

Phaedon cochleariae test

Active compound/active com-	Active compound/active compound	Kill in %
pound mixture	mixture concentration in ppm	
CI	20.00	0
fenpiclonil	·	
CN H ₃ CO OCH ₃	20.00	0
azoxystrobin		
CH ₃ O CH ₃ O CH ₃ O CH ₃ O CH ₃	20.00	0
metalaxyl		
CI	20.00	0
propiconazole		

Table F 3/3 (continued)

Phaedon cochleariae test

Active compound/active com-	Active compound/active compound	Kill in %
pound mixture	mixture concentration in ppm	
formula (I) + tebuconazole	1.25 + 20.00	100
	0.60 + 20.00	80
	0.30 + 20.00	80
formula (I) + triadimenol	1.25 + 20.00	100
	0.60 + 20.00	100
	0.30 + 20.00	100
formula (I) + bitertanol	1.25 + 20.00	100
	0.60 + 20.00	100
	0.30 + 20.00	70
formula (I) + fenpiclonil	1.25 + 20.00	100
	0.60 + 20.00	100
	0.30 + 20.00	70
formula (I) + azoxystrobin	1.25 + 20.00	100
i	0.60 + 20.00	90
	0.30 + 20.00	50
formula (I) + metalaxyl	1.25 + 20.00	100
	0.60 + 20.00	100
	0.30 + 20.00	50
formula (I) + propiconazole	1.25 + 20.00	100
	0.60 + 20.00	100
	0.30 + 20.00	90

Example G

Formula for the calculation of the efficacy of a combination of two active compounds

The expected activity for a given combination of two active compounds can be calculated as follows (cf. Colby, S.R., "Calculating Synergistic and Antagonistic Responses of Herbicide Combinations", Weeds 15, pages 20-22, 1967):

if

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- X is the efficacy, expressed in % of the untreated control, when applying the active compound A at an application rate of m g/ha,
- is the efficacy, expressed in % of the untreated control, when applying the active compound B at an application rate of n g/ha,
 - E is the efficacy, expressed in % of the untreated control, when applying the active compounds A and B at application rates of m and n g/ha,

then
$$E = X + Y - \frac{X \times Y}{100}$$
.

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If the actual fungicidal activity exceeds the calculated value, then the activity of the combination is superadditive, i.e. a synergistic effect exists. In this case, the efficacy which was actually observed must be greater than the value for the expected efficacy (E) calculated from the formula set out above.

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Phytophthora test (tomato) / protective

Solvent:

47 parts by weight of acetone

Emulsifier:

3 parts by weight of alkylaryl polyglycol ether

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To produce a suitable preparation of active compound, I part by weight of the active compound or the active compound mixture is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration, or a commercial formulation of active compound or active compound combination is diluted with water to the desired concentration.

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To test for protective activity, young plants are sprayed with the commercial active compound preparation at the stated application rate. After the spray coating has dried on, the plants are inoculated with an aqueous spore suspension of *Phytophthora infestans*. The plants are then placed in an incubation cabin at approximately 20°C and 100% relative atmospheric humidity.

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Evaluation is carried out 3 days after the inoculation. 0% means an efficacy which corresponds to that of the control, whereas an efficacy of 100% means that no infection is observed.

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The table below shows unambiguously that the activity which was found of the active compound combinations according to the invention is greater than the calculated efficacy (see above), i.e., a synergistic effect is present.

Table G/1
Phytophthora test (tomato) / protective

Active compound/active com-	Active compound/active	Efficacy in %
pound mixture	compound mixture in g/ha	calculated/found
O ₂ N NH	500	51
H ₃ C		
according to formula (I)		
CI	500	5
propiconazole		
formula (I) + propiconazole	500 + 500	53/69

Table G/2
Phytophthora test (tomato) / protective

Active compound/active compound mixture	Active compound/active compound mixture in g/ha	Efficacy in % calculated/found
O ₂ N NH	500	55
H ₃ C according to formula (I) (H ₃ C) ₃ C		
HO N CI	500	0
triadimenol		
formula (I) + triadimenol	500 + 500	55/63

Table G/3
Phytophthora test (tomato) / protective

Active compound/active	Active compound/active	Efficacy in %
compound mixture	compound mixture in g/ha	calculated/found
O ₂ N NH H ₃ C NH	50	19
according to formula (I)		
CH ₃ O CH ₃ O CH ₃ O CH ₃	50	42
metalaxyl	·	
formula (I) + metalaxyl	50 + 50	53/68

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Example H

Sphaerotheca test (cucumber) / protective

5 Solvent:

47 parts by weight of acetone

Emulsifier:

3 parts by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of the active compound or the active compound mixture is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration, or a commercial formulation of active compound or active compound combination is diluted with water to the desired concentration.

