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(54) **NOVEL IMINO DERIVATIVE, PROCESS FOR PRODUCTION OF THE SAME, AND PESTICIDE COMPRISING THE SAME**

NEUES IMINODERIVAT, VERFAHREN ZU SEINER HERSTELLUNG UND PESTIZID DAMIT
NOUVEAU DÉRIVÉ IMINO, PROCÉDÉ DE FABRICATION DE CELUI-CI ET PESTICIDE LE
COMPRENANT

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• **KEVIN A. FORD ET AL: "Chloropyridinyl
Neonicotinoid Insecticides: Diverse Molecular
Substituents Contribute to Facile Metabolism in
Mice", CHEMICAL RESEARCH IN TOXICOLOGY,
vol. 19, no. 7, 1 July 2006 (2006-07-01), pages
944-951, XP55003365, ISSN: 0893-228X, DOI:
10.1021/tx0600696**

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Description

[Technical Field]

5 **[0001]** The present invention relates to a novel imino derivative, a method for producing the same and a use thereof. More specifically, the invention relates to a novel imino derivative, a method for producing the derivative, and an insecticide containing the derivative as an active ingredient.

[Background Art]

10 **[0002]** An insecticide employed in the field of agriculture has conventionally been required to have a diversity of the characteristics. The characteristics required for an insecticide may for example be sustained effects and broad spectrum, safety when handling, easiness in using in combination with other drugs or formulation auxiliaries. Being less expensive is also required as a matter of course.

15 **[0003]** In conjunction with the invention, a synthesis of a carbonyliminothiazolidine as a conventional insecticide compound having an imino structure was reported (see Patent Literature 1). Also in Patent Literature 2, an insecticide compound having a carbonyliminoimidazolidine structure was reported (see Patent Literature 2).

[Citation List]

20 [Patent Literature]

[0004]

25 [PTL 1] Japanese Unexamined Patent Application Publication No. 63-150275
[PTL 2] Japanese Translation of PCT No. 2007-506674

[Summary of Invention]

30 [Technical Problem]

[0005] An insecticide employed in the field of the agriculture suffers from a problem which is the appearance of a pest which has acquired a resistance against a particular drug when such a drug has been employed for a prolonged period.

35 **[0006]** In order to prevent the appearance of such a drug-resistance pest, and also to combat against such a drug-resistance pest which has already appeared, a novel insecticide having various characteristics described above are still needed to be developed in these days.

[0007] While the synthesis of a 2-trifluoromethylcarbonyl-iminothiazolidine disclosed in Patent Literature 1 and a 3-pyridylcarbonyliminoimidazolidine disclosed in Patent Literature 2 described above as insecticide compounds having imino structures was reported, no other hopeful insecticide compounds have been found.

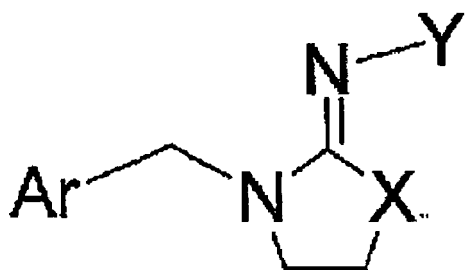
40 **[0008]** Accordingly, a primary objective of the present invention is to provide a novel imino derivative capable of being an insecticide compound excellent in such characteristics as sustained effects and broad spectrum.

[Solution to Problem]

45 **[0009]** In order to solve the problems described above, the invention provides an imino derivative represented by Formula (1):

[C.1]

50



55

• • • Formula (1)

wherein "Ar" denotes a 3-pyridyl group or a 5-thiazolyl group which have a substituent on the ring, wherein the substituent is selected from a halogen atom, a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, and a nitro group; "X" denotes a sulfur atom or CH₂;

when "Y" is COR₁, "R₁" denotes a hydrogen atom or a C1 to C5 alkyl group, a halogenated methyl except for a trifluoromethyl group, a C2 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C4)alkoxy(C1 to C5)alkyl group, a C1 to C3 alkoxy carbonyl group, a (C1 to C3)alkylsulfonyl (C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a cyano(C1 to C3)alkyl group, a substituted or unsubstituted phenoxy (C1 to C3)alkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted imidazolylmethyl group, a furanyl group, a morpholino group, a norbornenyl group, an adamantyl group, an isothiocyanatomethyl group, or substituted or unsubstituted heterocyclic or unsubstituted aromatic ring; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkylmethyl group, the phenoxy (C1 to C3)alkyl group, the pyridylmethyl group and the imidazolylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the heterocyclic or aromatic ring is substituted, each of the substituents is selected from a halogen atom, a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group and an acetylamino group;

when "Y" is CONHD, "D" denotes a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxy-carbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridylmethyl group a substituted or unsubstituted benzenesulfonyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the benzenesulfonyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, an acetylamino group, and a C1 to C3 alkyl group; when "Y" is CONHCOR₅, "R₅" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a substituted or unsubstituted phenyl group; when the phenol group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, and a C1 to C3 alkyl group;

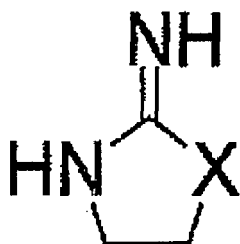
when "Y" is CO₂R₉, "R₉" denotes a hydrogen atom or a C1 to C7 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a tri(C1 to C3 alkyl)silyl(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a substituted or unsubstituted phenyl group, a substituted or unsubstituted phenylmethyl group, a substituted or unsubstituted furanylmethyl group, a substituted or unsubstituted thienylmethyl group, a substituted or unsubstituted pyridylmethyl group, a succinimide group, a 18-crown-6-methyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkylmethyl group, the furanylmethyl group, the thienylmethyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the phenyl group and the phenylmethyl group are substituted, each of the substituents is selected from a halogen atom, a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group, and an acetylamino group; and the above-mentioned carbon chains may be substituted with halogen.

[0010] This imino derivative has an excellent insecticidal activity against a broad range of agricultural pests and household pests.

[0011] The invention also provides a method for producing the imino derivative described above.

[0012] The first step of this production method is a step for producing a compound represented by Formula (4) by reacting a compound represented by Formula (2) with a compound (3) represented by Formula (3);

[C.2]

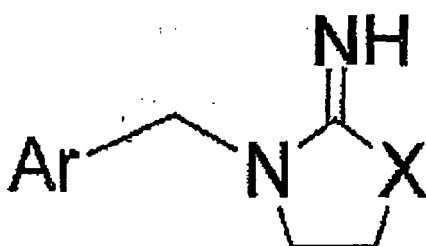


• • • Formula (2)

wherein "X" denotes a sulfur atom or CH₂.
[C.3]



wherein "Ar" has the same meaning as above; and "Z" denotes Cl or Br, I, OSO₂CH₃, OSO₂C₆H₅, OSO₂C₆H₄CH₃;
[C.4]



• • • Formula (4)

wherein "Ar" and "X" have the same meaning as above.

[0013] Then, the second step of this production method is a step (I) for producing a compound (1), which is an imino derivative represented by Formula (1) shown above wherein "Y" is defined similarly to "ACO", by reacting one compound selected from the group consisting of compounds represented by Formula (5), anhydrides represented by Formula (6) and carboxylic acids represented by Formula (7) with a compound represented by Formula (4) shown above;

[C.5]



[C.6]



Formula (6)
[C.7]



wherein "B" denotes a halogen atom of Cl or Br, I, an OCOA group (a group for med by allowing each of an oxygen atom and a group A to form a single bond with a carbonyl group (CO)) or a hydroxyl group;

when "ACO" is COR₁, "R₁" denotes a hydrogen atom or a C1 to C5 alkyl group, a halogenated methyl except for a trifluoromethyl group, a C2 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C4)alkoxy(C1 to C5)alkyl group, a C1 to C3 alkoxy carbonyl group, a (C1 to C3)alkylsulfonyl (C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a cyano(C1 to C3)alkyl group, a substituted or unsubstituted phenoxy (C1 to C3)alkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted imidazolylmethyl group, a furanyl group, a morpholino group, a norbornenyl group, an adamantyl group, an isothiocyanato-

methyl group, or substituted or unsubstituted heterocyclic or aromatic ring; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkylmethyl group, the phenoxy (C1 to C3)alkyl group, the pyridylmethyl group and the imidazolylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the heterocyclic or aromatic ring is substituted, each of the substituents is selected from a halogen atom and a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group and an acetylamino group; when "ACO" is CONHD, "D" denotes a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxycarbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted benzenesulfonyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the benzenesulfonyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, an acetylamino group, and a C1 to C3 alkyl group; when "ACO" is CO₂R₉, "R₉" denotes a hydrogen atom or a C1 to C7 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, a substituted or unsubstituted C6 to C10 aryl group, a substituted or unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a tri(C1 to C3 alkyl)silyl(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a substituted or unsubstituted phenyl group, a substituted or unsubstituted phenylmethyl group, a substituted or unsubstituted furanylmethyl group, a substituted or unsubstituted thienylmethyl group, a substituted or unsubstituted pyridylmethyl group, a succinimide group, a 18-crown-6-methyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkylmethyl group, the furanylmethyl group, the thienylmethyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the phenyl group and the phenylmethyl group are substituted, each of the substituents is selected from a halogen atom, a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group, and an acetetyl amino group; and the abovementioned carbon chains may be substituted with halogen.

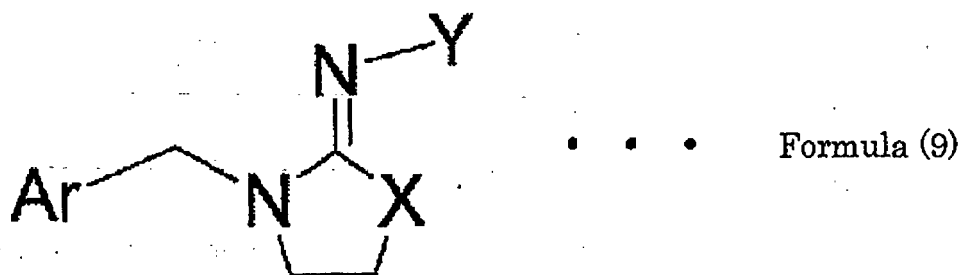
[0014] The second step of this production method may be Step (II) for producing an imino derivative represented by Formula (9) by reacting an isocyanate compound represented by Formula (8) with a compound represented by Formula (4) shown above;

[C.8]



Formula (8)

wherein "D" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxycarbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted benzenesulfonyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the benzenesulfonyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, an acetylamino group, and a C1 to C3 alkyl group; COR₅; and "R₅" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a substituted or unsubstituted phenyl group; when the phenyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, and a C1 to C3 alkyl group;



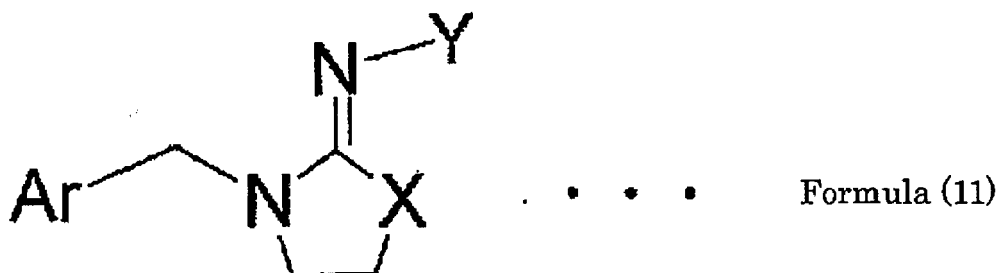
wherein "Ar", "X" and "Y" is CONHD and "D" has the same meaning as in claim 1.

15 **[0015]** The second step of this production method may further be Step (III) for producing an imino derivative represented by Formula (11) by reacting a formate represented by Formula (10) with a compound represented by Formula (4) shown above;

[C.10]



[C.11]



wherein "Ar", "X" and "Y" is CONHD and "D" has the same meaning as in claim 1.

35 **[0016]** The invention further provides an insecticide containing as an active ingredient the imino derivative described above.

[Advantageous Effect of the Invention]

40 **[0017]** According to the invention, a novel imino derivative capable of being an insecticide compound having excellent characteristics such as sustained effects and broad spectrum is provided.

[Brief Description of Drawings]

45 **[0018]**

[Fig. 1] Fig. 1 shows a reaction scheme of the first step of the method for producing a novel imino derivative according to the invention.

50 [Fig. 2] Fig. 2 shows a reaction scheme of the second step (I) of the method for producing the novel imino derivative according to the invention.

[Fig. 3] Fig. 3 shows a reaction scheme of the second step (II) of the method for producing the novel imino derivative according to the invention.

[Fig. 4] Fig. 4 shows a reaction scheme of the second step (III) of the method for producing the novel imino derivative according to the invention.

55 [Description of Embodiments]

[0019] Preferred embodiments for carrying out the invention are described below with referring to the figures. The

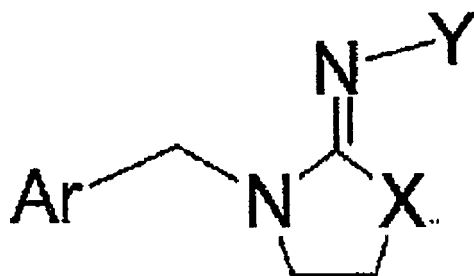
embodiments described below are only examples of representative embodiments of the invention and do not serve to allow the scope of the invention to be interpreted narrowly.

(A) Novel imino derivatives

[0020] A novel imino derivative according to the invention is represented by Formula (1). Hereinafter, this imino derivative is discussed in detail.

[0021]

[C.12]



• • • Formula (1)

[0022] In Formula (1), "Ar" a 3-pyridyl group or a 5-thiazolyl group.

[0023] The substituent on the heterocyclic ring is not limited particularly and may for example be a halogen atom (any of fluorine, chlorine, bromine and iodine), a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, and a nitro group. Preferably, a chlorine atom is employed as a substituent on the heterocyclic group.

[0024] In Formula (1), "X" denotes a sulfur atom or CH₂

[0025] In Formula (1), when "Y" is COR₁, "R₁" denotes a hydrogen atom or a C1 to C5 alkyl group, a halogenated methyl except for a trifluoromethyl group, a C2 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C4)alkoxy(C1 to C5)alkyl group, a C1 to C3 alkoxy carbonyl group, a (C1 to C3)alkylsulfonyl (C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a cyano(C1 to C3)alkyl group, a substituted or unsubstituted phenoxy (C1 to C3)alkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted imidazolylmethyl group, a furanyl group, a morpholino group, a norbornenyl group, an adamantyl group, an isothiocyanatomethyl group, or a substituted or unsubstituted heterocyclic or aromatic ring.

[0026] The C1 to C5 alkyl group may be any of primary, secondary and tertiary groups, and may for example be a methyl group, an ethyl group, an n-propyl group, an i-propyl group, an n-butyl group, an i-butyl group, a sec-butyl group, and a tert-butyl group. The alkyl group in the C2 to C5 halogenated alkyl group may be any of primary, secondary, and tertiary groups, and may for example be an ethyl group, an n-propyl group, an i-propyl group, an n-butyl group, an i-butyl group, a sec-butyl group, and a tert-butyl group. The C3 to C7 substituted or unsubstituted cycloalkyl group may for example be a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, and a cyclohexyl group. The substituent is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine and iodine) or a C1 to C3 alkyl group. The 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group is desirably a heterocycloalkyl group containing 1 to 2 heteroatoms such as oxygen atoms or sulfur atoms, nitrogen atoms. The substituent is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine and iodine) or a C1 to C3 alkyl group. The substituent substituting on the phenyl group in the substituted phenoxy(C1 to C3)alkyl group is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine and iodine) or a C1 to C3 alkyl group. The substituent attaching to the pyridine ring in the substituted pyridylmethyl group or the imidazole ring in the substituted imidazolylmethyl group is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a C1 to C3 alkyl group. Typical examples of the 5-membered and 6-membered heterocyclic ring include a quinoline ring, an indole ring, a pyridine ring, a pyrazine ring, a pyridazine ring, a pyrimidine ring, a thiophene ring, a thiazole ring, a tetrahydrofuran ring, and a furan ring. Those especially preferred are a 3-pyridyl group, a 5-thiazolyl group, and a 3-tetrahydrofuryl group. The substituent on the heterocyclic ring or an aromatic ring is not limited particularly and may for example be a halogen atom (any of fluorine, chlorine, bromine, and iodine), a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group, and an acetamino group.

[0027] It is preferred especially that "R₁" is a trichloromethyl group, a methyl group, a chloromethyl group, a difluoromethyl group, a methoxymethyl group, a 2-methylthioethyl group, a cyclopropyl group, a 2,2-difluorocyclopropyl group.

[0028] In Formula (1), when "Y" is CONHD "D" denotes a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, a C1 to C3 alkoxy group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxycarbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted benzenesulfonyl group; "NR₃R₄" may form a ring.

[0029] The C1 to C5 alkyl group may be any of primary, secondary, and tertiary groups, and may desirably be a methyl group, an ethyl group, an n-propyl group, an i-propyl group, an n-butyl group, an i-butyl group, a sec-butyl group, a tert-butyl group and the like. The C1 to C5 halogenated alkyl group may be any of primary, secondary, and tertiary groups, and may desirably be a methyl group, an ethyl group, an n-propyl group, an i-propyl group, an n-butyl group, an i-butyl group, a sec-butyl group, a tert-butyl group and the like. The C3 to C7 substituted or unsubstituted cycloalkyl group may for example be a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, and a cyclohexyl group. The substituent is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a C1 to C3 alkyl group. The 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group is desirably a heterocycloalkyl group containing 1 to 2 heteroatoms such as oxygen atoms or sulfur atoms, nitrogen atoms. The substituent is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a C1 to C3 alkyl group. The substituent attaching to the pyridine ring in the substituted pyridylmethyl group is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a C1 to C3 alkyl group. The substituent attaching to the benzene ring in the substituted benzene sulfonyl group is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a nitro group, a di(C1 to C4 alkyl)amino group, an acetamino group, a C1 to C3 alkyl group. When "NR₃R₄" forms a ring, each of R₃ and R₄ is preferably a C2 to C6 cyclic structure or a 3-membered to 7-membered cyclic structure containing, in the ring of the 3-membered to 7-membered ring represented as NR₃R₄, 1 to 2 heteroatoms such as oxygen atoms or sulfur atoms, nitrogen atoms. Especially preferably, "CONR₃R₄" has, as its "NHR₄" moiety, a methoxylamino group or an ethoxylamino group.

[0030] In Formula (1), when "Y" is CONHCOR₅, "R₅" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a substituted or unsubstituted C6 to C10 aryl group, a substituted or unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a substituted or unsubstituted phenyl group.

[0031] The C1 to C5 alkyl group may be any of primary, secondary and, tertiary groups, and may for example be a methyl group, an ethyl group, an n-propyl group, an i-propyl group, an n-butyl group, an i-butyl group, a sec-butyl group, and a tert-butyl group. The C1 to C5 halogenated alkyl group may be any of primary, secondary, and tertiary groups, and may desirably be a methyl group, an ethyl group, an n-propyl group, an i-propyl group, an n-butyl group, an i-butyl group, a sec-butyl group, a tert-butyl group and the like. The C3 to C7 substituted or unsubstituted cycloalkyl group may for example be a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, and a cyclohexyl group. The substituent is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a C1 to C3 alkyl group. The substituent attaching to the benzene ring in the substituted phenyl group is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a nitro group, a di(C1 to C4 alkyl)amino group, an acetamino group, a C1 to C3 alkyl group.

[0032] In Formula (1), when "Y" is CO₂R₉, "R₉" denotes a hydrogen atom or a C1 to C7 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, a substituted or unsubstituted C6 to C10 aryl group, a substituted or unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a tri(C1 to C3 alkyl)silyl(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a substituted or unsubstituted phenyl group, a substituted or unsubstituted phenylmethyl group, a substituted or unsubstituted furanylmethyl group, a substituted or unsubstituted thienylmethyl group, a substituted or unsubstituted pyridylmethyl group, a succinimide group, a 18-crown-6-methyl group. The C1 to C7 alkyl group may be any of primary, secondary, and tertiary groups, and may for example be a methyl group, an ethyl group, an n-propyl group, an i-propyl group, an n-butyl group, an i-butyl group, a sec-butyl group, a tert-butyl group, an n-pentyl group, and an n-hexyl group. The alkyl group in the C1 to C5 halogenated alkyl group may be any of primary, secondary, and tertiary groups, and may for example be a methyl group, an ethyl group, an n-propyl group, an i-propyl group, an n-butyl group, an i-butyl group, a sec-butyl group, and a tert-butyl group. The C3 to C7 substituted or unsubstituted cycloalkyl group may for example be a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, and a cyclohexyl group. The substituent is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a C1 to C3 alkyl group. The 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group is desirably a

heterocycloalkyl group containing 1 to 2 heteroatoms such as oxygen atoms or sulfur atoms, nitrogen atoms. The substituent is not limited particularly, and may for example be a halogen atom (any of chlorine, bromine, fluorine, and iodine) or a C1 to C3 alkyl group. The substituent substituting on the phenyl group in the substituted phenyl group is not limited particularly, and may for example be a halogen atom (any of fluorine, chlorine, bromine, and iodine), a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group, and an acetamino group. The substituent attaching to the phenyl ring, the furan ring, the thiophene ring, the pyridine ring, the imidazole ring in the substituted phenylmethyl group or the substituted furanyl methyl group, the substituted thienylmethyl group, the substituted pyridylmethyl group, the substituted imidazolylmethyl group is desirably a halogen atom (any of chlorine, bromine, fluorine, and iodine), a C1 to C3 alkyl group, and the like.