To test for protective activity, young plants are sprayed with the commercial active compound preparation at the stated application rate. After the spray coating has dried on, the plants are inoculated with an aqueous spore suspension of *Sphaerotheca fuliginea*. The plants are then placed in an incubation cabin at approximately 20°C and 100% relative atmospheric humidity.

Evaluation is carried out 10 days after the inoculation. 0% means an efficacy which corresponds to that of the control, whereas an efficacy of 100% means that no infection is observed.

The table below shows unambiguously that the activity which was found of the active compound combinations according to the invention is greater than the calculated efficacy (see above), i.e., a synergistic effect is present.

Table H/1
Sphaerotheca test (cucumber) / protective

Active compound/active com-	Active compound/active	Efficacy in %
pound mixture	compound mixture in g/ha	calculated/found
S N	10	0
O₂N NH		1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
H₃C´		
according to formula (I)		
CH₃ OH	10	70
N CI		
cyproconazole		
formula (I) + cyproconazole	10 + 10	70 / 80

Table H/2
Sphaerotheca test (cucumber) / protective

Active compound/active compound mixture	Active compound/active compound mixture in g/ha	Efficacy in % calculated/found
CI	500	0
O ₂ N NH NH	300	
according to formula (I)		
CI	500	50
CN		
fenpiclonil		
formula (I) + fenpiclonil	500+ 500	50/83

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Example I

Botrytis test (bean) / protective

5 Solvent:

47 parts by weight of acetone

Emulsifier:

3 parts by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of the active compound or the active compound mixture is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration, or a commercial formulation of active compound or active compound combination is diluted with water to the desired concentration.

To test for protective activity, young plants are sprayed with the active compound preparation at the stated application rate. After the spray coating has dried on, 2 small pieces of agar colonized by *Botrytis cinerea* are placed onto each leaf. The inoculated plants are placed in a dark chamber at approximately 20°C and 100% relative atmospheric humidity.

2 days after the inoculation, the size of the disease spots on the leaves is evaluated.

0% means an efficacy which corresponds to that of the control, whereas an efficacy of 100% means that no infection is observed.

The table below shows unambiguously that the activity which was found of the active compound combinations according to the invention is greater than the calculated efficacy (see above), i.e., a synergistic effect is present.

Table I/1

Botrytris test (bean) / protective

Active compound/active	Active compound/active	Efficacy in %
compound mixture	compound mixture in g/ha	calculated/found
O ₂ N NH	10	0
according to formula (I)	-	
CN CN	10	40
N H fludioxonil		
formula (I) + fludioxonil	10 + 10	40 / 83

Table I/2
Botrytris test (bean) / protective

Active compound/active compound mixture	Active compound/active compound mixture in g/ha	Efficacy in % calculated/found
S N	100	4
O ₂ N NH H ₃ C		
according to formula (I)	4 1	
N C(CH ₃) ₃	100	80
tebuconazole		
formula (I) + tebuconazole	100 + 100	81 / 94

Example J

Fusarium culmorum test (wheat) / seed treatment

The active compounds are applied as a dry dressing. They are prepared by extending the active compound in question with ground minerals to give a finely pulverulent mixture which ensures even distribution on the surface of the seeds.

For the dressing, the infected seeds are shaken with the dressing in a closed glass bottle for 3 minutes.

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 2×100 seeds of the wheat are sown at a depth of 1 cm into standard soil, and the wheat is cultivated in a greenhouse at a temperature of approximately 18°C and at a relative atmospheric humidity of approximately 95% in seed containers receiving 15 hours of light per day.

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The evaluation of the plants for symptoms is carried out approximately 3 weeks after sowding. 0% means an efficacy which corresponds to that of the control, whereas an efficacy of 100% means that no infection is observed.

Table J

Fusarium culmorum test (wheat) / seed treatment

Active compound/active	Active compound/active	Efficacy in %
compound mixture	compound mixture in g/ha	
O ₂ N NH	75	0
according to formula (I)		
O-CF ₂ CN H fludioxonil	75	0
(H ₃ C) ₃ C HO N CI triadimenol	7 5	13.5
formula (I) + fludioxonil	37.5 + 37.5	38.5
formula (I) + triadimenol	37.5 + 37.5	30.5

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Example K

Puccinia test (wheat) / protective

5 Solvent:

25 parts by weight of N,N-dimethylacetamide

Emulsifier:

0.6 parts by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of active compound or active compound combination is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration, or a commercial formulation of active compound or active compound combination is diluted with water to the desired concentration.

To test for protective activity, young plants are sprayed with the active compound preparation at the stated application rate. After the spray coating has dried on, the plants are sprayed with a conidia suspension of *Puccinia recondita*. The plants remain in an incubation cabin at 20°C and 100% relative atmospheric humidity for 48 hours.

The plants are then placed in a greenhouse at a temperature of approximately 20°C and a relative atmospheric humidity of 80% to promote the development of rust pustules.

Evaluation is carried out 10 days after the inoculation. 0% means an efficacy which corresponds to that of the control, whereas an efficacy of 100% means that no infection is observed.

Table K Puccinia (wheat)/ protective

Active compound/active com-	Active compound/active com-	Efficacy in %
pound mixture	pound mixture in g/ha	
O ₂ N NH H ₃ C according to formula (I)	18.75	0
(H ₃ C) ₃ C		
HO N CI	18.75	13
triadimenol		
formula (I) + triadimenol	9.375 + 9.375	63

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Example L

Pyrenophora teres test (barley) / protective

5 Solvent:

25 parts by weight of N,N-dimethylacetamide

Emulsifier:

0.6 parts by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, 1 part by weight of active compound or active compound combination is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration, or a commercial formulation of active compound or active compound combination is diluted with water to the desired concentration.

To test for protective activity, young plants are sprayed with the active compound preparation at the stated application rate. After the spray coating has dried on, the plants are sprayed with a conidia suspension of *Pyrenophora teres*. The plants remain in an incubation cabin at 20°C and 100% relative atmospheric humidity for 48 hours.

The plants are then placed in a greenhouse at a temperature of approximately 20°C and a relative atmospheric humidity of approximately 80%.

Evaluation is carried out 7 days after the inoculation. 0% means an efficacy which corresponds to that of the control, whereas an efficacy of 100% means that no infection is observed.

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 $\underline{ \textbf{Table L}}$ Pyrenophora teres test (barley) / protective

Active compound/active com-	Active compound/active com-	Efficacy in %
pound mixture	pound mixture in g/ha	calculated/found
CI	pound mixture in gina	carculated/found
	62.5	0
ş	18.75	20
	16.75	20
O ₂ N N		
H ₃ C NH		
according to formula (I)		
CI	62.5	40
N=NO N=NO		
triazoxides		
CN H ₃ CO OCH ₃	18.75	0
ll l		
azox ystrobin		
formula (I) + azoxystrobin	9.375 + 9.375	60
formula (I) + triazoxides	31.25 + 31.25	70

Example M

Erysiphe test (barley) / protective

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Solvent:

parts by weight of N,N-dimethylacetamide

Emulsifier:

0.6 parts by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of active compound or active compound combination is mixed with the stated amounts of solvent and emulsifier, and the concentrate is diluted with water to the desired concentration, or a commercial formulation of active compound or active compound combination is diluted with water to the desired concentration.

To test for protective activity, young plants are sprayed with the preparation of active 15 compound at the stated application rate.

After the spray coating has dried on, the plants are dusted with spores of Erysiphe graminis f.sp. hordei.

20 The plants are placed in a greenhouse at a temperature of approximately 20°C and a relative atmospheric humidity of approximately 80% to promote the development of mildew pustules.

Evaluation is carried out 7 days after the inoculation. 0% means an efficacy which 25 corresponds to that of the control, whereas an efficacy of 100% means that no infection is observed.

<u>Table M</u>
Erysiphe test (barley) / protective

Active compound/active	Active compound/active	Efficacy in %
compound mixture	compound mixture in g/ha	calculated/found
O ₂ N NH NH according to formula (I)	62.5	0
CH ₃ OCH ₃ OCH ₃ kresoxim-methyl	62.5	0.
formula (I) + kresoxim-methyl	31.25 + 31.25	75

Example N

Critical concentration test/root systemic action

5 Test insect:

Spodoptera frugiperda

Solvent:

4 parts by weight of acetone

Emulsifier:

I part by weight of alkylaryl polyglycol ether

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

The preparation of active compound is mixed intimately with soil. Here, the concentration of the active compound in the preparation is virtually immaterial; only the amount by weight of active compound per volume unit of soil, which is stated in ppm (mg/l), matters. The soil is filled into 250 ml pots and these are allowed to stand at 20°C.