[0033] It is especially preferable that "R₉" is an n-propyl group, an i-propyl group, a 2-chloroethyl group, an ethyl group, a 2-fluoroethyl group, a 2,2-difluoroethyl group, a 2-methoxyethyl group, a 2-methylthioethyl group, a 3-fluoropropyl group, a 2-propenyl group, an n-butyl group, an i-butyl group, a sec-butyl group, a 3-methyl-2-butenyl group, an n-pentyl group, a cyclopropylmethyl group, a 3-oxetanyl group, a 3-tetrahydrofuranylmethyl group, or a 4-tetrahydropyranylmethyl group.

[0034] The carbon chain in Formula (1) described above may be substituted with halogens.

[0035] A novel imino derivative according to the invention has excellent insecticidal activity, acaricidal activity and the like against a broad range of agricultural pests and household pests. Accordingly, it can be used as a pesticide containing such compounds as an active ingredient, such as an insecticide or acaricide against agricultural pests and household pests and as a domestic pest-controlling agent, such as termite-controlling agent, as well as a veterinary pharmaceutical.

(B) Methods for producing novel imino derivatives

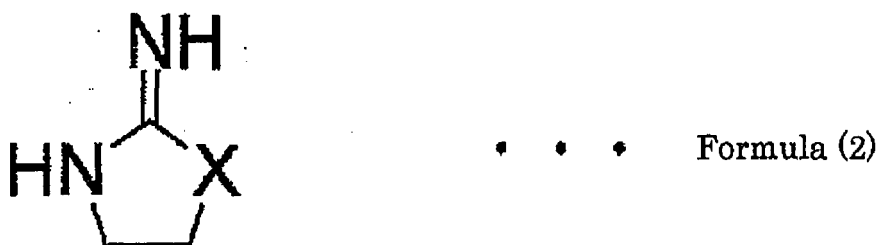
[0036] A method for producing a novel imino derivative according to the invention is described below.

(1) First step

[0037] In the first step of this production method, a compound represented by Formula (4) is produced by reacting a compound represented by Formula (2) with a compound (3) represented by Formula (3) in the presence of a base. The reaction scheme of the first step is shown in Fig. 1.

[0038]

[C.13]



wherein "X" denotes a sulfur atom or CH₂.

[0039]

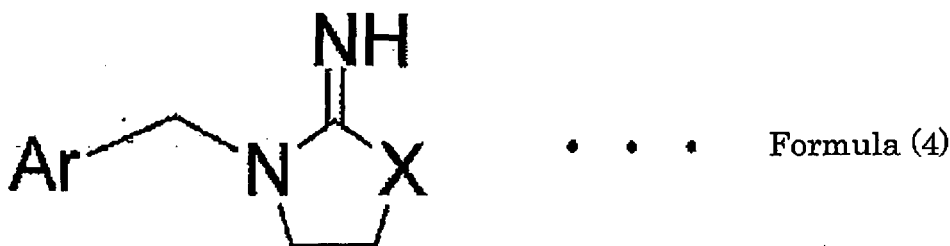
[C.14]



wherein "Ar" denotes a 3-pyridyl group or a 5-thiazolyl group which may have a substituent on the ring; "Z" denotes Cl or Br, I, OSO₂CH₃, OSO₂C₆H₅, OSO₂C₆H₄CH₃;

[0040]

[C.15]



10 wherein "Ar" denotes a 3-pyridyl group or a 5-thiazolyl group which may have a substituent on the ring; "X" denotes a sulfur atom or CH₂.

[0041] In the first step, 1 mole of the compound represented by Formula (4) can be produced by reacting 1 mole of the compound represented by Formula (2) with 1.0 to 1.5 moles of a halogenated compound represented by Formula (3).

15 [0042] In the first step, the amount of the compound (3) represented by Formula (3) to be added is preferably 1.0 to 1.1 moles per mole of the compound (2) described above.

[0043] This compound (4) may be synthesized by a known production method (for example, Journal of Medical Chemistry, 1999, 42(12), 2227-2234) or a method analogous thereto.

20 [0044] While the type of the base employed in the first step is not limited particularly, it is preferably an alkaline metal hydride such as sodium hydride (NaH), a carbonate such as potassium carbonate (K₂CO₃) or sodium carbonate (Na₂CO₃), an alkaline metal hydroxide such as potassium hydroxide (KOH), sodium hydroxide (NaOH), tertiary amines such as triethylamine [(C₂H₅)₃N] and the like.

25 [0045] It is preferred in the first step to use a solvent, which may for example be amides such as dimethyl formamide (DMF) and dimethyl acetamide, nitriles such as acetonitrile, sulfoxides such as dimethyl sulfoxide (DMSO), ethers such as diethyl ether and tetrahydrofuran (THF), aromatic hydrocarbons such as benzene, xylene, and toluene, alcohols such as ethanol and propanol, ketones such as acetone and methylethylketone, aliphatic hydrocarbons such as hexane, heptane, and octane, halogenated hydrocarbons such as dichloromethane, chloroform, chlorobenzene, and dichlorobenzene, and water, which may be employed alone or in combination of one or more thereof.

30 [0046] To the reaction mixture, a phase transfer catalyst including a quaternary ammonium salt such as tetrabutylammonium salt or a crown ether or analogues thereto may be added to proceed these reactions. In such a case, the solvent employed is not limited particularly, and the oil phase may be benzene, chloroform, dichloromethane, hexane, toluene and the like.

35 [0047] The reaction temperature in the first step is preferably 0 to 200°C. A reaction temperature below 0°C results in a slow reaction rate, while that exceeding 200°C results in a too rapid reaction rate, which is disadvantageous since it allows a side reaction to be proceeded easily. The reaction time in the first step may appropriately be selected based on the temperature condition and the pressure condition, and it is preferably within the range of 30 minutes to 24 hours.

(2-1) Second step (I)

40 [0048] In the second step of this production method, a compound (1), which is an imino derivative represented by Formula (1) shown above wherein "Y" is defined similarly to "ACO", is produced by reacting a compound (5) represented by Formula (5) with a compound represented by Formula (4) obtained in the first step in the presence of a base. The reaction scheme of the second step (I) is shown in Fig. 2.

[0049]

45 [C.16]

ACO-B

Formula (5)

50 wherein "B" denotes a halogen atom of Cl or Br, I, an OCOA group (a group formed by allowing each of an oxygen atom and a group A to form a single bond with a carbonyl group (CO)) or a hydroxyl group; when "ACO" is COR₁, "R₁" denotes a hydrogen atom or a C1 to C5 alkyl group, a halogenated methyl except for a trifluoromethyl group, a C2 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C4)alkoxy(C1 to C5)alkyl group, a C1 to C3 alkoxy carbonyl group, a (C1 to C3)alkylsulfonyl (C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a cyano(C1 to C3)alkyl group, a substituted or unsubstituted phenoxy (C1 to C3)alkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted

imidazolymethyl group, a furanyl group, a morpholino group, a norbornenyl group, an adamantyl group, an isothiocyanatomethyl group, or substituted or unsubstituted heterocyclic or aromatic ring;

when "ACO" is CONR_3R_4 , each of "R₃" and "R₄" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, a C1 to C3 alkoxy group, a substituted or unsubstituted C6 to C10 aryl group, a substituted or unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxycarbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted benzenesulfonyl group; "NR₃R₄" may form a ring;

when "ACO" is CO_2R_9 , "R₉" denotes a hydrogen atom or a C1 to C7 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, a substituted or unsubstituted C6 to C10 aryl group, a substituted or unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a tri(C1 to C3 alkyl)silyl(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a substituted or unsubstituted phenyl group, a substituted or unsubstituted phenylmethyl group, a substituted or unsubstituted furanylmethyl group, a substituted or unsubstituted thienylmethyl group, a substituted or unsubstituted pyridylmethyl group, a succinimide group, a 18-crown-6-methyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkylmethyl group, the phenoxy (C1 to C3)alkyl group, the pyridylmethyl group and the imidazolymethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the heterocyclic or aromatic ring is substituted, each of the substituents is selected from a halogen atom and a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group and an acetylamino group;

[0050] In the second step (I), 1 mole of the novel imino derivative represented by Formula (1) can be produced by reacting 1 mole of the compound represented by Formula (4) with 1 to 2 moles of a halogenated compound represented by Formula (5).

[0051] In the second step (I), the amount of the compound (5) represented by Formula (5) to be added is preferably 1.0 to 1.2 moles per mole of the compound represented by Formula (4).

[0052] While the type of the base employed in the second step (I) is not limited particularly, it is preferably an alkaline metal hydride such as sodium hydride (NaH), a carbonate such as potassium carbonate (K_2CO_3) or sodium carbonate (Na_2CO_3), an alkaline metal hydroxide such as potassium hydroxide (KOH), sodium hydroxide (NaOH), tertiary amines such as triethylamine $[(\text{C}_2\text{H}_5)_3\text{N}]$ and the like.

[0053] It is desirable in the second step (I) to use a solvent, which may for example be amides such as dimethyl formamide (DMF) and dimethyl acetamide, nitriles such as acetonitrile, sulfoxides such as dimethyl sulfoxide (DMSO), ethers such as diethyl ether and tetrahydrofuran (THF), aromatic hydrocarbons such as benzene, xylene, and toluene, alcohols such as ethanol and propanol, ketones such as acetone and methylethylketone, aliphatic hydrocarbons such as hexane, heptane, and octane, halogenated hydrocarbons such as dichloromethane, chloroform, chlorobenzene, and dichlorobenzene, and water, which may be employed alone or in combination of one or more thereof.

[0054] To the reaction mixture in the second step (I), a phase transfer catalyst including a quaternary ammonium salt such as tetrabutylammonium salt or a crown ether or analogues thereto may be added to proceed these reactions. In such a case, the solvent employed is not limited particularly, and the oil phase may be benzene, chloroform, dichloromethane, hexane, toluene and the like.

[0055] The reaction temperature in the second step (I) is preferably 0 to 200°C. A reaction temperature below 0°C results in a slow reaction rate, while that exceeding 200°C results in a too rapid reaction rate, which is disadvantageous since it allows a side reaction to be proceeded easily. While the reaction in the second step (I) can be conducted under reduced pressure, at ambient temperature, or under pressure, it is conducted preferably at ambient temperature. The reaction time in the second step (I) may appropriately be selected based on the temperature condition and the pressure condition, and it is preferably within the range of 30 minutes to 24 hours.

[0056] In the second step (I) of this production method, an imino derivative represented by Formula (1) shown above can be produced also by reacting an anhydride represented by Formula (6) or a carboxylic acid represented by Formula (7), instead of the halogenated compound represented by Formula (5), with a compound represented by Formula (4) obtained in the first step in the presence of a base.

[0057]

[C.17]



Formula (6)

[0058]

[C.18]



Formula (7)

[0059] When using a carboxylic acid represented by Formula (7), a dehydrating condensation agent such as dicyclohexyl carbodiimide, 1-alkyl-2-halopyridinium salt, 1,1-carbonium imidazole, 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride is employed in the presence of a base.

(2-2) Second step (II)

[0060] In the second step of this production method, an imino derivative represented by Formula (9) may be produced by reacting an isocyanate compound represented by Formula (8) with a compound represented by Formula (4) shown above in the presence of a base. The reaction scheme of the second step (II) is shown in Fig. 3.

[0061]

[C.19]

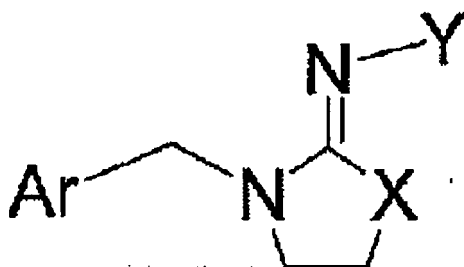


Formula (8)

wherein "D" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxycarbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted benzenesulfonyl group, when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the benzenesulfonyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, an acetvlamino group, and a C1 to C3 alkyl group ; COR₅; and "R₅" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a substituted or unsubstituted phenyl group; when the phenyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl amino group, and a C1 to C3 alkyl group;

[0062]

[C.20]



Formula (9)

wherein "Ar", "X" and "Y" is CONHD and "D" has the same meaning as in claim 1.

[0063] In the second step (II), 1 mole of the novel imino derivative represented by Formula (9) can be produced by reacting 1 mole of the compound represented by Formula (4) with 1 mole of an isocyanate compound represented by

Formula (8).

[0064] In the second step (II), the amount of the isocyanate compound represented by Formula (8) to be added is preferably 1.0 to 1.2 moles per mole of the compound represented by Formula (4) described above.

[0065] While the type of the base employed in the second step (II) is not limited particularly, it is preferably an alkaline metal hydride such as sodium hydride (NaH), a carbonate such as potassium carbonate (K_2CO_3) or sodium carbonate (Na_2CO_3), an alkaline metal hydroxide such as potassium hydroxide (KOH), sodium hydroxide (NaOH), tertiary amines such as triethylamine $[(C_2H_5)_3N]$ and the like.

[0066] It is desirable in the second step (II) to use a solvent, which may for example be amides such as dimethyl formamide (DMF) and dimethyl acetamide, nitriles such as acetonitrile, sulfoxides such as dimethyl sulfoxide (DMSO), ethers such as diethyl ether and tetrahydrofuran (THF), aromatic hydrocarbons such as benzene, xylene, and toluene, alcohols such as ethanol and propanol, ketones such as acetone and methylethylketone, aliphatic hydrocarbons such as hexane, heptane, and octane, halogenated hydrocarbons such as dichloromethane, chloroform, chlorobenzene, and dichlorobenzene, and water, which may be employed alone or in combination of one or more thereof.

[0067] To the reaction mixture in the second step (II), a phase transfer catalyst including a quaternary ammonium salt such as tetrabutylammonium salt or a crown ether or analogues thereto may be added to proceed these reactions. In such a case, the solvent employed is not limited particularly, and the oil phase may be benzene, chloroform, dichloromethane, hexane, toluene and the like.

[0068] The reaction temperature in the second step (II) is preferably 0 to 200°C. A reaction temperature below 0°C results in a slow reaction rate, while that exceeding 200°C results in a too rapid reaction rate, which is disadvantageous since it allows a side reaction to be proceeded easily. While the reaction in the second step (II) can be conducted under reduced pressure, at ambient temperature, or under pressure, it is preferably conducted at ambient temperature. The reaction time in the second step (II) may appropriately be selected based on the temperature condition and the pressure condition, and it is preferably within the range of 30 minutes to 24 hours.

(2-3) Second step (III)

[0069] Moreover, in the second step of this production method, an imino derivative represented by Formula (11) may be produced by reacting a formate represented by Formula (10) with a compound represented by Formula (4) shown above in the presence of a base. The reaction scheme of the second step (III) is shown in Fig. 4.

[0070]

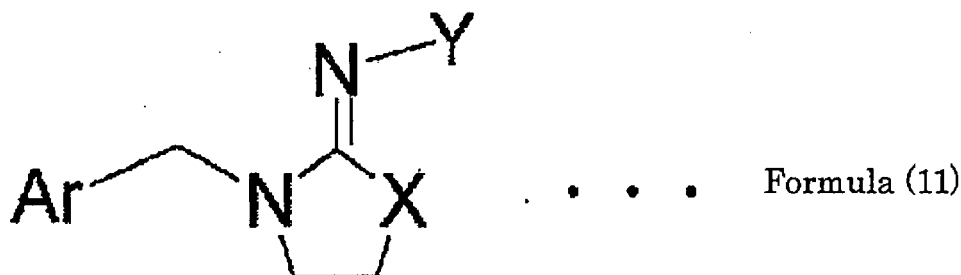
[C.21]

HCO_2Et

Formula (10)

[0071]

[C.22]



wherein "Ar", "X" and "Y" denotes CHO.

[0072] In the second step (III), 1 mole of the novel imino derivative represented by Formula (11) can be produced by reacting 1 mole of the compound represented by Formula (4) with 1 mole of a formate represented by Formula (10).

[0073] In the second step (III), the amount of the formate represented by Formula (10) to be added is preferably 1 to 1.2 moles per mole of the compound represented by Formula (4) described above.

[0074] While the type of the base employed in the second step (III) is not limited particularly, it is preferably an alkaline metal hydride such as sodium hydride (NaH), a carbonate such as potassium carbonate (K_2CO_3) or sodium carbonate (Na_2CO_3), an alkaline metal hydroxide such as potassium hydroxide (KOH), sodium hydroxide (NaOH), tertiary amines

such as triethylamine $[(C_2H_5)_3N]$ and the like.

[0075] It is desirable in the second step (III) to use a solvent, which may for example be amides such as dimethyl formamide (DMF) and dimethyl acetamide, nitriles such as acetonitrile, sulfoxides such as dimethyl sulfoxide (DMSO), ethers such as diethyl ether and tetrahydrofuran (THF), aromatic hydrocarbons such as benzene, xylene, and toluene, alcohols such as ethanol and propanol, ketones such as acetone and methylethylketone, aliphatic hydrocarbons such as hexane, heptane, and octane, halogenated hydrocarbons such as dichloromethane, chloroform, chlorobenzene, and dichlorobenzene, and water, which may be employed alone or in combination of one or more thereof.

[0076] To the reaction mixture in the second step (III), a phase transfer catalyst including a quaternary ammonium salt such as tetrabutylammonium salt or a crown ether or analogues thereto may be added to proceed these reactions. In such a case, the solvent employed is not limited particularly, and the oil phase may be benzene, chloroform, dichloromethane, hexane, toluene and the like.

[0077] The reaction temperature in the second step (III) is preferably 0 to 200°C. A reaction temperature below 0°C results in a slow reaction rate, while that exceeding 200°C results in a too rapid reaction rate, which is disadvantageous since it allows a side reaction to be proceeded easily. While the reaction in the second step (III) can be conducted under reduced pressure, at ambient temperature, or under pressure, it is preferably conducted at ambient temperature. The reaction time in the second step (III) may appropriately be selected based on the temperature condition and the pressure condition, and it is preferably within the range of 30 minutes to 24 hours.

[0078] The first step and the second steps (I) to (III) can also be conducted continuously while skipping a step for isolating the intermediate which is a compound represented by Formula (4). In such a case, after completing the first step, certain amounts of the halogenated compound represented by Formula (5) and the base employed in the second step are added, and a certain reaction temperature is established, whereby allowing the reaction of the second step to be carried out consecutively.

[0079] A compound (1) wherein "Y" is "CONR₃R₄" [CONR₃R₄ in which R₃ is a hydrogen or a C1 to C5 alkyl group and R₄ is a C1 to C3 alkoxy group, preferably CON(hydrogen or methyl)(C1 to C3-alkoxy) group, more preferably, methoxylaminocarbonyl, ethoxylaminocarbonyl] can be produced by reacting a compound wherein "Y" is "CO₂R₉" in which "R₉" is a substituted or unsubstituted phenyl group (for example, 4-nitrophenyl group) with an O-alkylhydroxylamine, an N,O-dialkylhydroxylamine or salts thereof in the presence of a base (for example, sodium acetate, or potassium carbonate) in a solvent (for example, acetonitrile).

[0080] By using a production method according to the invention described above, a novel imino derivative represented by Formula (I) according to the invention can be produced using as a starting material a substance which has a relatively high versatility.

(C) Insecticides

[0081] A novel imino derivative according to the invention can be preferably employed as an insecticide. An agro-horticultural insecticide containing a novel imino derivative represented by Formula (1) as an active ingredient is discussed below.

[0082] Agricultural pests against which an inventive insecticide can exert its controlling effect are listed below. Lepidoptera pests may for example be oriental corn borer (*Ostrinia furnacalis*), Oriental armyworm moth (*Pseudaletia separata*), pink borer (*Sesamia inferens*), corn earworm (*Heliothis* sp.), turnip moth (*Agrotis segetum*), apple leafminer (*Phyllonorycter ringoniella*), diamondback moth (*Plutella xylostella*), rice leafroller (*Cnaphalocrocis medinalis*), yellow stemborer (*Scirpophaga incertulas*), beet armyworm (*Spodoptera exigua*), rice stem borer (*Chilo suppressalis*), common cutworm (*Spodoptera litura*), soybean pod borer (*Leguminivora glycinivorella*), peach fruit moth (*Carposina niponensis*), common cabbage worm (*Pieris rapae crucivora*), summer fruit tortrix (*Adoxophyes orana fasciata*), and cabbage armyworm (*Mamestra brassicae*). Hemiptera pests may for example be greenhouse whitefly (*Trialeurodes vaporariorum*), sweetpotato whitefly (*Bemisia tabaci*), green rice leafhopper (*Nephotettix cincticeps*), brown rice planthopper (*Nilaparvata lugens*), turnip aphid (*Lipaphis erysimi*), citrus psyllid (*Diaphorina citri*), cottony citrus scale (*Pulvinaria aurantii*), green peach aphid (*Myzus persicae*), arrowhead scale (*Unaspis yanonensis*), and cotton aphid (*Aphis gossypii*). Coleoptera pests may for example be bean beetle (*Callosobruchus chinensis*), rice leaf beetle (*Oulema oryzae*), rice water weevil (*Lissorhoptrus oryzophilus*), cucurbit leaf beetle (*Aulacophora femoralis*), striped flea beetle (*Phyllotreta striolata*), Maize weevil (*Sitophilus zeamais*), cigarette beetle (*Lasioderma serricorne*), twenty-eight-spotted ladybird (*Epilachna vigintioctopunctata*), soybean beetle (*Anomala rufocuprea*), Brown powder-post beetle (*Lyctus brunneus*), and Japanese beetle (*Popillia japonica*). Thysanoptera pests may for example be onion thrips (*Thripstabaci*), melon thrips (*Thrips palmi*), and rice thrips (*Stenchaetothrips biformis*). Diptera pests may for example be house mosquito (*Culex pipiens*), House Fly (*Musca domestica*), rice leafminer (*Agromyza oryzae*), Melon fly (*Dacus*(*Zeugodacus*) *cucurbitae*), soybean pod gall midge (*Asphondylia* sp.), onion maggot (*Delia antiqua*), seedcorn maggot (*Delia platura*), and oriental fruit fly (*Dacus*(*Bactrocera*) *dorsalis*). Dytiscoptera pests may for example be German cockroach (*Batella germanica*), Oriental termite (*Coptotermes formosanus*), and Japanese subterranean termite (*Reticulitermes speratus*). Orthoptera pests may for example be Mi-

gratory locust (*Locusta migratoria*), Acari pests may for example be two-spotted spider mite (*Tetranychus urticae*), kanzawa spider mite (*Tetranychus kanzawai*), and citrus red mite (*Panonychus citri*), acarian pests parasiting animals may for example be soft tick (*Ornithodoros*), fleas may for example be cat flea (*Ctenocephalides felis*) and dog flea (*Ctenocephalides canis*), lice may for example be poultry lice (*Menopon*), and Trematoda, Cestoda, Nematoda, and Coccidia may also be exemplified.

[0083] Methods for using the insecticides according to the invention is not limited particularly, and it may be used as it is or as being diluted to a desired concentration with a diluent such as water. It is also possible to use several types of 6-membered heterocyclic derivatives according to the invention in combination, and a mixture with other pesticides may also be employed as long as the effects of the insecticides are not affected adversely. Other pesticides which may be employed in a mixture are not limited particularly, and may for example be other fungicides, insecticides, acaricides, herbicides, plant growth regulators, fertilizers and the like.

[0084] When using an inventive insecticide as being diluted, the concentration of the novel imino derivative is preferably within the range of 0.001 to 1.0%. The treatment rate of the inventive insecticide is adjusted in such a manner that 1 ha of an agro-horticultural site such as a field, a paddy field, an orchard, and a greenhouse is treated preferably with 20 to 5000 g, more preferably with 50 to 1000 g of a novel imino derivative. Such concentration and treatment rate are not limited to those specified above, and may appropriately be increased or decreased depending on the dosage form, the timing of use, the method of use, the site of use, the crops to be treated and the like.

[0085] The insecticide according to the invention is used as a formulation in which a novel imino derivative and a carrier are mixed. If necessary, various formulation auxiliaries, such as a surfactant, a spreading agent, a depositing agent, a thickening agent, and a stabilizer, may further be incorporated to formulate a dosage form such as a wettable powder, a granule, and a flowable formulation.

[0086] In the insecticide according to the invention, when a novel imino derivative is mixed with a carrier, the amount of the carrier is usually within the range of 0.1 to 80% by weight. The carriers for the formulations mentioned above include microparticulate or granulate solid carriers such as kaolin, attapulgite, bentonite, acid clay, pyrophyllite, talc, kieselguhr, calcite, ground walnut powder, urea, ammonium sulfate, and synthetic hydrated silicon oxide, aromatic hydrocarbons such as xylene and naphthalene, alcohols such as isopropanol, ethylene glycol, and cellosolve, ketones such as acetone, cyclohexanone, and isophorone, vegetable oils such as soybean oil and cottonseed oil, liquid carriers such as dimethyl sulfoxide, acetonitrile, and water.

[0087] The surfactant employed for emulsification, dispersion, and spreading may for example be an anionic surfactant such as an alkyl sulfate salt, an alkyl(aryl)sulfonate, a dialkylsulfosuccinate, a polyoxyethylene alkylaryl ether phosphate salt, a naphthalenesulfonic acid formaldehyde condensate, and a polycarboxylic acid-type polymer, and a nonionic surfactant such as a polyoxyethylene alkyl ether, a polyoxyethylene alkylaryl ether, a polyoxyethylene polyoxypropylene block copolymer, and a sorbitan fatty acid ester. The formulation auxiliaries may for example be a lignin sulfonate, alginate, a polyvinylalcohol, gum arabic, CMC (carboxymethyl cellulose), PAP (acidic isopropyl phosphate), and xanthane gum.

[0088] The insecticide capable of being mixed with an insecticide according to the invention may for example be an organic phosphorus-based insecticide (fenitrothion, malathion, acephate, diazinon and the like), a carbamate-based insecticide (benfuracarb, methomyl, carbosulfan and the like), a pyrethroid-based insecticide (allethrin, permethrin, fenvalerate, etofenprox, silafluofen and the like), a nereistoxin-based insecticide (cartap, thiocyclam and the like), a neonicotinoid-based insecticide (imidacloprid, acetamiprid, thiamethoxam, dinotefuran and the like), an IGR agent (diflubenzuron, cyromazine and the like), fipronil, emamectin benzoate, pyridalyl, propylene glycol fatty acid monoester, and a BT agent, a bactericide capable of being mixed may for example be a copper fungicide (inorganic copper, organic copper, nonylphenylether copper sulfonate and the like), an inorganic sulfur agent, an organic sulfur fungicide (maneb, manzeb, amobam, propineb, thiuram and the like), an organic phosphorus-based fungicide (fosethyl, tolclofos-methyl and the like), a melanin biosynthesis inhibitor (fthalide, tricyclazole, pyroquilon, carpropamid, diclocymet, fenoxanil), a benzimidazol-based fungicide (thiophanate-methyl, benomyl, diethofencarb, and the like), a dicarboxyimide agent (iprodione, procymidone and the like), an acid amid-based fungicide (mepronil, flutolanil, boscalid, furametpyr, thifluzamide, metalaxyl- and the like), a sterol biosynthesis inhibitor (triflumizole, tebuconazole, ipconazole, metconazole, epoxiconazole and the like), a strobilurin-based fungicide (azoxystrobin, kresoxim-methyl, trifloxystrobin and the like), an anilinopyrimidine-based fungicide (mepanipyrim, cyprodinil and the like), an antibacterial agent (oxolinic acid, tecloftalam and the like), an antibiotic fungicide (kasugamycin, polyoxins, streptomycin and the like), probenazole, ferimzone, TPN, fludioxonil, iminoctadine acetate, cyazofamid, and cyflufenamid, an acaricide capable of being mixed may for example be bifentazate, milbemectin, and etoxazole, a herbicide capable of being mixed may for example be a phenoxy acid-based herbicide (2,4-d, clomeprop, and fluzafop), a carbamate-based herbicide (benthiocarb, molinate, pyributicarb and the like), an acid amide-based herbicide (pretilachlor, diflufenican, mefenacet, cafenstrole, asulam and the like), an urea-based herbicide (daimuron, isouron and the like), a sulfonyl urea-based herbicide (imazosulfuron, thifensulfuron-methyl, nicosulfuron, halosulfuron methyl and the like), a triazine-based herbicide (atrazine, simetryn, simazine, triaziflam and the like), a diazine-based herbicide (bentazone, bromacil and the like), a diazole-based herbicide (pyrazolate, oxadiazon

and the like), a bipyridylium-based herbicide (paraquat and the like), a dinitroaniline-based herbicide (trifluralin, pendimethalin, oryzalin and the like), an aromatic carboxylic acid-based herbicide (fentrazamide, imazapyr and the like), a pyrimidyl benzoate-based herbicide (bispiribac sodium and the like), a fatty acid-based herbicide (tetrapion and the like), an amino acid-based herbicide (glyphosate, glufosinate and the like), a nitrile-based herbicide (chlorthiamid and the like), a cyclogexadione-based herbicide (sethoxydim, clethodim and the like), a phenylphthalimide-based herbicide (chlorphthalim and the like), butamifos, pentoxazone, and benzobicyclon, a plant growth regulator capable of being mixed may for example be uniconazole p, daminozide, paraffin, wax, and benzylaminopurine, although the agents capable of being mixed or being a composite formulation with an inventive plant anthrax controller are not limited to those listed above.

Examples

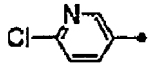
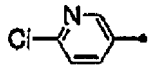
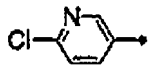
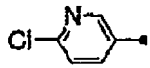
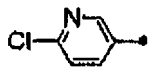
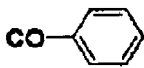
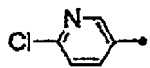
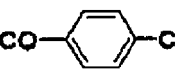
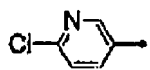
[0089] Production Examples and Formulation Examples, as well as Experimental Examples of the novel imino derivatives and insecticides according to the invention are represented below whereby a detailed description of the invention is made. The invention is not limited to the below-indicated Production Examples and the like, as long as there is no departure from its scope. The compounds employed in the following Production Examples and the like may be any commercially available ones as appropriate. The measurement of the physical data of respective intended compound obtained in respective Examples was conducted under the condition indicated in "Table 1" shown below.

[Table 1]

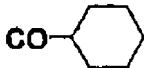
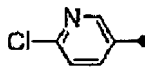
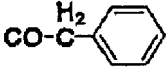
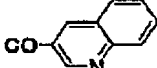
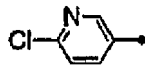
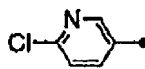
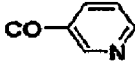
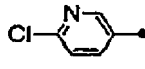
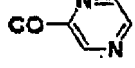
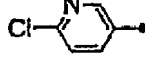
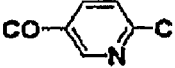
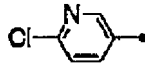
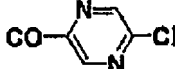
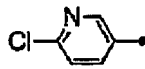
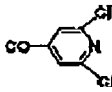
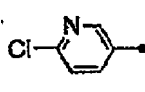
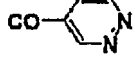
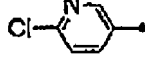
Melting point	Measured using BUECHI Model B-545
IR (Infrared spectroscopy)	Measured by KBr tablet method using HITACHI, Ltd. Model 270-30
¹ H-NMR (Proton nuclear magnetic resonance spectroscopy)	Measured by JEOL Ltd. device (400 MHz) using internal standard TMS (tetramethylsilane) and solvent DMSO-d ₆ (tritylated dimethylsulfoxide)

[0090] The intended compounds obtained in Examples of novel imino derivatives represented by Formula (1) shown above had the chemical structures and the melting points shown in "Table 2" to "Table 11", and the measured values of IR spectra and NMR spectra shown in "Table 12" to "Table 22". Among those, some Examples are detailed below with regard to Production Examples, Formulation Examples, and Experimental Examples.

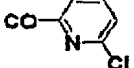
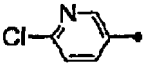
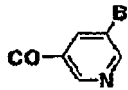
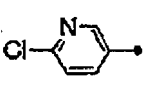
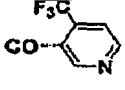
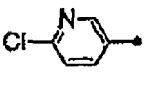
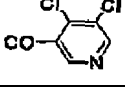
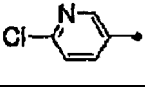
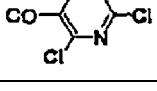
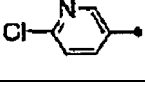
[Table 2]

Compound No.	Y	Ar	X	Melting Point (°C)
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2	-COCH ₂ CF ₃		S	88-90
3	-COC(CH ₃) ₃		S	45
4	-COC(CF ₃) ₂ CH ₃		S	114-116
5	-COCCl ₃		S	90
6			S	133
7			S	186
8			S	175

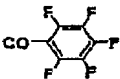
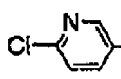
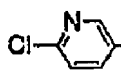
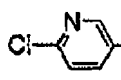
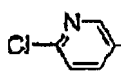
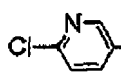
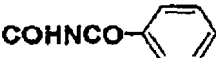
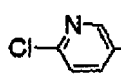
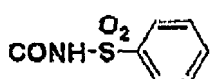
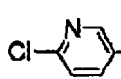
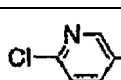
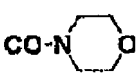
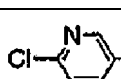
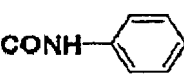
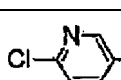
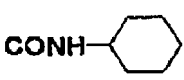
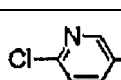
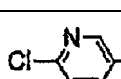
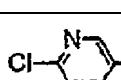
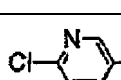
(continued)

Compound No.	Y	Ar	X	Melting Point (°C)
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10			S	157-158
11			S	180-182
12			S	185-188
13			S	117-118
14			S	157-158
15			S	160-162
16			S	134-136
17			S	165-168
18			S	204-206

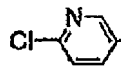
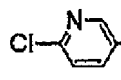
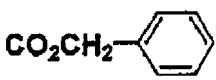
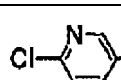
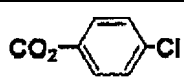
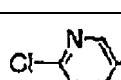
[Table 3]

19			S	128
20			S	156-158
21			S	123-125
22			S	180-181
23			S	179-182

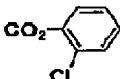
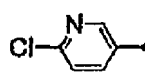
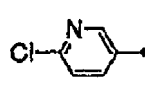
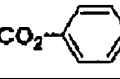
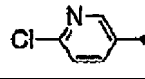
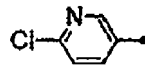
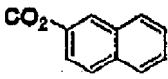
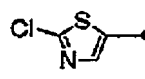
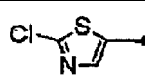
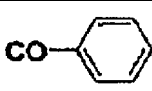
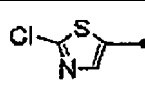
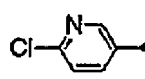
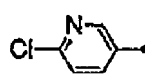
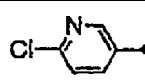
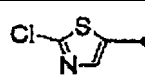
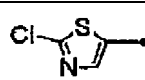
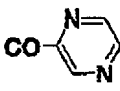
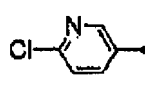
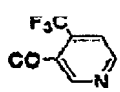
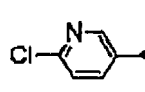
(continued)

24			S	233
28	CONH ₂		S	158-159
29	CONHCH ₃		S	175-177
30	CONHCH ₂ CO ₂ CH ₂ CH ₃		S	166-167
31	CONHCOCH ₂ Cl		S	165-166
32			S	179-181
33			S	191-193
34	CONHOH(CH ₃) ₂		S	86
35			S	133-134
36			S	170-172
37			S	74-76
42	CO ₂ OH ₃		S	128-130
43	CO ₂ CH ₂ CH ₂ CH ₃		S	97-99
44	CO ₂ CH(CH ₃) ₂		S	141-143

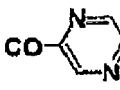
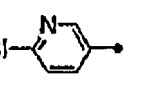
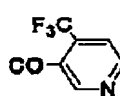
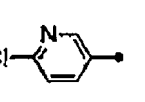
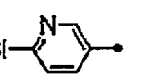
[Table 4]

45	CO ₂ CH ₂ CH ₂ Cl		S	106-108
46	CO ₂ CH ₂ CH ₂ OCH ₃		S	69-71
47			S	94-95
48			S	114-115

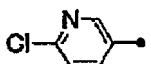
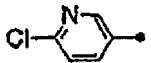
(continued)

49			S	Oil
50			S	207-209
51			S	169
52	CO ₂ C ₆ H ₁₃		S	43
53			S	193
56	COCCl ₃		S	Oil
57			S	120
58	CHO		S	99-100
59	CHO		CH ₂	Oil
60	CHO		NH	Oil
61	CHO		S	76
62	CHO		CH ₂	Oil
63			CH ₂	104-106
64			CH ₂	79-80

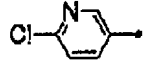
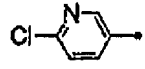
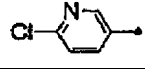
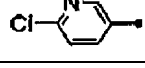
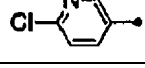
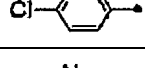
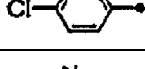
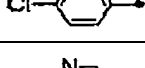
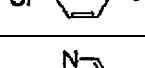
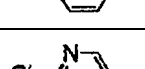
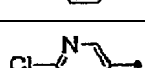
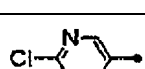
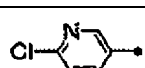
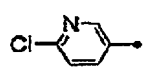
[Table 5]

66			NH	172-174
66			NH	137-139
67	-COCH(CF ₃) ₂		S	118-119

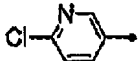
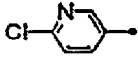
(continued)

70			S	Oil
71			S	138

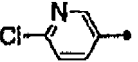
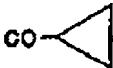
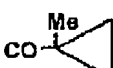
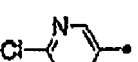
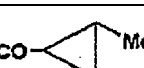
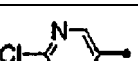
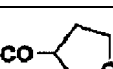
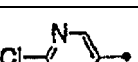
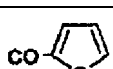
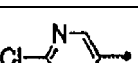
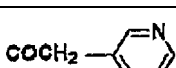
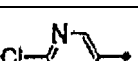
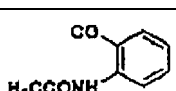
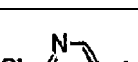
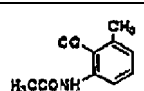
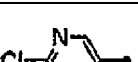
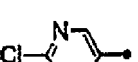
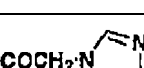
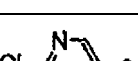
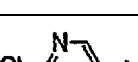
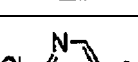
[Table 6]

Compound No.	Y	Ar	x	Melting Point (°C)
72	-COCH ₃		S	108-110
73	-COCH ₂ Cl		S	111-112
74	-COCHCl ₂		S	124-126
75	-COCH ₂ Br		S	95-97
76	-COCHF ₂		S	61
77	-COCClF ₂		S	86-88
78	-COCH ₂ OCH ₃		S	103-104
79	-COCH ₂ CN		S	122
80	-COCH ₂ OC ₆ H ₅		S	167
81	-COCH ₂ CH ₂ OCH ₃		S	56-57
82	-COCH ₂ NCS		S	147-149
83	-COCH=CH ₂		S	92-93
84	-COCCl=CCl ₂		S	98-100
85	-COCH ₂ CH=CHCH ₃		S	75-77
86	COCH ₂ -C≡CCH ₃		S	132-134
87	-COCCH ₃ (CF ₃) ₂		S	114-116

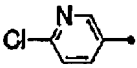
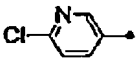
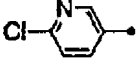
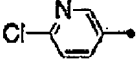
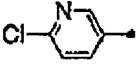
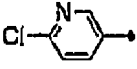
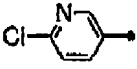
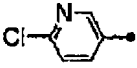
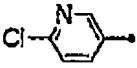
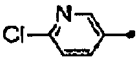
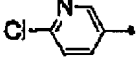
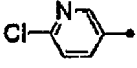
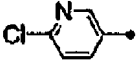
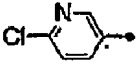
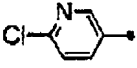
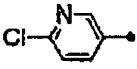
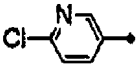
(continued)

Compound No.	Y	Ar	x	Melting Point (°C)
88	-COCO ₂ CH ₂ CH ₃		S	88-89
89	-COCH ₂ SCH ₃		S	Oil

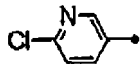
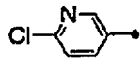
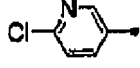
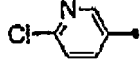
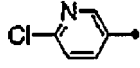
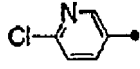
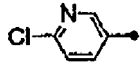
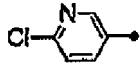
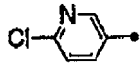
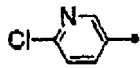
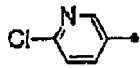
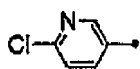
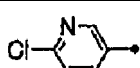
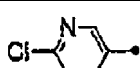
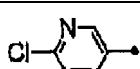
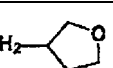
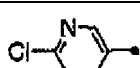
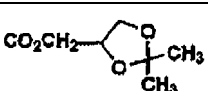
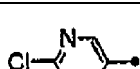
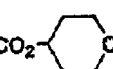
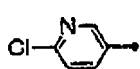
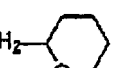
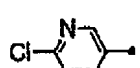
[Table 7]