Immediately after preparation, 3 maize coms are placed into each pot. 14 days after the preparation, the test insects are placed into an insect bracket and attached to the plant. 4 days after the infection, the efficacy of the active compound is determined in % by counting the dead and live test insects. The efficacy is 100% if all test insects have been killed; it is 0% if the same number of test insects is alive as in the untreated control.

<u>Table N</u>

Spodoptera frugiperda test

Active compound/active compound	Kill in % at active compound/active
mixture	compound mixture concentration in ppm
O ₂ N NH H ₃ C NH	0.15 ppm = 50%
according to formula (I)	
CI OH S H	5.00 ppm = 0%
compound of group (49) compound according to formula (I) + compound of group (49)	0.15 ppm + 5.00 ppm = 80%

Example O

Critical concentration test/soil insects

Test insect:

Diabrotica balteata - larvae in the soil

5 Solvent:

4 parts by weight of acetone

Emulsifier:

1 part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, I part by weight of active compound is mixed with the stated amount of solvent, the stated amount of emulsifier is added and the concentrate is diluted with water to the desired concentration. Here, the concentration of the active compound in the preparation is almost irrelevant; only the amount by weight of active compound per volume unit of soil, which is stated in ppm (mg/l), matters. The soil is filled into 0.5 l pots and these are allowed to stand at 20°C.

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Immediately after preparation, 5 maize corns are placed into each pot. After 3 days, the test insects are placed into the treated soil. After a further 7 days, the efficacy is determined. The efficacy is calculated from the number of maize plants which have emerged.

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Active compounds, application rates and results are shown in the table below:

Table O

Soil insecticides

Diabrotica balteata - larvae in the soil

Active compound	Kill in % at active compound concen-
(constitution)	trations in ppm
O ₂ N NH S NH	0.15 ppm = 0%
according to formula (I)	
FON N CI N SO ₂ CH ₃	20.00 ppm = 70%
according to (50)	
formula (I)	0.15 ppm
+ compound according to (50)	+20.00 ppm = 70 %

Table P

Soil insecticides

Diabrotica balteata - larvae in the soil

Active compound	Kill in % at active compound concen-
(constitution)	trations in ppm
O ₂ N NH H ₃ C NH formula (I)	0.30 ppm = 50%
F O-CH ₃	20.00 ppm = 0%
according to (48)	
formula (I)	0.15 ppm
+ compound according to (48)	+20.00 ppm = 90 %

Throughout this specification and the claims which follow, unless the context requires otherwise, the word "comprise", and variations such as "comprises" or "comprising", will be understood to imply the inclusion of a stated integer or step or group of integers or steps but not the exclusion of any other integer or step or group of integers or steps.

The reference in this specification to any prior publication (or information derived from it), or to any matter which is known, is not, and should not be taken as an acknowledgment or admission or any form of suggestion that that prior publication (or information derived from it) or known matter forms part of the common general knowledge in the field of endeavour to which this specification relates.

The claims defining the invention are as follows:

1. A synergistic composition comprising a compound of the general formula (I)

and Fluquinconazole, Triadimenol, Difenoconazole, Tebuconazole or Azoxystrobin.

- 2. The composition according to Claim 1, which comprises from 0.1 to 10 parts by weight of Fluquinconazole, Triadimenol, Difenoconazole, Tebuconazole or Azoxystrobin per part by weight of the compound of the general formula (I).
- 3. A process for controlling fungi and insects, said process comprising allowing a composition of Claim 1 or Claim 2 to act on the fungi and insects, and/or their habitat.
 - 4. Use of a composition according to Claim 1 or Claim 2 for controlling fungi and insects.
- 5. A process for preparing a composition comprising a compound of the general formula (I) as defined in Claim 1 and Fluquinconazole, Triadimenol, Difenoconazole, Tebuconazole or Azoxystrobin, wherein the compound of the general formula (I) and Fluquinconazole, Triadimenol, Difenoconazole, Tebuconazole or Azoxystrobin are mixed with extenders and/or surfactants.