90	-COCH ₂ CH ₂ SCH ₃		S	88-89
91	-COCH ₂ CH ₂ SO ₂ CH ₃		S	139-141
92			S	83-84
93			S	92-93
94			S	56-59
95			S	72-74
96			S	123-125
100			S	156-157
101			S	157-158
102			S	153-155
103			S	102-103
104			S	172-173
106	-CONHOCH ₃		S	124-126
107	-CONHOCH ₂ CH ₃		S	93-94

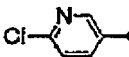
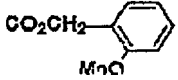
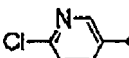
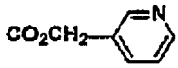
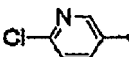
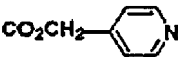
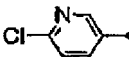
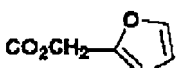
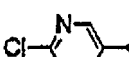
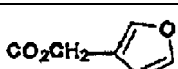
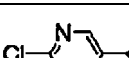
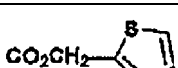
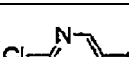
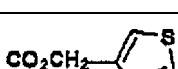
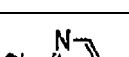
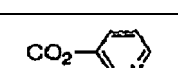
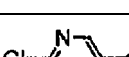
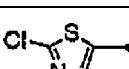
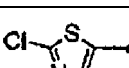
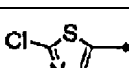
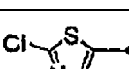
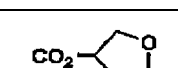
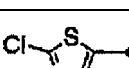
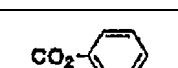
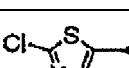
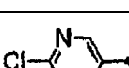
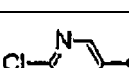
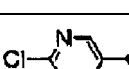
[Table 8]

110	CONHCH_2 		S	
111	$-\text{CONHCH}_2\text{CH}_2\text{OCH}_3$		S	151-152
113	$-\text{CO}_2\text{CH}_2\text{CH}_3$		S	112-114
114	$-\text{CO}_2\text{CH}_2\text{CH}_2\text{F}$		S	144
115	$-\text{CO}_2\text{CH}_2\text{CHF}_2$		S	113
116	$-\text{CO}_2\text{CH}_2\text{CF}_3$		S	88-90
117	$-\text{CO}_2\text{CH}_2\text{CCl}_3$		S	112-114
118	$-\text{CO}_2\text{CH}_2\text{CH}_2\text{OCH}_3$		S	69-71
119	$-\text{CO}_2\text{CH}_2\text{CH}_2\text{SCH}_3$		S	75-76
120	$-\text{CO}_2\text{CH}_2\text{CH}_2\text{Si}(\text{CH}_3)_3$		S	185-186
121	$-\text{CO}_2\text{CH}_2\text{CH}_2\text{OCH}_2\text{CF}_3$		S	48
122	$-\text{CO}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{F}$		S	73
123	$-\text{CO}_2\text{CH}(\text{CH}_2\text{F})_2$		S	138-140
124	$-\text{CO}_2\text{CH}(\text{CF}_3)\text{CH}_3$		S	130-182
125	$-\text{CO}_2\text{CH}(\text{CF}_3)_2$		S	161-163
126	$-\text{CO}_2\text{CH}_2\text{CH}=\text{CH}_2$		S	81-83
127	$-\text{CO}_2\text{CH}_2\text{CCl}=\text{CCl}_2$		S	125-127
128	$-\text{CO}_2(\text{CH}_2)_3\text{CH}_3$		S	95-96

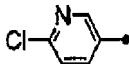
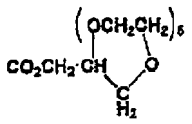
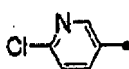
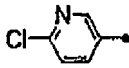
[Table 9]

129	$-\text{CO}_2\text{CH}_2\text{CH}(\text{CH}_3)_2$		S	62-63
130	$-\text{CO}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_3$		S	77-78
131	$-\text{CO}_2\text{C}(\text{CH}_3)_3$		S	166
132	$-\text{CO}_2(\text{CH}_2)_4\text{F}$		S	93-94
133	$-\text{CO}_2(\text{CH}_2)_3\text{CF}_3$		S	87-88
134	$-\text{CO}_2\text{CH}_2\text{CF}_2\text{CF}_2\text{CF}_3$		S	75
135	$-\text{CO}_2\text{CH}_2\text{CH}=\text{C}(\text{CH}_3)_2$		S	69-61
136	$-\text{CO}_2(\text{CH}_2)_4\text{CH}_3$		S	59-61
137	$-\text{CO}_2(\text{CH}_2)_5\text{CH}_3$		S	43
138	$-\text{CO}_2\text{CH}_2\text{C}(\text{CH}_3)_3$		S	162-163
139	CO_2CH_2 		S	117-118
140	CO_2CH_2 		S	70-71
141	CO_2 		s	128
142	CO_2 		S	124-125
143	CO_2 		S	102-103
144	CO_2CH_2 		S	103
145	CO_2CH_2 		S	Oil
146	CO_2 		s	101-102
147	CO_2CH_2 		S	101-102

[Table 10]

148			S	167-168
149			S	98-100
150			S	87-89
151			S	Oil
152			S	77-79
153			S	94-96
154			S	50-52
155			S	90-92
156			S	104-105
158	-CO ₂ CH ₃		S	120-121
159	-CO ₂ CH ₂ CH ₃		S	131-133
160	-CO ₂ CH(CH ₃) ₂		S	102-105
161	-CO ₂ (CH ₂) ₃ CH ₃		S	78-79
162			S	Oil
163			S	127-129
167	-CO ₂ CH(CH ₃) ₂		CH ₂	Oil
168	-COCF ₃		CH ₂	58-59
169	-CO ₂ CH ₃		NH	164-166

[Table 11]

170	-CO ₂ CH ₃		NCH ₃	82-83
171	-CO ₂ CH ₂ C≡CH		S	116-118
172			S	Oil
173	-CO ₂ CH ₂ CH ₂ CH=CH ₂		S	88-90

[Table 12]

Compound No.	IR (KBr, ν , cm^{-1})	NMR (DMSO- d_6 , δ , ppm)
1	1656, 1542, 1462, 1422	3.29(2H,t), 3.75(2H,t), 4.86(2H,s), 7.35(1H,d), 7.68(1H,dd), 8.35 (1H,d)
2	1638, 1536, 1463, 1410	3.18(2H,t), 3.68(2H,q), 3.63(2H,t), 4.82(2H,s), 7.34(1H,d), 7.66(1H, dd), 8.34 (1H,d)
3	1630, 1602, 1527, 1462, 1402	1.23(9H,s), 3.1 (2H,dd), 3.74 (2H,dd), 4.82(2H,s), 7.32(1H,d), 7.67 (1H,dd), 8.38 (1H,d)
4	1640, 1634, 1426, 1402	1.70(3H,s), 3.22(2H,t), 3.67(2H,), 4.84(2H,s), 7.34(1H,d), 7.65(1H d), 8.33(1H d)
5	1657, 1536, 1460, 1402, 888	3.27(2H,dd), 3.74(2H,dd), 4.87(2H,s), 7.84(2H,d), 7.78(1H,dd), 8.40(2H, d)
6	1616, 1522, 1455, 1401	3.19(2H,dd), 3.62(2H,dd), 4.98(2H,s), 7.32(1H,d), 7.43(2H, m), 7.50(1H,m), 7.72(1H,dd), 8.26(2H,m), 8.40(1H,d)
7	1618, 1527, 1457, 1402	3.20(2H,dd), 3.63(2H,dd), 4.97(2H,s), 7.33(1H,d), 7.39(2H, d), 7.68 (1H, dd), 8.19(2H,d), 8.39(1H,d)
8	1620, 1531, 1457, 1408	2.40(3H,s), 3.18(2H,t), 3.60(2H,t), 4.97(2H,s), 7.22(2H,d), 7.82(1H, d), 7.73 (1H,dd), 8.16(2H,d), 8.39(1H,d)
9	1630, 1528, 1460, 1406	1.25-2.00(10H,m), 2.39(1H,m), 8.10(2H,dd), 3.53(2H,dd), 4.82(2H,s), 7.32(2H,d), 7.64(1H,dd), 8.35(2H,d)
10	1726, 1631, 1531, 1397	3.08(2H,dd), 3.51(2H,dd), 3.78(2H,s), 4.71(2H,s), 7.18(1H, d), 7.21-7.32(5H, m), 7.38(1H,dd), 8.24(1H, d)
11	1728, 1626, 1614, 1589, 1529, 1457	3.24(2H,dd), 3.68(2H,dd), 5.03(2H,s), 7.34(1H,d), 7.58(1H, m), 7.74(1H, dd), 7.78-7.81(2H,m), 7.93-8.18(1H,m), 8.42 (1H,d), 8.99(1H,d), 9.73 (1H,d)
12	1587 1551, 1506, 1459, 1425, 1402	3.16(2H,dd), 3.57(2H,dd), 4.94(2H,s), 7.17(1H,m), 7.30(1H, d), 7.42(1H,d), 7.73(1H,dd), 8.04(1H,d), 8.41(1H,d), 10.8(1H,bs)
13	1625, 1524, 1460, 1402, 1282	3.22(2H,t), 3.66(2H,t), 4.98(2H,s), 7.32(1H,d), 7.36(1H,m), 7.71(1H,bd), 8.39(1H,bs), 8.47(1H,m), 8.72(1H,m), 8.98(1H,s)

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(continued)

Compound No.	IR (KBr, ν , cm^{-1})	NMR (DMSO-d_6 , δ , ppm)
14	1717, 1680, 1637, 1530, 1460, 1410	3.25(2H, dd), 3.68(2H, dd), 5.00(2H, s), 7.33(1H, d), 7.75(1H, dd), 8.42(1H, d), 8.68(1H, d), 8.74(1H, d), 8.49(1H, d)

[Table 13]

15	1627, 1528, 1461, 1405, 1352, 1297	3.23(2H, dd), 3.67(2H, dd), 4.96(2H, s), 7.32(1H, d), 7.39(1H, d), 7.67(1H, bd), 8.37(1H, bs), 8.40(1H, d), 9.22(1H, s)
16	1720, 1639, 1523, 1430, 1407, 1384	3.23(2H, dd), 3.73(2H, dd), 4.97(2H, s), 7.34(1H, d), 7.75(1H, dd), 8.42(1H, d), 8.69(1H, s), 9.34(1H, s)
17	1727, 1622, 1530, 1458, 1400	3.26(2H, dd), 3.70(2H, dd), 4.98(2H, s), 7.37(1H, d), 7.65(1H, dd), 7.96(2H, s), 8.36(1H, d)
18	1727, 1626, 1534, 1461, 1406	3.30(2H, dd), 3.82(2H, dd), 5.05(2H, s), 7.52(1H, d), 7.87(1H, dd), 8.19(1H, m), 8.48(1H, d), 9.44(1H, m), 9.74(1H, d)
19	1634, 1561, 1531, 1460, 1412	3.22(2H, dd), 3.69(2H, dd), 4.95(2H, s), 7.35(1H, d), 7.44(1H, d), 7.77(1H, dd), 8.01(1H, dd), 8.17(1H, d), 8.40(1H, d)
20	1726, 1624, 1581, 1462, 1402	8.24(2H, dd), 3.67(2H, dd), 4.98(2H, s), 7.36(1H, d), 7.68(1H, bd), 8.38(1H, bs), 8.59(1H, bs), 8.77(1H, bs), 9.35(1H, bs)
21	1720, 1630, 1532, 1461, 1402	3.24(2H, dd), 3.65(2H, dd), 4.89(2H, s), 7.84(1H, d), 7.69(1H, dd), 7.62(1H, dd), 8.34(1H, d), 8.83(1H, d), 9.18(1H, d)
22	1629, 1524, 1458, 1402	3.24(2H, dd), 3.68(2H, dd), 4.97(2H, s), 7.34(1H, d), 7.67(1H, dd), 8.38(1H, dd), 8.50(1H, d), 9.09(1H, d)
23	1727, 1634, 1566, 1527, 1459, 1402	3.24(2H, dd), 3.68(2H, dd), 4.92(2H, s), 7.31-7.35(2H, m), 7.56(1H, dd), 8.25(1H, d), 8.36(1H, d)
24	1648, 1622, 1529, 1496, 1404	3.25(2H, t), 3.68(2H, t), 4.85(2H, s), 7.35(2H, d), 7.58(1H, dd), 8.34(2H, d)
28	1655, 1611, 1558, 1880, 1312	8.10(2H, t), 3.54(2H, t), 4.75(2H, s), 7.33(2H, d), 7.68(1H, d), 8.35(1H, d)
29	1620, 1611, 1578, 1525, 1459, 1436	2.87(3H, d), 3.09(2H, t), 3.48(2H, t), 4.70(2H, s), 5.23(1H, bs), 7.30(2H, d), 7.60(1H, dd), 8.31(1H, d),
30	1742, 1610, 1578, 1531, 1460, 1290	1.22(3H, t), 3.10(2H, m), 3.56(2H, m), 3.82(2H, s), 4.10(2Hq), 4.74(2H, s), 7.20(1H, bs), 7.40(1H, m), 7.78(1H, m), 8.41(1H, bs)
31	1726, 1700, 1655, 1552, 1475, 1459	3.18(2H, m), 3.71(2H, m), 4.60(2H, s), 4.78(2H, s), 4.78(1H, bs), 7.91(1H, m), 8.51(1H, bs), 10.44(1H, bs)
32	1718, 1549, 1497, 1483, 1469, 1438	3.17(2H, dd), 3.61(2H, dd), 4.79(2H, s), 7.34(1H, m), 7.44-7.46(2H, m), 7.53(1H, m), 7.68(1H, m), 7.83(2H, m), 8.35(1H, bs), 8.65(1H, bs)
33	1661, 1628, 1551, 1449, 1423, 1400	3.07(2H, m), 3.58(2H, m), 4.59(2H, s), 7.18-7.58(5H, m), 7.70(1H, m), 7.85(1H, m), 8.36(1H, bs)

[Table 14]

34	1632, 1615, 1580, 1531, 1457, 1280	1.18(6H,d), 3.08(2H,t), 5.47(2H,t), 4.71(2H,s), 5.14(1H,bs), 7.30(2H,d), 7.60(1H,d), 8.32(1H,d)
35	1634, 1567, 1457, 1418, 1273, 1233	3.10(2H,dd), 3.52(2H,dd), 3.63-3.72(8H,m), 4.75(2H,s), 7.32(1H,d), 7.61(1H,dd), 8.33(1H,d)
36	1648, 1562, 1513, 1466, 1437, 1311	8.13(2H,dd), 3.50(2H,dd), 4.76(2H,s), 5.21(1H,bs), 7.03(1H,m), 7.20(1H,bs), 7.26-7.40(3H,m), 7.50-7.65(3H,m), 8.32(1H,bs)
37	1684, 1628, 1562, 1508, 1460, 1385	1.1-2.1(10H,m), 3.09(2H,m), 3.47(2H,m), 3.66(1H,m), 4.73(2H,s), 5.19(2H,bs), 7.30(1H,m), 7.62(1H,m), 8.34(1H,bs)
42	1662, 1555, 1459, 1440, 1264	3.18(H,dd), 3.60(2H,dd), 3.79(3H,s), 4.81(2H,s), 7.82(1H,d), 7.67(1H,dd), 8.31(1H,d)
43	1564, 1543, 1463, 1429, 1415, 1260	0.97(3H,t), 1.74(2H,tq), 3.15(2H,t), 3.56(2H,t), 4.12(2H,t), 4.81(2H,s), 7.32(1H,d), 7.65(1H,dd), 8.31(2H,d)
44	1652, 1545, 1465, 1415, 1262, 1239	1.32(6H,d), 3.14(2H,t), 3.54(2H,t), 4.82(2H,s), 5.01(1H,sep), 7.35(1H,d), 7.62(1H,dd), 8.2.31(1H,d)
45	1657, 1563, 1434, 1265	3.18(2H,dd), 3.60(2H,dd), 3.75(2H,t), 4.41(2H,t), 4.82(2H,s), 7.32(1H,bs), 7.65(1H,dd), 8.32(1H,bs)
46	1657, 1572, 1558, 1460, 1436, 1420,	3.16(2H,dd), 3.39(3H,s), 3.57(2H,dd), 3.67(2H,t), 4.32(2H,t), 4.81(2H,s), 7.32(1H,d), 7.65(4H,dd), 8.30(1H,d)
47	1686, 1664, 1541, 1470, 1455, 1428	3.15(2H,t), 3.55(2H,t), 4.78(2H,s), 5.20(2H,s), 7.26-7.43(6H,m), 7.63(1H,dd), 8.29(1H,d)
48	1692, 1551, 1214	3.20(2H,t), 3.64(2H,t), 4.83(2H,s), 7.14(2H,d), 7.31-7.36(3H,m), 7.66(1H,dd), 8.35(1H,d)
49	1684, 1548, 1420, 1213, 1141	3.20(2H,dd), 3.64(2H,dd), 4.80(2H,s), 7.16(1H,m), 7.24-7.34(3H,m), 7.42(1H,m), 7.65(1H,m), 8.31(1H,d)
50	1730, 1717, 1551, 1420, 1221	2.53(2H,m), 2.76(4H,s), 3.85(2H,m), 4.81(2H,s), 7.53(1H,d), 7.78(1H,dd), 8.39(1H,bs)
51	1637, 1570, 1460, 1445, 1423, 1265	3.19(2H,dd), 8.62(2H,dd), 4.84(2H,s), 7.19-7.25(3H,m), 7.34-7.40(1H,m), 7.64(1H,dd), 8.33(1H,d)
52	1656, 1552, 1464, 1446, 1426, 1266	0.88(3H,t), 1.30-1.40(6H,m), 1.68-1.72(2H,m), 3.15(2H,t), 3.56(2H,t), 4.16(2H,t), 4.81(2H,s), 7.32(1H,d), 7.64(1H,dd), 8.31(1H,t)
53	1664, 1546, 1271, 1232, 1208, 1157	3.25(2H,dd), 3.74(2H,dd), 4.80(2H,s), 7.29-7.35(2H,m), 7.46-7.55(2H,m), 7.61-7.70(2H,m), 7.84-7.88(3H,m), 8.37(1H,bs)

[Table 15]

56	1656,1535,1524,1408	3.29(2H,t),3.79 (2H,t),4.96(2H,s),7.53(1H,s)
57	1617,1522,1445,1407	3.20(2H,dd),3.68(2H,dd),5.02(2H,s),7.40-7.54(3H,m),7.51(1H,s),8.32(2H,m)
58	1617,1536,1401,1380	3.25(2H,dd),3.64(2H,t),4.85(2H,s),7.34(1H,d),7.69(1H,dd),8.33(1H,d),8.97(1Hs)
59	1687,1639,1593,1459,1386	2.17(2H,m),3.06(4H,m),3.60(2H,t),4.82(2H,s),7.37(1H,d),7.72(1H,dd),8.30(1H,s),8.35(1H,d)
60	1730,1690,1567,1462,1388,1371, 1286	3.39(2H,t),3.69(2H,t),4.58(2H,s),7.33(1H,d),7.67(1H,d),8.32(1H,s),8.58(1Hs),8.61(1H, bs)
61	1620,1520,1411,1383	3.25(2H,t),3.69(2H, dd),4.89(2H,s),7.48(1H,s),8.97(1H,s)
62	1684,1600,1416,1352	2.18(2H,m),3.06(2H m),3.67(2H,t),5.07 2H, s),7.63(1H,s),8.36(1H, s)
63	1729,1650,1555,1479,1469	2.14(2H,m),3.29(2H,t),3.71(2H,t),4.84(2H,s),7.33(1H,d),7.73(1H,d d),8.39(1H,d),8.66(1H,d),8.71(1H,bs),9.47(1H,s)
64	1637,1560,1489,1460,1318	3.14(2H,m),3.27(2H,t),3.44(2H,t),4.74(2H,s),7.32(1H,d),7.56(1H,d),7.62(1H,dd),8.32(1H,d),8.79(1H,d),9.11(1H,s)
65	1627,1606,1576,1551,1469	3.47(2H,t),3.78(2H,dd),4.72(2H,s),7.33(1H,d),7.72(1H,bs),8.38(1H ,bs),8.64(1H,d),8.65(1H,bs),8.71(1H,d),8.91(1H,bs),9.52(1H,s)
66	1611,1567,1463,1380,1347	8.45(2H,t),3.76(2H,t),4.62(2H,s),7.33(1H,d),7.56(1H,d),7.64(1H,d d),8.32(1H,d),8.71(1H,bs),8.77(1H,d),9.12(1H,s)
67	1639,1531,1412,1305	3.23(2H,t),3.62(2H,t),4.04(1H, m) 4.83(2H, s),7.34 (1H,d),7.65(1 H,dd),8.33(1H,d)
70	1632,1525,1463,1401	1.31-2.05(4H,m),2.90(1H,m),3.08-3.28(3H,m),3.28(1H,m),3.54(2H ,t),4.82(2H,s),5.90(1H,m),6.15(1H,m),7.33(1H,m),7.66(1H,m),8.36 (1H,m)
71	2903,2848,1637,1525,1455	1.68-2.04(15H,m),3.11(2H,dd),3.54(2H,dd),4.83(2H,s),7.32(1H,d),7.67(1H,dd),8.37(1H,d)

[Table 16]

Compound No.	IR (KBr, ν , cm^{-1})	NMR (DMSO- d_6 , δ , ppm)
72	1627, 1525, 1396, 1270	2.23(3H,t), 3.11(2H,t), 3.51(2H,t), 4.79(2H,s), 7.30(1H,d), 7.63(1H,dd), 8.32(1H,d)
73	1644, 1533, 1419, 793	8.18(2H,t), 3.64(2H,t), 4.21(2H,s), 4.83(2H,s), 7.33(1H,d), 7.69(1H,bs), 8.35(1H,bs)
74	1646, 1584, 1427, 818	3.23(2H,dd), 3.70(2H,dd), 4.86(2H,s), 6.06(1H,s), 7.34(1H,d), 7.76(1H,dd), 8.39(1H,d)
75	1640, 1532, 1416, 1290, 1260, 1183	3.17(2H,t), 3.63(2H,t), 4.02(2H,s), 4.83(2H,s), 7.34(2H,d), 7.32(1H,d), 7.72(1H,dd), 8.36(1H,d)
76	1654, 1548, 1431, 1241, 1109	3.24(2H,t), 3.68(2H,t), 4.87(2H,s), 5.94(1H,t), 7.35(2H,d), 7.68(1H,bs), 8.34(1H,bs)
77	1660, 1541, 1418, 1141, 1112, 959	3.28(2H,t), 3.75(2H,t), 4.86(2H,s), 7.34(2H,d), 7.73(1H,dd), 8.38(2H,d)
78	1653, 1538, 1456, 1423, 1238	3.15(2H,t), 3.48(3H,s), 3.57(2H,t), 3.78(2H,s), 4.16(2H,s), 4.80(2H,s), 7.33(1H,d), 7.63(1H,d), 8.24(1H,s)
79	2361, 1642, 1529, 1409, 1389	3.21(2H,t), 3.52(2H,s), 3.67(2H,t), 4.85(2H,s), 7.35(1H,d), 7.39(1H,bs), 7.67(1H,bs), 8.87(1H,bs)
80	1662, 1542, 1417, 1239, 754	3.13(2H,t), 3.56(2H,t), 4.73(2H,s), 4.74(2H,s), 6.90-6.95(3H,m), 7.22-7.27(3H,m), 7.45(1H,dd), 8.27(1H,d)
81	1726, 1635, 1531, 1460, 1406, 1289	2.75(2H,t), 3.12(2H,t), 3.35(3H,s), 3.56(2H,t), 3.74(2H,t), 4.82(2H,s), 7.31(1H,m), 7.68(1H,dd), 8.34(1H,d)
82	2152, 1615, 1530, 1429, 1191, 1129	3.21(2H,t), 3.65(2H,t), 3.95(2H,s), 4.86(2H,s), 7.35(1H,d), 7.67(1H,dd), 8.35(1H,d)
83	1640, 1592, 1537, 1406, 1241	3.16(2H,t), 3.59(2H,t), 4.87(2H,s), 5.79(1H,dd), 6.40(1H,dd), 6.47(1H,dd), 7.33(1H,d), 7.68(1H,bs), 8.37(1H,bs)
84	1626, 1528, 1462, 1410, 1386, 1242	3.22(2H,t), 3.67(2H,t), 4.86(2H,s), 7.34(1H,d), 7.68(1H,dd), 8.38(1H,d)
85	1634, 1530, 1464, 1437, 1403, 1233	1.69(3H,d), 3.11(2H,t), 3.19(2H,m), 3.54(2H,t), 4.80(2H,s), 5.56(1H,m), 5.67(1H,m), 7.31(1H,d), 7.67(1H,dd), 8.34(1H,d)
86	2235, 1598, 1530, 1413, 1279, 1248	2.01(3H,s), 3.17(2H,t), 3.62(2H,t), 4.86(2H,bs), 7.34(1H,d), 7.69(1H,dd), 8.35(1H,d)

[Table 17]

87	1640, 1543, 1462, 1402	1.70(3H,s), 3.22(2H,t), 3.67(2H,t), 4.84(2H,s), 7.34(1H,d), 7.65(1H, dd), 8.33(1H,d)
88	1733, 1634, 1540, 1460, 1411, 1390	1.38(3H,t), 3.22(2H,t), 3.69(2H,t), 4.33(2H,q), 4.89(2H,s), 7.34(1H,d), 7.77(1H,dd), 8.37(1H,d)
89	1628, 1530, 1460, 1406, 1281, 1246	2.19(3H,s), 3.16(2H,t), 3.36(2H,s), 3.60-(2H,t), 4.83(2H,s), 7.32(1H,d), 7.74(1H,dd), 8.36(1H,d)
90	1635, 1533, 1401, 1243, 1130, 1104	2.13(3H,s), 2.80(2H,m), 3.13(2H,t), 3.55(2H,t), 4.82(2H,t), 4.82(2H,s), 7.32(1H,d), 7.65(1H,dd), 8.34(1H,d)
91	1651, 1557, 1456, 1442, 1289, 1237	3.01(3H,s), 3.21(2H,t), 3.39(2H,t), 3.61(2H,t), 4.57(2H,t), 4.78(2H,s), 7.33(1H,d), 7.63(1H,dd), 8.31(1H,d)
92	1619, 1535, 1458, 1405	0.85(2H,m), 1.05(2H,m), 1.84(1H,m), 3.11(2H,t), 3.54(2H,t), 4.81(2H,s), 7.33(1H, d), 7.65(1H,dd), 8.34(1H,d)
93	1620, 1532, 1460, 1407, 1125	0.70(2H,m), 1.30(2H,m), 1.35(3H,s), 3.10(2H,t), 3.53(2H,t), 4.75(2H,s), 7.32(1H,d), 7.65(1H,dd), 8.32(1H,d)
94	1622, 1532, 1460, 1407	0.68(1H,m), 1.12(3H,d), 1.24(1H,m), 1.45(1H,m), 1.58(1H,m), 3.09(2H,t), 3.52(2H,t), 4.78(1H,d), 4.82(1H,d), 7.33(1H,d), 7.64(1H,dd), 8.34(1H,d)
95	1629, 1529, 1411, 1238, 1222	2.14(1H,m), 2.28(1H,m), 3.15(2H,t), 3.24(1H,m), 3.56(2H,t), 3.84(1H, m), 3.89(1H,m), 3.98(1H,m), 4.02(1H,m), 4.82(2H,s), 7.34(1H,d), 7.64(1H,dd), 8.34(1H,d)
96	1627, 1531, 1468, 1417, 1292, 1098	3.18(2H,t), 3.62(2H,t), 4.90(2H,s), 6.50(1H,dd), 7.21(1H,d), 7.32(1H, d), 7.58(1H,bs), 7.40(1H,dd), 8.40(1H, d)
100	1628, 1537, 1458, 1396, 1236, 1189	3.11(2H,t), 3.54(2H,t), 3.77(2H,s), 4.73(2H,s), 7.23(2H,m), 7.43(1H,m) , 7.64(1H,bd), 8.26(1H,bs), 8.47(1H,bs), 8.55(1H,bs)
101	1682, 1618, 1592, 1518, 1505, 1335	2.25(3H,s), 3.22(2H,t), 3.65(2H,t), 4.96(2H,s), 7.05(1H,m), 7.33(1H,d) , 7.49(1H,m), 7.69(1H,m), 8.38(1H,d), 8.43(1H,m), 8.70(1Hm), 12.0(1H ,bs)
102	1683, 1609, 1632, 1536, 1506, 1281	2.22(3H,s), 2.29(3H,s), 3.19(2H,t), 3.62(2H,t), 4.92(2H,s), 7.13(1H,t), 7.32(1H,d), 7.36(1H,d), 7.69(1H,m), 8.09(1H,d), 8.36(1H,bs), 10.1(1H

[Table 18]

103	1626, 1536, 1460, 1427, 1371, 1129	1.69(1H,m), 2.15(1H,m), 2.66(1H,m), 3.15(2H,m), 3.57(2H,m), 4.73(1H,d), 4.91(1H,d), 7.34(1H,d), 7.64(1H,dd), 8.34(1H, d)
104	2959, 2930, 1729, 1536, 1287, 1181	3.17(2H,t), 3.60(2H,t), 4.68(2H,s), 4.77(2H,s), 6.98(1H,s), 7.08(1H,s), 7.31(1H,d), 7.42(1H,dd), 7.53(1H,s), 8.26(1H,d)
106	3222, 1620, 1568, 1439, 1238, 1100	3.14(2H,t), 3.54(2H,t), 3.79(3H,s), 4.71(2H,s), 7.32(1H,d), 7.61(1H,dd), 7.81(1H,bs), 8.32(1H,d)
107	3435, 1632, 1565, 1272, 1109, 1085	1.27(3H,t), 3.13(2H,t), 3.54(2H,t), 3.99(2H,q), 4.71(2H,s), 7.31(1H,d), 7.63(1H,dd), 7.85(1H,bs), 8.31(1H,d)
110	3298, 1608, 1570, 1508, 1436, 1239	3.10(2H,t), 3.48(2H,t), 4.60(2H,d), 4.74(2H,s), 6.24(1H,bs), 7.16(1H, m), 7.80-7.35(2H,m), 7.65(1H,m), 8.32(1H,bs), 8.54(1H,bd)
111	3306, 1607, 1578, 1531, 1469, 1279	3.08(2H,t), 3.37(3H,s), 3.50(6H,m), 4.71(2H,s), 5.58(1H,bs), 7.31(1H, d), 7.62(1H,dd), 8.31(1H,d)
113	1664, 1548, 1460, 1416, 1264, 1240	1.34(3H,t), 3.16(2H,t), 3.57(2H,t), 4.22(2H,q), 4.81(2H,s), 7.32(1H,d), 7.65(1H, dd), 8.31(1H,d)
114	1641, 1557, 1405, 1322, 1286, 1143	3.18(2H,t), 3.59(2H,t), 4.41(2H,dt), 4.67(2H,dt), 4.82(2H,s), 7.33(1H, d), 7.65(1H,dd), 8.32(1H,d)
115	1662, 1551, 1462, 1428, 1388, 1267	8.20(2H,t), 3.63(2H,t), 4.35(2H,td), 4.81(2H,s), 6.02(1H,tt), 7.33(1H, d), 7.64(1H,bd), 8.32(1H,bs)
116	1678, 1547, 1459, 1291, 1237, 1141	3.21(2H,t), 3.63(2H,t), 4.55(2H,q), 4.82(2H,s), 7.34(1H,d), 7.66(1H, bd), 8.33(1H,bs)
117	1668, 1536, 1460, 1443, 1429, 1231	3.21(2H,t), 3.63(2H,t), 4.55(2H,q), 4.85(2H,s), 7.34(1H,d), 7.68(1H,d d), 8.33(1H,d)
118	1657, 1572, 1558, 1460, 1436, 1260	3.16(2H,dd), 3.39(3H,s), 3.57(2H,dd), 3.67(2H,t), 4.32(2H,t), 4.81(2H, s), 7.32(1H,d), 7.65(1H,dd), 8.30(1H,d)
119	1660, 1551, 1459, 1435, 1264, 1240	2.17(3H,s), 2.81(2H,t), 3.17(2H,t), 3.39(3H,s), 3.58(2H,t), 4.32(2H,t), 4.81(2H,s), 7.32(1H,d), 7.65(1H,dd), 8.31(1H,d)

[Table 19]

120	2923,2582,1742,1663,1655,1263	0.06(9H,s),1.05(2H,t),3.11(2H,t),3.52(2H,t),4.21(2H,t),4.76(2H,s),7.26(1H,d),7.61(1H,bd),8.26(1H,bs)
121	1667,1553,1459,1421,1265,1240	3.18(2H,t),3.58(2H,t),3.91(4H,overlap),4.34(2H,t),4.81(2H,s),7.32 (1H,d)7.66(1H, bd),8.31(1H,bs)
122	1667,1543,1465,1429,1414,1260	2.10(2H,dt),3.17(2H,dd),3.59(2H,dd),4.29(2H,t),4.57(2H,dt),4.81 (2H,s),7.32(1H,d),7.65(1H,dd),8.31(1H,d)
123	1654,1558,1542,1457,1419,1260	3.17(2H,dd),3.59(2H,dd),4.61(2H,bs),4.71(2H,bs),4.81(2H,s),5.20(1H,m),7.32 (1H, d),7.64(1H,bd),8.31(1H,bs)
124	1668,1543,1466,1428,1414,1235	1.46(3H,d),3.19(2H,t),3.61(2H,t),4.80(1H,d),4.84(1H,d),5.31(1H,m),7.33(1H,d),7.65(1H,bd),8.32(1H,bs)
125	1698,1546,1552,1428,1348,1198	3.25(2H,dd),3.68(2H,dd),4.84(2H,s),5.31(1H,m),7.36(1H,d),7.66 (1H,bd),8.33(1H,bs)
126	1655,1545,1465,1431,1415,1210	3.17(2H,t),3.58(2H,t),4.67(2H,dt),4.81(2H,s),5.25(1H,dd),5.37(1H, ddd),6.02(1H,m),7.32 (1H,d),7.65(1H,dd),8.31 (1H,d)
127	1660,1543,1460,1442,1416,1204	3.19(2H,t),3.61(2H,t),4.83(2H,s),5.03(2H,s),7.33(1H,d),7.65(1H, dd),8.31(1H,d)
128	1659,1560,1456,1435,1389,1157	0.94(3H,t),1.41(2H,m),1.70(2H,m),3.15(2H,dd),3.56(2H,dd),4.17(2H,t),4.81(2H,s),7.32(1H,d),7.65(1H,bd),8.31(1H,bs)
129	1657,1550,1462,1419,1264,1146	0.96(6H,d),3.15(2H,t),3.56(2H,t),3.95(2H,d),4.82(2H,s),7.32(1H,d),7.65(1H,dd),8.31(1H,d)
130	1653,1542,1464,1428,1237,1144	0.94(3H,t),1.29(2H,d),1.56-1.78(2H,m),3.15(2H,t),3.54(2H,t),4.78 -4.86(1H,m),4.82(2H,s),7.32(1H,d),7.63(1H,dd),8.31(1H,d)
131	1653,1560,1464,1419,1280,1105	1.54(9H,s),3.15(2H,t),3.51(2H,t),4.81(2H,s),7.32(1H,d),7.63(1H,bd),8.31(1H,bs)
132	1660,1559,1456,1435,1389,1155	1.85(4H,m),3.17(2H,t),3.58(2H,d),4.21(2H,m),4.47(2H,bd),4.82(2 H,s),7.32(1H,d),7.65(1H,bd),8.32(1H,bs)
133	1662,1552,1459,1422,1254,1148	1.98(2H,m),2.23(2H,m),3.18(2H,t),3.59(2H,t),4.23(2H,t),4.84(2H,s),7.33(1H,d),7.65(1H,dd),8.32(1H, d)
134	1680,1552,1459,1232,1148,1103	3.21(2H,t),3.64(2H,t),4.68(2H,m),4.82(2H,s),7.33(1H,d),7.67(1H,b d),8.33(1H,bs)
135	1660,1571,1557,1458,1260,1102	1.73(3H,s),1.75(3H,t),3.16(2H,t),3.56(2H,t),4.67(2H,d),4.81(2H,s),5.44(1H,m),7.32(1H,d),7.65(1H,dd),8.30(1H,d)

[Table 20]

136	1660, 1561, 1456, 1436, 1330, 1155	0.90(3H,t), 1.36(4H,m), 1.72(2H,m), 3.15(2H,dd), 3.56(2H,dd), 4.16(2H,t), 4.81(2H,s), 7.32(1H,d), 7.64(1H,dd), 8.31(1H,d)
137	1856, 1552, 1464, 1426, 1266, 1234	0.88(3H,t), 1.30-1.40(6H,m), 1.68-1.72(2H,m), 3.15(2H,t), 3.56(2H,t), 4.16(2H,t), 4.81(2H,s), 7.32(1H,d), 7.64(1H,dd), 8.31(1H,d)
138	1661, 1546, 1458, 1432, 1271, 1150	0.98(9H,s), 3.15(2H,dd), 3.56(2H,dd), 3.90(2H,s), 4.82(2H,s), 7.32(1H,d), 7.68(1H,dd), 8.32(1H,d)
139	1662, 1550, 1459, 1431, 1263, 1145	0.31(2H,m), 0.58(2H,m), 1.24(1H,m), 3.16(2H,t), 3.56(2H,t), 4.00(2H,t), 4.83(2H,s), 7.32(1H,d), 7.64(1H,dd), 8.32(1H,bs)
140	1666, 1550, 1458, 1446, 1265, 1146	2.68(1H,m), 2.84(1H,m), 3.18(2H,dd), 8.30(1H,m), 3.56(2H,dd), 4.10(1H,m), 4.38(1H,m), 4.80(2H,s), 7.33(1H,d), 7.66(1H,dd), 8.32(1H,d)
141	1664, 1548, 1459, 1418, 1268, 1143	1.61(1H,m), 1.78(1H,m), 2.19(2H,m), 2.39(2H,m), 3.15(2H,t), 3.56(2H,t), 4.82(2H,s), 5.02(1H,m), 7.32(1H,d), 7.64(1H,dd), 8.32(1H,bs)
142	1662, 1542, 1507, 1458, 1419, 1233	3.17(2H,t), 3.60(2H,t), 4.77(2H,m), 4.82(2H,s), 4.91(2H,m), 5.46(1H,t), 7.34(1H,d), 7.65(1H,dd), 8.33(1H,d)
143	1650, 1547, 1463, 1430, 1263, 1238	1.58(2H,m), 1.72-1.85(4H,m), 1.93(2H,m), 3.14(2H,t), 3.54(2H,t), 4.81(2H,s), 5.16(1H,m), 7.32(1H,d), 7.64(1H,dd), 8.32(1H,d)
144	1665, 1550, 1462, 1420, 1212, 1068	1.68(1H,m), 2.06(1H,m), 2.69(1H,m), 3.17(2H,dd), 3.58(2H,dd), 3.63(1H,m), 3.75(1H,m), 3.87(2H,m), 4.09(1H,m), 4.16(1H,m), 4.81(2H,s), 7.33(1H,d), 7.64(1H,dd), 8.32(1H,d)
145	1665, 1552, 1460, 1444, 1238, 1151	1.35(3H,s), 1.42(3H,s), 3.18(2H,dd), 3.62(2H,dd), 3.79(1H,m), 4.10(1H,m), 4.20(2H,d), 4.40(1H,m), 4.81(2H,s), 7.33(1H,d), 7.64(1H,dd), 8.32(1H,bs)
146	1660, 1546, 1458, 1264, 1237, 1150	1.78(2H,m), 1.99(2H,m), 3.16(2H,t), 3.52(2H,m), 3.57(2H,t), 3.99(2H,m), 4.82(2H,s), 4.89(1H,m), 7.32(1H,d), 7.64(1H,dd), 8.32(1H,d)
147	1661, 1543, 1423, 1236, 1144, 1054	1.25-1.65(5H,m), 1.85(1H,m), 3.15(2H,t), 3.46(1H,m), 3.64(2H,t), 3.65(1H,m), 3.99(1H,m), 4.12(2H,d), 4.82(2H,s), 7.32(1H,d), 7.65(1H,dd), 8.30(1H,d)
148	1698, 1522, 1322, 1398, 1143, 1110	3.29(2H,t), 3.78(2H,t), 4.79(2H,s), 7.41(2H,d), 7.47(1H,dd), 8.25(2H,d), 8.35(1H,d)
149	1654, 1550, 1444, 1421, 1291, 1154	3.15(2H,dd), 3.54(2H,dd), 3.82(3H,s), 4.80(2H,s), 5.27(2H,s), 6.86(1H,d), 6.93(1H,m), 7.27(1H,m), 7.30(1H,d), 7.39(1H,d), 7.63(1H,dd), 8.29(1H,d)

5			
10			
15			
20			
25			3.18(2H,t), 3.58(2H,t), 4.80(2H,s), 5.22(2H,s), 7.27(1H,m), 7.32(1H,d), 7.62(1H,dd), 7.78(1H,m), 8.31(1H,d), 8.56(1H,m), 8.69(1H,s)
30			
35			
40		1654, 1544, 1469, 1439, 1287, 1131	
45			
50	150		
55			

[Table 21]

151		3.18(2H,t),3.52(2H,t),4.59(2H,s),5.22(2H,s),7.30(1H,d),7.45(2H,dd),8.32(1H,d),8.64(2H,dd)
152	1669,1558,1457,1420,1260,1039	3.16(2H,dd),3.56(2H,dd),4.79(2H,s),5.15(2H,s),6.34(1H,dd),6.44(1H,d),7.31(1H,d),7.40(1H,d),7.63(1H,dd),8.29(1H,d)
153	1657,1557,1457,1428,1240,1146	3.16(2H,t),3.57(2H,t),4.79(2H,s),5.01(2H,s),6.50(1H,s),7.31(1H,d),7.38(1H,bs),7.52(1H,bs),7.62(1H,dd),8.30(1H,d)
154	1660,1551,1458,1440,1237,1146	3.15(2H,t),3.55(2H,t),4.77(2H,s),5.34(2H,s),6.96(1H,t),7.13-7.80(3H,overlap),7.62(1H,dd),8.28(1H,d)
155	1650,1546,1460,1440,1263,1146 -	3.16(2H,t),3.57(2H,t),4.79(2H,s),5.20(2H,s),7.16(1H,d),7.27-7.32(3H,overlap),7.62(1H,dd),8.30(1H,d)
156	1682,1548,1461,1418,1240,1104	3.22(2H,dd),3.67(2H,dd),4.84(2H,s),7.30-7.34(1H,m),7.36(1H,d),7.57(1H,m),7.67(1H,dd),8.34(1H,d),8.46(1H,m),8.53(1H,m)
158	1667,1547,1528,1443,1256,1128	3.19(2H,t),3.66(2H,t),3.81(3H,s),4.85(2H,s),7.46(1H,s)
169	1664,1552,1529,1432,1237,1192	1.36(3H,t),3.18(2H,t),3.63(2H,t),4.24(2H,q),4.86(2H,s),7.46(1H,s)
160	1667,1550,1530,1418,1250,1048	1.33(6H,d),3.16(2H,t),3.62(2H,t),4.87(2H,s),5.03(1H,e),7.45(1H,s)
161	1661,1550,1528,1446,1238,1186	0.95(3H,t),1.44(2H,m),1.72(2H,m),3.18(2H,t),3.64(2H,t),4.19(2H,q),4.86(2H,s),7.46(1H,s)
162	1661,1549,1527,1429,1252,1052	1.70(2H,m),2.07(1H,m),2.71(1H,m),3.18(2H,dd),3.65(2H,dd),3.78(1H,m),3.88(2H,m),4.11(1H,m),4.17(1H,m),4.86(2H,s),7.45(1H,s)
163	1680,1550,1528,1412,1248,1183,	3.19(2H,dd),3.68(2H,dd),4.89(2H,s),7.21(3H,overlap),7.37(2H,m),7.47(1H,s)
167	1671,1587,1495,1460,1252,1081	1.31(6H,d),2.05(2H,m),3.10(2H,t),3.32(2H,m),4.69(2H,s),4.94(1H,sep),7.31(1H,d),7.63(1H,dd),8.30(1H,d)
168	1667,1566,1498,1461,1391,1183	2.14(2H,m),3.22(2H,m),3.53(2H,m),4.74(2Hs),7.35(1H,d),7.72(1H,dd),8.36(1H,d)
169	8370,1730,1648,1599,1460,1138	3.35(2H,dd),3.64(2H,dd),3.71(8H,s),4.55(2H,s),7.31(1H,d),7.65(1H,dd),7.94(1H,bs),8.29(1H,d)

[Table 22]

170	1655, 1522, 1459, 1389, 1294, 1081	2.92(3H,s), 3.34(2H,dd), 3.55(2H,dd), 3.70(3H,s), 4.54(2H,s), 7.31(1H,d), 7.70(1H,dd), 8.28(1H,d)
171	1658, 1550, 1421, 1265, 1240, 1144	3.07(2H,m), 3.58(2H,m), 4.59(2H,s), 7.18-7.58(5H,m), 7.70(1H,m), 7.78(1H,m), 8.36(1H,bs)
172	2904, 1663, 1459, 1352, 1239, 1064	3.16(2H,dd), 3.57(2H,dd), 3.63-3.90(23H,m), 4.24(2H,d), 4.80(2H,s), 7.32(1H,d), 7.40(1H,d), 7.65(1H,dd), 8.31(1H,d)
173	1659, 1543, 1459, 1415, 1261, 1146	2.49(2H,m), 3.16(2H,t), 3.56(2H,t), 4.22(2H,t), 4.80(2H,s), 5.14(2H,m), 5.84(1H,m), 7.32(1H,d), 7.64(1H,dd), 8.31(1H,d)

<Production Example 1>

3-(2-Chloro-5-pyridinylmethyl)-2-(benzoylimino)-thiazolidine ("Table 2", Compound No.6)

[0091] 3-(2-Chloro-5-pyridinylmethyl)-2-iminothiazolidine (227 mg, 1.1 mmol) and triethylamine (110 mg, 1.1 mmol) were dissolved in 10ml of acetonitrile, into which, benzoyl chloride (141 mg, 1.0 mmol) dissolved in acetonitrile 5ml was added under cooling with ice, and stirred at room temperature overnight. After distilling the solvent off under reduced pressure followed by extraction with ethyl acetate, the ethyl acetate layer was washed with 2N-hydrochloric acid, saturated aqueous sodium bicarbonate, and water in this order, and the ethyl acetate layer was dehydrated and dried, and then the ethyl acetate layer was distilled off, and remaining oil was purified by a silica gel column chromatography to obtain the intended substance. The crude crystal of the intended substance was washed with hexane. Yield: 160 mg (65%). Melting point: 133°C.

<Production Example 2>

3-(2-Chloro-5-pyridinylmethyl)-2-(phenylacetylimino)-thiazolidine ("Table 2", Compound No.10)

[0092] To a solution of 3-(2-Chloro-5-pyridinylmethyl)-2-iminothiazolidine (132 mg, 0.5 mmol) and l'ethyl-3-(3'dimethylaminopropyl)carbodiimide hydrochloride (96 mg, 0.5 mmol) and 4-dimethylaminopyridine (65 mg, 0.54 mmol) in 3 ml of dichloromethane, phenylacetic acid (68 mg, 0.5 mmol) was added in portions, and stirred at room temperature overnight. After distilling the solvent off, the resultant crude crystal was washed with 1% hydrochloric acid, recrystallized from ethyl acetate thereby obtaining the intended substance. Yield: 100 mg (58%). Melting point: 157-158°C.

<Production Example 3>

3-(2-Chloro-5-thiazolylmethyl)-2-(benzoylimino)-thiazolidine ("Table 4", Compound No.57)

[0093] 3-(2-Chloro-5-thiazolylmethyl)-2-iminothiazolidine (233 mg, 1.0 mmol) and sodium carbonate (120 mg, 1.1 mmol) were dissolved in 10 ml of water, into which, benzoyl chloride (141 mg, 1.0 mmol) dissolved in 5ml of chloroform was added under cooling with ice, and thereafter stirred at room temperature for 2 hours. The chloroform layer was washed successively with 1% hydrochloric acid and water, dehydrated and dried, chloroform was distilled off, and remaining oil was purified by a silica gel column chromatography to obtain the intended substance. The crude crystal of the intended substance was recrystallized from ethyl acetate. Yield: 101 mg (30%). Melting point: 120°C.

<Production Example 5>

[[3-(2-Chloro-5-pyridinylmethyl)]-2-(3H)-thiazolidinilidene] urea("Table 3", Compound No.28)

[0094] 3-(2-Chloro-5-pyridinylmethyl)-2-iminothiazolidine (227 mg, 1.1 mmol) and trimethyl isocyanate (110 mg, 1.1 mmol) were dissolved in 20 ml of acetonitrile, into which, several drops of triethylamine were added, and heated under reflux for 9 hours with stirring. After distilling the solvent off, the residue was combined with 5 ml of 2N hydrochloric acid, and allowed to stand overnight. The precipitating intended substance was recovered by filtration, and recrystallized from ethyl acetate. Yield: 104 mg (39%). Melting point: 158-159°C.

<Production Example 6>

3-(2-Chloro-5-pyridinylmethyl)-2-(methoxycarbonylimino)-thiazolidine ("Table 3", Compound No.42)

[0095] 3-(2-Chloro-5-pyridinylmethyl)-2-iminothiazolidine (227 mg, 1.1 mmol) and methyl chloroformate (165 mg, 1.2 mmol) were dissolved in 20 ml of acetonitrile, into which, potassium carbonate (114 mg, 1 mmol) was added in portions and stirred at room temperature overnight. The reaction solution was made free of solids, and the solids were washed with ethyl acetate, and from the wash the solvent was distilled off, and the resultant intended substance was recrystallized from ethyl acetate. Yield: 120 mg (42%). Melting point: 128-130°C.

<Production Example 7>

3-(2-Chloro-5-pyridinylmethyl)-2-(formylimino)-thiazolidine ("Table 4", Compound No.58)

5 **[0096]** A mixture of 3-(2-Chloro-5-pyridinylmethyl)-2-iminothiazolidine (227 mg, 1.1 mmol) and 10 ml of ethyl formate was heated at the boiling point of ethyl formate for 5 hours with stirring. After distilling the excessive ethyl formate off followed by purification by a silica gel column chromatography, the resultant intended solid was washed with diethyl ether. Yield: 32 mg (13%). Melting point: 99-100°C.

10 <Formulation Example 1>

1. Powder formulation

15 **[0097]** 3 Parts by weight of the imino derivative of Compound No.1 (Table 1), 40 parts by weight of a clay, 57 parts by weight of a talc were ground respectively and mixed to obtain a powder formulation.

<Formulation Example 2>

20 2. Wettable powder

[0098] 50 Parts by weight of the imino derivative of Compound No.6 (Table 1), 5 parts by weight of lignin sulfonate, 3 parts by weight of an alkyl sulfonate, 42 parts by weight of a kieselguhr were ground respectively and mixed to obtain a wettable powder.

25 <Formulation Example 3>

3. Granule formulation

30 **[0099]** 5 Parts by weight of the imino derivative of Compound No.21 (Table 3), 43 parts by weight of a bentonite, 45 parts by weight of a clay, 7 parts by weight of lignin sulfonate were mixed uniformly, combined with water and kneaded, and then subjected to an extruding granulator to fabricate into granules, which was dried to obtain a granule formulation.

<Formulation Example 4>

35 4. Emulsifiable concentrate

[0100] 20 Parts by weight of the imino derivative of Compound No.71 (Table 5), 10 parts by weight of a polyoxyethylene alkylalyl ether, 3 parts by weight of a polyoxyethylene sorbitan monolaurate, 67 parts by weight of a xylene were mixed uniformly while dissolving to obtain an emulsifiable concentrate.

40

<Experimental Example 1>

1. Cotton aphid controlling effect

45 **[0101]** The insecticides formulated in Production Examples were used under a practical condition to validate the pest controlling effects.

[0102] A cucumber cotyledon was dipped in a 100 µg/ml treatment solution obtained by a 5000-fold dilution with water of the wettable powder prepared in accordance with Formulation Example 2. This cotyledon was placed on a moistened filter paper placed on the bottom of a petri dish of 9 cm in diameter. On this treated leaf, 30 apterous viviparous female insects of the cotton aphid were allowed to inhabit, and then the petri dish was covered with the stopper, which was allowed to stand in a room whose temperature was constant at 25°C. After 120 hours, the number of viable insects was counted.

50 **[0103]** As a result, Compound Nos. 5, 9, 14, 21, 42, 43, 44, 45, 58, 61, 64, 65, 72, 73, 76, 78, 81, 90, 92, 103, 106, 113, 114, 115, 116, 118, 119, 122, 126, 128, 129, 130, 132, 135, 136, 139, 142, 144, and 146, as in the form of a wettable powder prepared in accordance with Formulation Example 2 at an active ingredient concentration of 100 µg/ml exhibited an effect as excellent as a lethality of 50% or higher.

55

<Experimental Example 2>

2. Green peach aphid controlling effect

[0104] On a 2-leaf stage Chinese cabbage seedling, 30 adult insects of green peach aphid were allowed to inhabit, and immobilized by allowing to stand for several days. Thereafter, the overground part was cut off, was dipped in a 100 $\mu\text{g/ml}$ treatment solution obtained by a 5000-fold dilution with water of the wettable powder prepared in accordance with Formulation Example 2. This was placed on a moistened filter paper placed on the bottom of a petri dish of 9 cm in diameter, which was allowed to stand in a room whose temperature was constant at 25°C. After 120 hours, the number of viable insects was counted.

[0105] As a result, Compound Nos. 5, 6, 7, 9, 10, 11, 13, 14, 16, 18, 20, 21, 24, 29, 36, 37, 43, 44, 45, 64, 65, 72, 73, 76, 78, 90, 92, 103, 106, 113, 114, 115, 116, 118, 119, 122, 126, 128, 129, 130, 135, 136, 139, 142, 144, 146, and 160, as in the form of a wettable powder prepared in accordance with Formulation Example 2 at an active ingredient concentration of 100 $\mu\text{g/ml}$, exhibited an effect as excellent as a lethality of 80% or higher.

<Experimental Example 3>

3. Greenhouse whitefly controlling effect

[0106] The overground part of the 1-leaf stage cucumber was dipped in a 100 $\mu\text{g/ml}$ treatment solution obtained by a 5000-fold dilution with water of the wettable powder prepared in accordance with Formulation Example 2, and air-dried, and then transferred to a plastic container having a stopper. This was allowed to be inhabited with 20 adult insects of the greenhouse whitefly, covered with the stopper, and allowed to stand in a room whose temperature was constant at 25°C. After 120 hours, the number of viable insects was counted.

[0107] As a result, Compound Nos. 5, 6, 7, 9, 10, 11, 12, 13, 16, 17, 18, 20, 24, 29, 37, 43, 44, 45, 57, 64, 72, 73, 76, 78, 92, 113, 114, 115, 116, 118, 119, 122, 126, 128, 129, 130, 135, 136, 139, 144, and 146, as in the form of a wettable powder prepared in accordance with Formulation Example 2 at an active ingredient concentration of 100 $\mu\text{g/ml}$ exhibited an effect as excellent as a lethality of 50% or higher.

<Experimental Example 4>

4. Two-spotted spider mite controlling effect

[0108] A kidney bean leaf was allowed to be inhabited with 10 adult insects of two-spotted spider mite, and then dipped in a 100 $\mu\text{g/ml}$ treatment solution obtained by a 5000-fold dilution with water of the wettable powder prepared in accordance with Formulation Example 2. After air drying, the cut end was wrapped with a cotton wet with water thereby supplying water to the common bean leaf. After 120 hours of allowing to stand in a room whose temperature was constant at 25°C, the number of viable insects was counted.

[0109] As a result, Compound Nos. 5, 6, 7, 9, 10, 11, 17, 24, 82, 90, 91, 107, 113, 135, 142, 171, and 173, as in the form of a wettable powder prepared in accordance with Formulation Example 2 at an active ingredient concentration of 100 $\mu\text{g/ml}$ exhibited an effect as excellent as a lethality of 50% or higher.

[0110] In the above-indicated Experimental Examples 1 to 4, especially, Compound Nos. 5, 43, 44, 45, 72, 73, 76, 78, 90, 92, 103, 106, 113, 114, 115, 118, 119, 122, 126, 128, 129, 130, 135, 136, 139, 142, 144, and 146 exhibited excellent pest controlling effects.

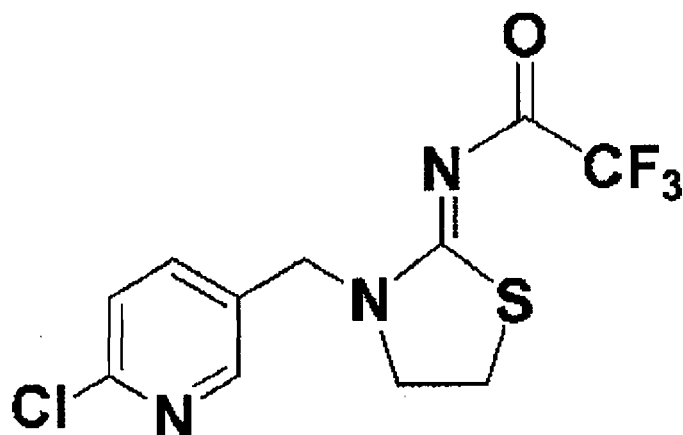
<Comparative Example 1>

5. Greenhouse whitefly controlling effect

[0111] For the purpose of comparison with the insecticide according to the invention, the pest controlling effect of the compound represented by Formula (12) disclosed in the above-indicated Patent Literature (Japanese Unexamined Patent Application Publication No. 63-150275) was validated.

[0112]

[C-23]



Formula (12)

[0113] This compound was prepared into a wettable powder in accordance with Formulation Example 2. The over-ground part of the 1-leaf stage cucumber was dipped in a 100 $\mu\text{g/ml}$ treatment solution obtained by a 5000-fold dilution with water of the prepared wettable powder, and air-dried, and then transferred to a plastic container having a stopper. This was allowed to be inhabited with 20 adult insects of the greenhouse whitefly, covered with the stopper, and allowed to stand in a room whose temperature was constant at 25°C. After 120 hours, the number of viable insects was counted.

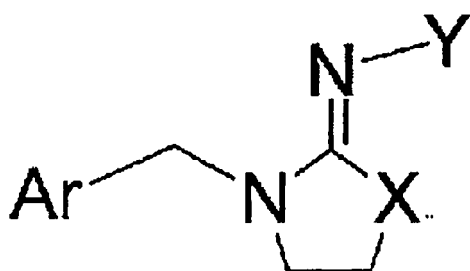
[0114] As a result, at an active ingredient concentration of 100 $\mu\text{g/ml}$, the lethality was 25%. On the contrary, the insecticide according to the invention exhibited the lethality in the similar experiment (see Experimental Example 3) of 50% or higher at an active ingredient concentration of 100 $\mu\text{g/ml}$. Based on these findings, the insecticide according to the invention was revealed to have an excellent pest controlling effect.

[Industrial Applicability]

[0115] A novel imino derivative according to the invention and an insecticide containing the same as an active ingredient exhibit an excellent controlling effect on agro-horticultural and household pests such as Hemiptera pests. Accordingly, the invention can widely be utilized in the fields of agro-horticultural production, livestock, and household sanitation, and can contribute a lot to such fields of the industry.

Claims

1. An imino derivative represented by Formula (I):



Formula (1)

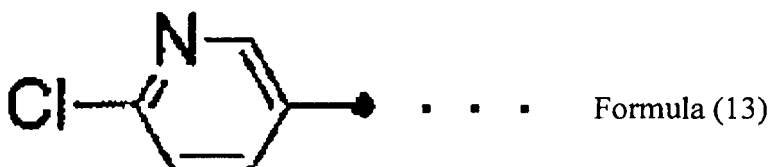
wherein "Ar" denotes a 3-pyridyl group or a 5-thiazolyl group which have a substituent on the ring, wherein the substituent is selected from a halogen atom, a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, and a nitro group; "X" denotes a sulfur atom or CH_2 ; when "Y" is COR_1 , "R₁" denotes a hydrogen atom or a C1 to C5 alkyl group, a halogenated methyl except for a trifluoromethyl group, a C2 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C4)alkoxy(C1 to C5)alkyl group, a C1 to C3 alkoxy carbonyl group, a (C1 to C3)alkylsulfonyl (C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a cyano(C1 to C3)alkyl group, a substituted

or unsubstituted phenoxy (C1 to C3)alkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted imidazolylmethyl group, a furanyl group, a morpholino group, a norbornenyl group, an adamantyl group, an isothiocyanatomethyl group, or substituted or unsubstituted heterocyclic or unsubstituted aromatic ring; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkylmethyl group, the phenoxy (C1 to C3)alkyl group, the pyridylmethyl group and the imidazolylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the heterocyclic or aromatic ring is substituted, each of the substituents is selected from a halogen atom, a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group and an acetylamino group;

when "Y" is CONHD, "D" denotes a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxy-carbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted benzenesulfonyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the benzenesulfonyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, an acetylamino group, and a C1 to C3 alkyl group; when "Y" is CONHCOR₅, "R₅" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a substituted or unsubstituted phenyl group; when the phenyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, and a C1 to C3 alkyl group;

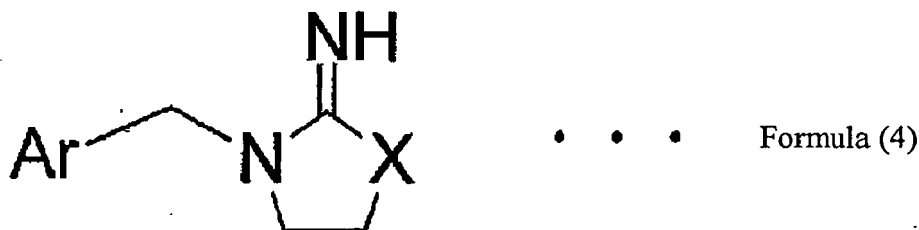
when "Y" is CO₂R₉, "R₉" denotes a hydrogen atom or a C1 to C7 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a tri(C1 to C3 alkyl)silyl(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a substituted or unsubstituted phenyl group, a substituted or unsubstituted phenylmethyl group, a substituted or unsubstituted furanylmethyl group, a substituted or unsubstituted thienylmethyl group, a substituted or unsubstituted pyridylmethyl group, a succinimide group, a 18-crown-6-methyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkylmethyl group, the furanylmethyl group, the thienylmethyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the phenyl group and the phenylmethyl group are substituted, each of the substituents is selected from a halogen atom, a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group, and an acetylamino group; and the abovementioned carbon chains may be substituted with halogen.

2. The imino derivative according to Claim 1 wherein "Ar" in Formula (1) is represented by Formula (13) or (14).



3. A method for producing an imino derivative represented by Formula (1) shown above by reacting one compound

selected from the group consisting of compounds represented by Formula (5), anhydrides represented by Formula (6), and carboxylic acids represented by Formula (7) with a compound represented by Formula (4):



wherein "Ar" and "X" have the same meaning as in claim 1;

15 **ACO-B** Formula (5)

ACOOCOA Formula (6)

20 **ACOOH** Formula (7)

wherein "B" denotes a halogen atom of Cl or Br, I, an OCOA group (a group formed by allowing each of an oxygen atom and a group A to form a single bond with a carbonyl group (CO)) or a hydroxyl group;

when "ACO" is COR₁, "R₁" denotes a hydrogen atom or a C1 to C5 alkyl group, a halogenated methyl except for a trifluoromethyl group, a C2 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C4)alkoxy(C1 to C5)alkyl group, a C1 to C3 alkoxy carbonyl group, a (C1 to C3)alkylsulfonyl (C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkylmethyl group, a cyano(C1 to C3)alkyl group, a substituted or unsubstituted phenoxy (C1 to C3)alkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted imidazolylmethyl group, a furanyl group, a morpholino group, a norbornenyl group, an adamantyl group, an isothiocyanatomethyl group, or substituted or unsubstituted heterocyclic or aromatic ring; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkylmethyl group, the phenoxy (C1 to C3)alkyl group, the pyridylmethyl group and the imidazolylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the heterocyclic or aromatic ring is substituted, each of the substituents is selected from a halogen atom and a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group and an acetylamino group;

when "ACO" is CONHD, "D" denotes a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxy-carbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridylmethyl group, a substituted or unsubstituted benzenesulfonyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group and the pyridylmethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the benzenesulfonyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, an acetylamino group, and a C1 to C3 alkyl group;

when "ACO" is CO₂R₉, "R₉" denotes a hydrogen atom or a C1 to C7 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, a substituted or unsubstituted C6 to C10 aryl group, a substituted or unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a (C1 to C3)alkylthio(C1 to C3)alkyl group, a tri(C1 to C3 alkyl)silyl(C1 to C3)alkyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl-methyl group, a substituted or unsubstituted phenyl group, a substituted or unsubstituted phenylmethyl group, a substituted or unsubstituted furanylmethyl group, a substituted or unsubstituted thienylmethyl group, a substituted or unsubstituted pyridylmethyl group, a succinimide group, a 18-crown-6-methyl group; when the C3 to C7 cycloalkyl

group, the 3-membered to 7-membered heterocycloalkyl group, the 3-membered to 7-membered heterocycloalkyl-methyl group, the furanymethyl group, the thienymethyl group and the pyridymethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the phenyl group and the phenylmethyl group are substituted, each of the substituents is selected from a halogen atom, a C1 to C4 alkyl group, a C1 to C4 halogenated alkyl group, a C1 to C4 alkoxy group, a di(C1 to C4 alkyl)amino group, a nitro group, and an acetylamino group; and the abovementioned carbon chains may be substituted with halogen.

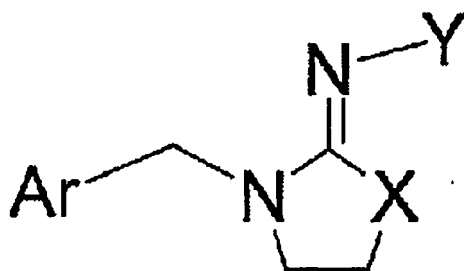
4. A method for producing an imino derivative represented by Formula (9) by reacting an isocyanate compound represented by Formula (8) with a compound represented by Formula (4) shown above:



Formula (8)

wherein "D" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, a C3 to C5 alkynyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a (C1 to C3)alkoxy(C1 to C3)alkyl group, a C1 to C3 alkoxycarbonylmethyl group, a C3 to C7 substituted or unsubstituted cycloalkyl group, a 3-membered to 7-membered substituted or unsubstituted heterocycloalkyl group, a substituted or unsubstituted pyridymethyl group, a substituted or unsubstituted benzenesulfonyl group; when the C3 to C7 cycloalkyl group, the 3-membered to 7-membered heterocycloalkyl group and the pyridymethyl group are substituted, each of the substituents is selected from a halogen atom and a C1 to C3 alkyl group; when the benzenesulfonyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, an acetylamino group, and a C1 to C3 alkyl group;

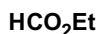
COR₅; and "R₅" denotes a hydrogen atom or a C1 to C5 alkyl group, a C1 to C5 halogenated alkyl group, a C2 to C5 alkenyl group, a C2 to C5 halogenated alkenyl group, an unsubstituted C6 to C10 aryl group, an unsubstituted (C6 to C10)aryl(C1 to C3)alkyl group, a substituted or unsubstituted phenyl group; when the phenyl group is substituted, each of the substituents is selected from a halogen atom, a nitro group, a di(C1 to C4 alkyl)amino group, and a C1 to C3 alkyl group;



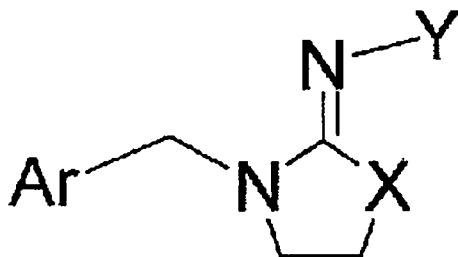
• • • Formula (9)

wherein "Ar", "X" and "Y" is CONHD and "D" has the same meaning as in claim 1.

5. A method for producing an imino derivative represented by Formula (11) by reacting a formate represented by Formula (10) with a compound represented by Formula (4) shown above:



Formula (10)



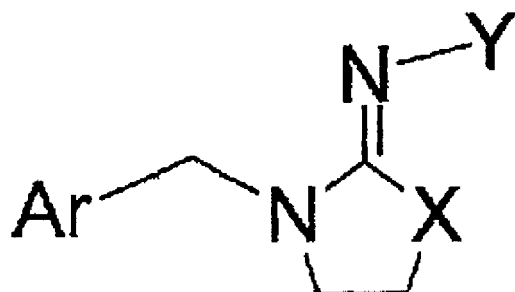
• • • Formula (11)

wherein "Ar", "X" and "Y" denotes CHO.

6. An insecticide containing as an active ingredient an imino derivative according to Claim 1.

Patentansprüche

1. Iminoderivat, repräsentiert durch Formel (1):



Formel (1)

worin "Ar" eine 3-Pyridylgruppe oder eine 5-Thiazolylgruppe bezeichnet, welche am Ring einen Substituenten tragen, wobei der Substituent ausgewählt ist unter einem Halogenatom, einer C1- bis C4-Alkylgruppe, einer halogenierten C1- bis C4-Alkylgruppe, einer C1- bis C4-Alkoxygruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe und einer Nitrogruppe; "X" bezeichnet ein Schwefelatom oder ein CH₂;

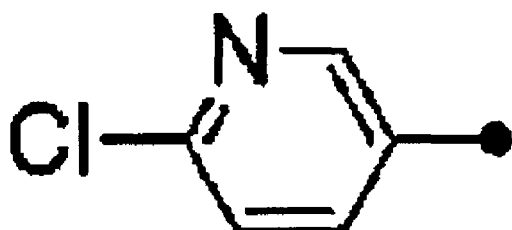
wenn "Y" COR₁ ist, bezeichnet "R₁" ein Wasserstoffatom oder eine C1- bis C5-Alkylgruppe, ein halogeniertes Methyl, ausgenommen eine Trifluormethylgruppe, eine halogenierte C2- bis C5-Alkylgruppe, eine C2- bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine C3- bis C5-Alkylgruppe, eine unsubstituierte C6- bis C10-Arylgruppe, eine unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C4)Alkoxy(C1 bis C5)alkyl-Gruppe, eine C1- bis C3-Alkoxycarbonylgruppe, eine (C1 bis C3)Alkylsulfonyl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkylthio(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte C3- bis C7-Cycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylmethyl-Gruppe, eine Cyano(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte Phenoxy(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte Pyridylmethylgruppe, eine substituierte oder unsubstituierte Imidazolylmethylgruppe, eine Furanylgruppe, eine Morpholinogruppe, eine Norbornenylgruppe, eine Adamantylgruppe, eine Isothiocyanatomethyl-Gruppe oder einen substituierten oder unsubstituierten heterocyclischen oder einen unsubstituierten aromatischen Ring; wenn die C3- bis C7-Cycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylmethyl-Gruppe, die Phenoxy(C1 bis C3)alkylGruppe, die Pyridylmethylgruppe und die Imidazolylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom und einer C1- bis C3-Alkylgruppe; wenn der heterocyclische oder aromatische Ring substituiert ist, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom, einer C1- bis C4-Alkylgruppe, einer halogenierten C1- bis C4-Alkylgruppe, einer C1- bis C4-Alkoxygruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe, einer Nitrogruppe und einer Acetylaminogruppe;

wenn "Y" CONHD ist, dann bezeichnet "D" eine C1- bis C5-Alkylgruppe, eine halogenierte C1- bis C5-Alkylgruppe, eine C2- bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine C3- bis C5-Alkylgruppe, eine unsubstituierte C6- bis C10-Arylgruppe, eine unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkoxy(C1 bis C3)alkyl-Gruppe, eine C1- bis C3-Alkoxycarbonylmethyl-Gruppe, eine substituierte oder unsubstituierte C3- bis C7-Cycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylgruppe, eine substituierte oder unsubstituierte Pyridylmethylgruppe, eine substituierte oder unsubstituierte Benzolsulfonylgruppe; wenn die C3- bis C7-Cycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylgruppe und die Pyridylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom und einer C1- bis C3-Alkylgruppe; wenn die Benzolsulfonylgruppe substituiert ist, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom, einer Nitrogruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe, einer Acetylaminogruppe und einer C1- bis C3-Alkylgruppe;

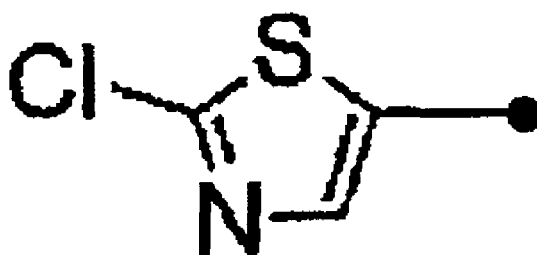
wenn "Y" CONHCOR₅ ist, dann bezeichnet "R₅" ein Wasserstoffatom oder eine C1- bis C5-Alkylgruppe, eine halogenierte C1- bis C5-Alkylgruppe, eine C2- bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine unsubstituierte C6- bis C10-Arylgruppe, eine unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte Phenylgruppe; wenn die Phenylgruppe substituiert ist, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom, einer Nitrogruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe und einer C1- bis C3-Alkylgruppe; wenn "Y" CO₂R₉ ist, dann bezeichnet "R₉" ein Wasserstoffatom oder eine C1- bis C7-

Alkylgruppe, eine halogenierte C1- bis C5-Alkylgruppe, eine C2- bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine C3- bis C5-Alkynylgruppe, eine unsubstituierte C6- bis C10-Arylgruppe, eine unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkoxy(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkylthio(C1 bis C3)alkyl-Gruppe, eine Tri(C1- bis C3-alkyl)silyl(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte C3- bis C7-Cycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylmethyl-Gruppe, eine substituierte oder unsubstituierte Phenylgruppe, eine substituierte oder unsubstituierte Phenylmethylgruppe, eine substituierte oder unsubstituierte Furanylmethylgruppe, eine substituierte oder unsubstituierte Thienylmethylgruppe, eine substituierte oder unsubstituierte Pyridylmethylgruppe, eine Succinimidgruppe, eine [18]Krone-6-methyl-Gruppe; wenn die C3- bis C7-Cycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylmethyl-Gruppe, die Furanylmethylgruppe, die Thienylmethylgruppe und die Pyridylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom und einer C1- bis C3-Alkylgruppe; wenn die Phenylgruppe und die Phenylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom, einer C1- bis C4-Alkylgruppe, einer halogenierten C1- bis C4-Alkylgruppe, einer C1- bis C4-Alkoxygruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe, einer Nitrogruppe und einer Acetylaminogruppe; und die oben genannten Kohlenstoffketten können mit Halogen substituiert sein.

2. Iminoderivat nach Anspruch 1, worin "Ar" in Formel (1) repräsentiert wird durch Formel (13) oder (14).

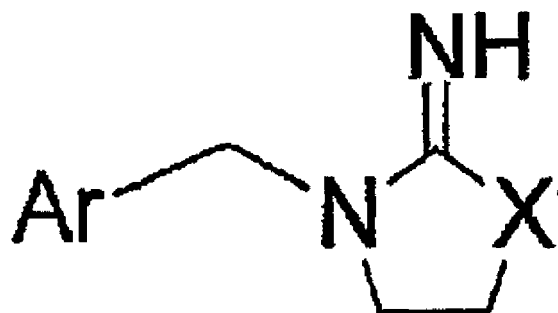


Formel (13)



Formel (14)

3. Verfahren zur Darstellung eines Iminoderivats, welches repräsentiert wird durch die oben abgebildete Formel (1), durch Reaktion einer Verbindung, ausgewählt aus der Gruppe bestehend aus Verbindungen, welche repräsentiert werden durch Formel (5), Anhydriden, welche repräsentiert werden durch Formel (6), sowie Carbonsäuren, welche repräsentiert werden durch Formel (7), mit einer Verbindung, welche repräsentiert wird durch Formel (4):



Formel (4)

worin "Ar" und "X" die gleiche Bedeutung besitzen wie in Anspruch 1;

ACO-B Formel (5)

5 ACOOCoA Formel (6)

ACOOH Formel (7)

10 worin "B" ein Halogenatom aus Cl, Br oder I bezeichnet, eine OCOA-Gruppe (eine Gruppe, gebildet, indem es jeweils einem Sauerstoffatom und einer Gruppe A ermöglicht wird, eine Einfachbindung mit einer Carbonylgruppe (CO) zu bilden), oder eine Hydroxylgruppe;

wenn "ACO" COR₁ ist, dann bezeichnet "R₁" ein Wasserstoffatom oder eine C1- bis C5-Alkylgruppe, ein halogeniertes Methyl, ausgenommen eine Trifluormethylgruppe, eine halogenierte C2- bis C5-Alkylgruppe, eine C2-bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine C3- bis C5-Alkylgruppe, eine unsubstituierte C6- bis C10-Arylgruppe, eine unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C4)Alkoxy(C1 bis C5)alkyl-Gruppe, eine C1- bis C3-Alkoxy-carbonylgruppe, eine (C1 bis C3)Alkylsulfonyl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkylthio(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte C3- bis C7-Cycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylmethyl-Gruppe, eine Cyano(C1-C3)alkyl-Gruppe, eine substituierte oder unsubstituierte Phenoxy(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte Pyridylmethylgruppe, eine substituierte oder unsubstituierte Imidazolylmethylgruppe, eine Furanylgruppe, eine Morpholinogruppe, eine Norbornenylgruppe, eine Adamantylgruppe, eine Isothiocyanatomethyl-Gruppe oder einen substituierten oder unsubstituierten, heterocyclischen oder aromatischen Ring; wenn die C3- bis C7-Cycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylmethyl-Gruppe, die Phenoxy(C1 bis C3)alkyl-Gruppe, die Pyridylmethylgruppe und die Imidazolylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom und einer C1-bis C3-Alkylgruppe; wenn der heterocyclische oder aromatische Ring substituiert ist, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom und einer C1- bis C4-Alkylgruppe, einer halogenierten C1- bis C4-Alkylgruppe, einer C1- bis C4-Alkoxygruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe, einer Nitrogruppe und einer Acetylaminogruppe;

30 wenn "ACO" CONHD ist, dann bezeichnet "D" eine C1- bis C5-Alkylgruppe, eine halogenierte C1- bis C5-Alkylgruppe, eine C2- bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine C3- bis C5-Alkylgruppe, eine unsubstituierte C6- bis C10-Arylgruppe, eine unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkoxy(C1 bis C3)alkyl-Gruppe, eine C1- bis C3-Alkoxy-carbonylmethyl-Gruppe, eine substituierte oder unsubstituierte C3- bis C7-Cycloalkyl-Gruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylgruppe, eine substituierte oder unsubstituierte Pyridylmethylgruppe, eine substituierte oder unsubstituierte Benzolsulfonylgruppe; wenn die C3- bis C7-Cycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylgruppe und die Pyridylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom und einer C1- bis C3-Alkylgruppe; wenn die Benzolsulfonylgruppe substituiert ist, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom, einer Nitrogruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe, einer Acetylaminogruppe und einer C1- bis C3-Alkylgruppe;

40 wenn "ACO" CO₂R₉ ist, dann bezeichnet "R₉" ein Wasserstoffatom oder eine C1- bis C7-Alkylgruppe, eine halogenierte C1- bis C5-Alkylgruppe, eine C2- bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine C3- bis C5-Alkylgruppe, eine substituierte oder unsubstituierte C6- bis C10-Arylgruppe, eine substituierte oder unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkoxy(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkylthio(C1 bis C3)alkyl-Gruppe, eine Tri(C1- bis C3-alkyl)silyl(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte C3- bis C7-Cycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylmethyl-Gruppe, eine substituierte oder unsubstituierte Phenylgruppe, eine substituierte oder unsubstituierte Phenylmethylgruppe, eine substituierte oder unsubstituierte Furanylmethylgruppe, eine substituierte oder unsubstituierte Thienylmethylgruppe, eine substituierte oder unsubstituierte Pyridylmethylgruppe, eine Succinimidgruppe, eine [18]Krone-6-methyl-Gruppe; wenn die C3-bis C7-Cycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylmethyl-Gruppe, die Furanylmethylgruppe, die Thienylmethylgruppe und die Pyridylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom und einer C1- bis C3-Alkylgruppe; wenn die Phenylgruppe und die Phenylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom, einer C1- bis C4-Alkylgruppe, einer halogenierten C1- bis C4-Alkylgruppe, einer C1- bis C4-Alkoxygruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe, einer Nitrogruppe und einer Acetylaminogruppe; und

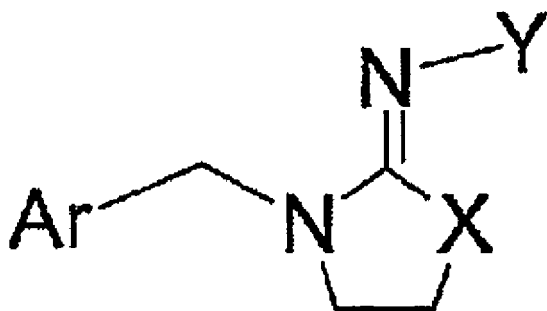
55 die oben genannten Kohlenstoffketten können mit Halogen substituiert sein.

4. Verfahren zur Darstellung eines Iminoderivats, repräsentiert durch Formel (9), durch Reaktion einer Isocyanat-Verbindung, repräsentiert durch Formel (8), mit einer Verbindung, repräsentiert durch die oben abgebildete Formel (4):



worin "D" ein Wasserstoffatom bezeichnet oder eine C1- bis C5-Alkylgruppe, eine halogenierte C1- bis C5-Alkylgruppe, eine C2- bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine C3- bis C5-Alkynylgruppe, eine unsubstituierte C6- bis C10-Arylgruppe, eine unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine (C1 bis C3)Alkoxy(C1 bis C3)alkyl-Gruppe, eine C1- bis C3-Alkoxycarbonylmethyl-Gruppe, eine substituierte oder unsubstituierte C3- bis C7-Cycloalkylgruppe, eine 3-gliedrige bis 7-gliedrige, substituierte oder unsubstituierte Heterocycloalkylgruppe, eine substituierte oder unsubstituierte Pyridylmethylgruppe, eine substituierte oder unsubstituierte Benzolsulfonylgruppe; wenn die C3- bis C7-Cycloalkylgruppe, die 3-gliedrige bis 7-gliedrige Heterocycloalkylgruppe und die Pyridylmethylgruppe substituiert sind, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom und einer C1- bis C3-Alkylgruppe; wenn die Benzolsulfonylgruppe substituiert ist, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom, einer Nitrogruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe, einer Acetylaminogruppe und einer C1- bis C3-Alkylgruppe;

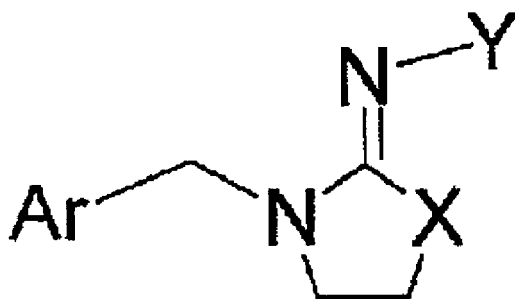
COR₅; und "R₅" bezeichnet ein Wasserstoffatom oder eine C1- bis C5-Alkylgruppe, eine halogenierte C1- bis C5-Alkylgruppe, eine C2- bis C5-Alkenylgruppe, eine halogenierte C2- bis C5-Alkenylgruppe, eine unsubstituierte C6- bis C10-Arylgruppe, eine unsubstituierte (C6 bis C10)Aryl(C1 bis C3)alkyl-Gruppe, eine substituierte oder unsubstituierte Phenylgruppe; wenn die Phenylgruppe substituiert ist, dann ist der jeweilige Substituent ausgewählt unter einem Halogenatom, einer Nitrogruppe, einer Di(C1- bis C4-alkyl)amino-Gruppe und einer C1- bis C3-Alkylgruppe;



Formel (9)

worin "Ar", "X" und "Y" CONHD sind und "D" die gleiche Bedeutung besitzt wie in Anspruch 1.

5. Verfahren zur Darstellung eines Iminoderivats, repräsentiert durch Formel (11), durch Reaktion eines Formiats, repräsentiert durch Formel (10), mit einer Verbindung, repräsentiert durch die oben abgebildete Formel (4):



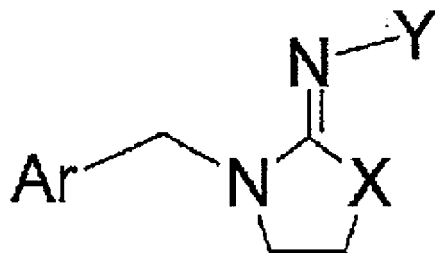
Formel (11)

worin "Ar", "X" und "Y" CHO bezeichnen.

6. Insektizid, enthaltend als einen Wirkstoff ein Iminoderivat nach Anspruch 1.

Revendications

1. Dérivé d'imino représenté par la formule (I) :



Formule (1)

dans laquelle « Ar » désigne un groupe 3-pyridyle ou un groupe 5-thiazolyle qui a un substituant sur le cycle, dans laquelle le substituant est choisi parmi un atome d'halogène, un groupe alkyle en C₁ à C₄, un groupe alkyle halogéné en C₁ à C₄, un groupe alcoxy en C₁ à C₄, un groupe di(alkyle en C₁ à C₄)amino, et un groupe nitro ; « X » désigne un atome de soufre ou de CH₂ ; quand « Y » est COR₁, « R₁ » désigne un atome d'hydrogène ou un groupe alkyle en C₁ à C₅, un groupe méthyle halogéné à l'exception d'un groupe trifluorométhyle, un groupe alkyle halogéné en C₂ à C₅, un groupe alcényle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe alcynyle en C₃ à C₅, un groupe aryle en C₆ à C₁₀ non substitué, un groupe (aryle en C₆ à C₁₀) (alkyle en C₁ à C₃) non substitué, un groupe (alcoxy en C₁ à C₄) (alkyle en C₁ à C₅), un groupe (alcoxy en C₁ à C₃)carbonyle, un groupe (alkylsulfonyl en C₁ à C₃) (alkyle en C₁ à C₃), un groupe (alkylthio en C₁ à C₃) (alkyle en C₁ à C₃), un groupe cycloalkyle substitué ou non substitué en C₃ à C₇, un groupe hétérocycloalkyle substitué ou non substitué de 3 membres à 7 membres, un groupe hétérocycloalkylméthyle substitué ou non substitué de 3 membres à 7 membres, un groupe cyanoalkyle (en C₁ à C₃), un groupe phénoxyalkyle (en C₁ à C₃) substitué ou non substitué, un groupe pyridylméthyle substitué ou non substitué, un groupe imidazolylméthyle substitué ou non substitué, un groupe furanyle, un groupe morpholino, un groupe norbornényle, un groupe adamantyle, un groupe isothiocyanatométhyle, ou un cycle aromatique non substitué ou hétérocyclique substitué ou non substitué ; quand le groupe cycloalkyle en C₃ à C₇, le groupe hétérocycloalkyle de 3 membres à 7 membres, le groupe hétérocycloalkylméthyle de 3 membres à 7 membres, le groupe phénoxyalkyle (en C₁ à C₃), le groupe pyridylméthyle et le groupe imidazolylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène et un groupe alkyle en C₁ à C₃ ; quand le cycle hétérocyclique ou aromatique est substitué, chacun des substituants est choisi parmi un atome d'halogène, un groupe alkyle en C₁ à C₄, un groupe alkyle en C₁ à C₄ halogéné, un groupe alcoxy en C₁ à C₄, un groupe di(alkyle en C₁ à C₄)amino, un groupe nitro et un groupe acétylamino ; quand « Y » est CONHD, « D » désigne un groupe alkyle en C₁ à C₅, un groupe alkyle halogéné en C₁ à C₅, un groupe alcényle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe alcynyle en C₃ à C₅, un groupe aryle en C₆ à C₁₀ non substitué, un groupe (aryle en C₆ à C₁₀) (alkyle en C₁ à C₃) non substitué, un groupe (alcoxy en C₁ à C₃) (alkyle en C₁ à C₃), un groupe alcoxycarbonylméthyle en C₁ à C₃, un groupe cycloalkyle substitué ou non substitué en C₃ à C₇, un groupe hétérocycloalkyle substitué ou non substitué de 3 membres à 7 membres, un groupe pyridylméthyle substitué ou non substitué, un groupe benzènesulfonyl substitué ou non substitué ; quand le groupe cycloalkyle en C₃ à C₇, le groupe hétérocycloalkyle de 3 membres à 7 membres et le groupe pyridylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène et un groupe alkyle en C₁ à C₃ ; quand le groupe benzènesulfonyl est substitué, chacun des substituants est choisi parmi un atome d'halogène, un groupe nitro, un groupe di(alkyle en C₁ à C₄)amino, un groupe acétylamino, et un groupe alkyle en C₁ à C₃ ; quand « Y » est CONHCOR₅, « R₅ » désigne un atome d'hydrogène ou un groupe alkyle en C₁ à C₅, un groupe alkyle halogéné en C₁ à C₅, un groupe alcényle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe aryle en C₆ à C₁₀ non substitué, un groupe (aryle en C₆ à C₁₀) (alkyle en C₁ à C₃) non substitué, un groupe phényle substitué ou non substitué ; quand le groupe phényle est substitué, chacun des substituants est choisi parmi un atome d'halogène, un groupe nitro, un groupe di(alkyle en C₁ à C₄)amino et un groupe alkyle en C₁ à C₃ ;

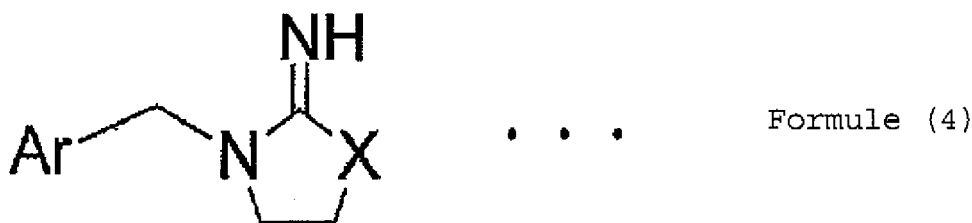
quand « Y » est CO₂R₉, « R₉ » désigne un atome d'hydrogène ou un groupe alkyle en C₁ à C₇, un groupe alkyle halogéné en C₁ à C₅, un groupe alcényle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe alcynyle en C₃ à C₅, un groupe aryle en C₆ à C₁₀ non substitué, un groupe (aryle en C₆ à C₁₀) (alkyle en C₁ à C₃) non substitué, un groupe (alcoxy en C₁ à C₃) (alkyle en C₁ à C₃), un groupe (alkylthio en C₁ à C₃) (alkyle en C₁ à C₃), un groupe tri(alkyle en C₁ à C₃)silyl(alkyle en C₁ à C₃), un groupe cycloalkyle substitué ou non substitué en C₃ à C₇, un groupe hétérocycloalkyle substitué ou non substitué de 3 membres à 7 membres, un groupe hétérocycloalkylméthyle substitué ou non substitué de 3 membres à 7 membres, un groupe phényle substitué ou non substitué, un groupe phénylméthyle substitué ou non substitué, un groupe furanylméthyle substitué ou non substitué, un

groupe thiénylméthyle substitué ou non substitué, un groupe pyridylméthyle substitué ou non substitué, un groupe succinimide, un groupe 18-couronne-6-méthyle ; quand le groupe cycloalkyle en C₃ à C₇, le groupe hétérocycloalkyle de 3 membres à 7 membres, le groupe hétérocycloalkylméthyle de 3 membres à 7 membres, le groupe furanylméthyle, le groupe thiénylméthyle et le groupe pyridylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène et un groupe alkyle en C₁ à C₃ ; quand le groupe phényle et le groupe phénylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène, un groupe alkyle en C₁ à C₄, un groupe alkyl en C₁ à C₄ halogéné, un groupe alcoxy en C₁ à C₄, un groupe di(alkyle en C₁ à C₄)amino, un groupe nitro et un groupe acétylamino ; et les chaînes carbonées susmentionnées peuvent être substituées par un halogène.

2. Dérivé d'imino selon la revendication 1, dans lequel « Ar » dans la formule (1) est représenté par la formule (13) ou (14).



3. Procédé de production d'un dérivé d'imino représenté par la formule (1) ci-dessus consistant à faire réagir un composé choisi dans le groupe constitué par les composés représentés par la formule (5), les anhydrides représentés par la formule (6) et les acides carboxyliques représentés par la formule (7) avec un composé représenté par la formule (4) :



dans laquelle « Ar » et « X » ont la même signification que dans la revendication 1 ;



dans lesquelles « B » désigne un atome d'halogène de Cl ou Br, I, un groupe OCOA (un groupe formé en permettant à chacun parmi un atome d'oxygène et un groupe A de former une simple liaison avec un groupe carbonyle (CO)) ou un groupe hydroxyle ;

quand « ACO » est COR₁, « R₁ » désigne un atome d'hydrogène ou un groupe alkyle en C₁ à C₅, un groupe méthyle halogéné à l'exception d'un groupe trifluorométhyle, un groupe alkyl halogéné en C₂ à C₅, un groupe alcényle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe alcynyle en C₃ à C₅, un groupe aryle en C₆ à C₁₀ non substitué, un groupe (aryle en C₆ à C₁₀) (alkyle en C₁ à C₃) non substitué, un groupe (alcoxy en C₁ à C₄) (alkyle en C₁ à C₅), un groupe (alcoxy en C₁ à C₃) carbonyle, un groupe (alkylsulfonyle en C₁ à C₃) (alkyle en C₁ à C₃), un groupe (alkylthio en C₁ à C₃) (alkyle en C₁ à C₃), un groupe cycloalkyle substitué ou non substitué en C₃ à C₇, un

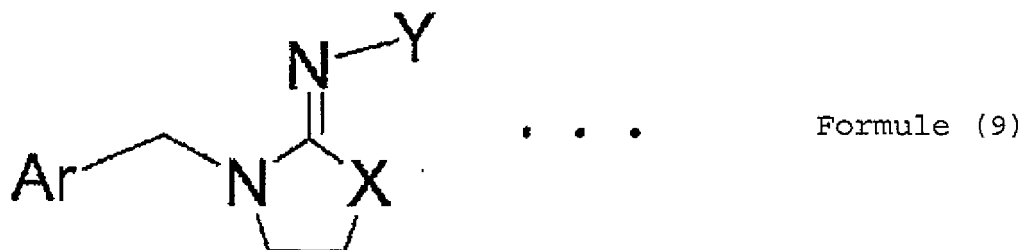
groupe hétérocycloalkyle substitué ou non substitué de 3 membres à 7 membres, un groupe hétérocycloalkylméthyle substitué ou non substitué de 3 membres à 7 membres, un groupe cyanoalkyle (en C₁ à C₃), un groupe phénoxyalkyle (en C₁ à C₃) substitué ou non substitué, un groupe pyridylméthyle substitué ou non substitué, un groupe imidazolylméthyle substitué ou non substitué, un groupe furanyle, un groupe morpholino, un groupe norbornényle, un groupe adamantyle, un groupe isothiocyanatométhyle, ou un cycle hétérocyclique ou aromatique substitué ou non substitué ; quand le groupe cycloalkyle en C₃ à C₇, le groupe hétérocycloalkyle de 3 membres à 7 membres, le groupe hétérocycloalkylméthyle de 3 membres à 7 membres, le groupe phénoxyalkyle (en C₁ à C₃), le groupe pyridylméthyle et le groupe imidazolylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène et un groupe alkyle en C₁ à C₃ ; quand le cycle hétérocyclique ou aromatique est substitué, chacun des substituants est choisi parmi un atome d'halogène, un groupe alkyle en C₁ à C₄, un groupe alkyle en C₁ à C₄ halogéné, un groupe alcoxy en C₁ à C₄, un groupe diamino(alkyle en C₁ à C₄), un groupe nitro et un groupe acétylamino ; quand « ACO » est CONHD, « D » désigne un groupe alkyle en C₁ à C₅, un groupe alkyle halogéné en C₁ à C₅, un groupe alcényle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe alcynyle en C₃ à C₅, un groupe aryle en C₆ à C₁₀ non substitué, un groupe (aryle en C₆ à C₁₀) (alkyle en C₁ à C₃) non substitué, un groupe (alcoxy en C₁ à C₃) (alkyle en C₁ à C₃), un groupe alcoxycarbonylméthyle en C₁ à C₃, un groupe cycloalkyle substitué ou non substitué en C₃ à C₇, un groupe hétérocyclolalkyle substitué ou non substitué de 3 membres à 7 membres, un groupe pyridylméthyle substitué ou non substitué, un groupe benzènesulfonyl substitué ou non substitué ; quand le groupe cycloalkyle en C₃ à C₇, le groupe hétérocycloalkyle de 3 membres à 7 membres et le groupe pyridylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène et un groupe alkyle en C₁ à C₃ ; quand le groupe benzènesulfonyl est substitué, chacun des substituants est choisi parmi un atome d'halogène, un groupe nitro, un groupe di(alkyle en C₁ à C₄)amino, un groupe acétylamino, et un groupe alkyle en C₁ à C₃ ; quand « ACO » est CO₂R₉, « R₉ » désigne un atome d'hydrogène ou un groupe alkyle en C₁ à C₇, un groupe alkyle halogéné en C₁ à C₅, un groupe alcényle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe alcynyle en C₃ à C₅, un groupe aryle en C₆ à C₁₀ substitué ou non substitué, un groupe (aryle en C₆ à C₁₀)(alkyle en C₁ à C₃) substitué ou non substitué, un groupe (alcoxy en C₁ à C₃)(alkyle en C₁ à C₃), un groupe (alkylthio en C₁ à C₃)(alkyle en C₁ à C₃), un groupe tri(alkyle en C₁ à C₃)silyl(alkyle en C₁ à C₃), un groupe cycloalkyle substitué ou non substitué en C₃ à C₇, un groupe hétérocycloalkyle substitué ou non substitué de 3 membres à 7 membres, un groupe hétérocycloalkylméthyle substitué ou non substitué de 3 membres à 7 membres, un groupe phényle substitué ou non substitué, un groupe phénylméthyle substitué ou non substitué, un groupe furanylméthyle substitué ou non substitué, un groupe thiénylméthyle substitué ou non substitué, un groupe pyridylméthyle substitué ou non substitué, un groupe succinimide, un groupe 18-couronne-6-méthyle ; quand le groupe cycloalkyle en C₃ à C₇, le groupe hétérocycloalkyle de 3 membres à 7 membres, le groupe hétérocycloalkylméthyle de 3 membres à 7 membres, le groupe furanylméthyle, le groupe thiénylméthyle et le groupe pyridylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène et un groupe alkyle en C₁ à C₃ ; quand le groupe phényle et le groupe phénylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène, un groupe alkyle en C₁ à C₄, un groupe alkyle en C₁ à C₄ halogéné, un groupe alcoxy en C₁ à C₄, un groupe di(alkyle en C₁ à C₄)amino, un groupe nitro et un groupe acétylamino ; et les chaînes carbonées susmentionnées peuvent être substituées par un halogène.

4. Procédé de production d'un dérivé d'imino représenté par la formule (9) consistant à faire réagir un composé isocyanate représenté par la formule (8) avec un composé représenté par la formule (4) illustrée ci-dessus :



dans laquelle « D » désigne un atome d'hydrogène ou un groupe alkyle en C₁ à C₅, un groupe alkyle halogéné en C₁ à C₅, un groupe alcényle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe alcynyle en C₃ à C₅, un groupe aryle en C₆ à C₁₀ non substitué, un groupe (aryle en C₆ à C₁₀) (alkyle en C₁ à C₃) non substitué, un groupe (alcoxy en C₁ à C₃) (alkyle en C₁ à C₃), un groupe (alcoxy en C₁ à C₃)carbonylméthyle, un groupe cycloalkyle substitué ou non substitué en C₃ à C₇, un groupe hétérocycloalkyle substitué ou non substitué de 3 membres à 7 membres, un groupe pyridylméthyle substitué ou non substitué, un groupe benzènesulfonyl substitué ou non substitué ; quand le groupe cycloalkyle en C₃ à C₇, le groupe hétérocycloalkyle de 3 membres à 7 membres, et le groupe pyridylméthyle sont substitués, chacun des substituants est choisi parmi un atome d'halogène et un groupe alkyle en C₁ à C₃ ; quand le groupe benzènesulfonyl est substitué, chacun des substituants est choisi parmi un atome d'halogène, un groupe nitro, un groupe di(alkyle en C₁ à C₄)amino, un groupe acétylamino, et un groupe alkyle en C₁ à C₃ ; COR₅ ; et « R₅ » désigne un atome d'hydrogène ou un groupe alkyle en C₁ à C₅, un groupe alkyle halogéné en C₁ à C₅, un groupe alcynyle en C₂ à C₅, un groupe alcényle halogéné en C₂ à C₅, un groupe aryle en C₆ à C₁₀ non substitué, un groupe (aryle en C₆ à C₁₀) (alkyl en C₁ à C₃) non substitué, un groupe phényle substitué ou non

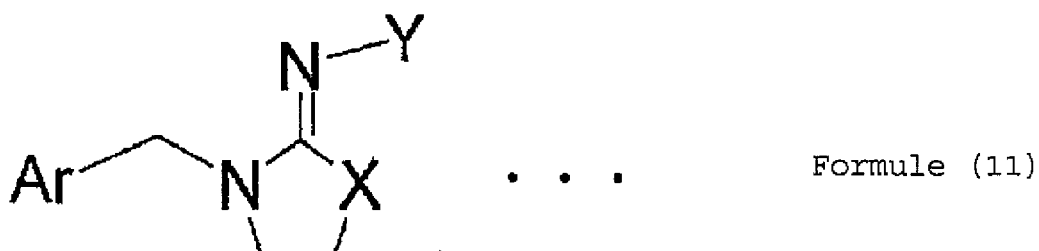
substitué ; quand le groupe phényle est substitué, chacun des substituants est choisi parmi un atome d'halogène, un groupe nitro, un groupe di(alkyle en C₁ à C₄)amino, et un groupe alkyle en C₁ à C₃ ;



15 dans laquelle « Ar », « X » et « Y » sont CONHD et « D » a la même signification que dans la revendication 1.

5. Procédé de production d'un dérivé d'imino représenté par la formule (11) en faisant réagir un formiate représenté par la formule (10) avec un composé représenté par la formule (4) illustrée ci-dessus :

20 HCO₂Et Formule (10)



30 dans laquelle « Ar », « X » et « Y » désignent CHO.

6. Insecticide contenant comme ingrédient actif un dérivé d'imino selon la revendication 1.

Fig. 1

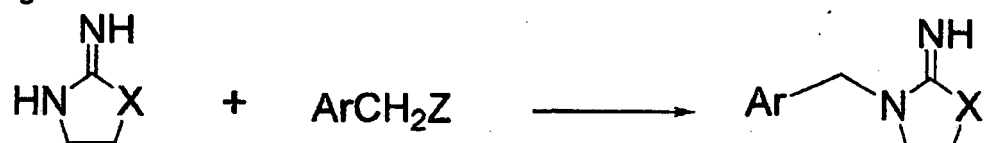


Fig. 2

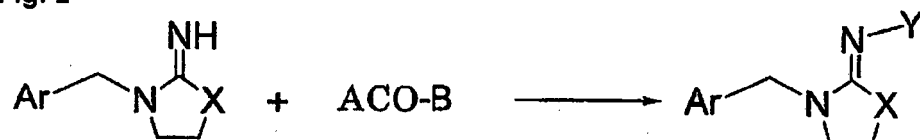


Fig. 3

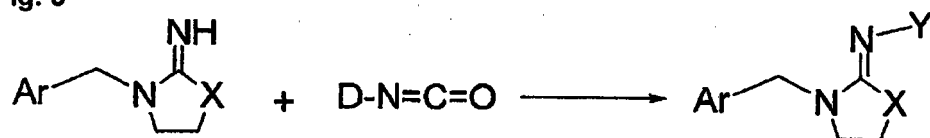
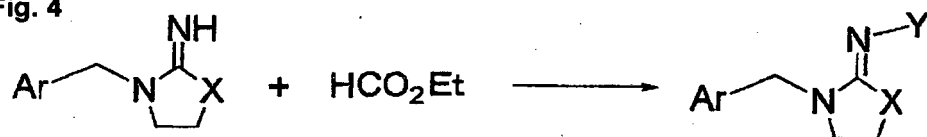


Fig. 4



REFERENCES CITED IN THE DESCRIPTION

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